# Maximizing Ozone Signals Among Chemical, Meteorological, and Climatological Variability

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22 Abstract23

24 The detection of meteorological, chemical, or other signals in modeled or observed air quality 25 data – such as an estimate of a temporal trend in surface ozone data, or an estimate of the mean 26 ozone of a particular region during a particular season – is a critical component of modern 27 atmospheric chemistry. However, the magnitude of a surface air quality signal is generally small 28 compared to the magnitude of the underlying chemical, meteorological, and climatological 29 variabilities (and their interactions) that exist both in space and in time, and which include 30 variability in emissions and surface processes. This can present difficulties for both policy-31 makers and researchers as they attempt to identify the influence or 'signal' of climate trends (e.g. 32 any pauses in warming trends), the impact of enacted emission reductions policies (e.g. United 33 States NO<sub>x</sub> State Implementation Plans), or an estimate of the mean state of highly variable data 34 (e.g. summertime ozone over the Northeastern United States). Here we examine the scale-35 dependence of the variability of simulated and observed surface ozone data within the United 36 States and the likelihood that a particular choice of temporal or spatial averaging scales produce 37 a misleading estimate of a particular ozone signal. Our main objective is to develop strategies 38 that reduce the likelihood of overconfidence in simulated ozone estimates. We find that while 39 increasing the extent of both temporal and spatial averaging can enhance signal detection 40 capabilities by reducing the 'noise' from variability, a strategic combination of particular 41 temporal and spatial averaging scales can maximize signal detection capabilities over much of 42 the Continental US. For signals that are large compared to the meteorological variability (e.g. 43 strong emissions reductions), shorter averaging periods and smaller spatial averaging regions 44 may be sufficient, but for many signals that are smaller than or comparable in magnitude to the 45 underlying meteorological variability, we recommend temporal averaging of 10 - 15 years 46 combine with some level of spatial averaging (up to several hundred kilometers). If this level of 47 averaging is not practical (e.g. the signal being examined is at a local scale), we recommend 48 some exploration of the spatial and temporal variability to provide context and confidence in the 49 robustness of the result. These results are consistent between simulated and observed data, and 50 within a single model with different sets of parameters. The strategies selected in this study are 51 not limited to surface ozone data and could potentially maximize signal detection capabilities 52 within a broad array of climate and chemical observations or model output.

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#### 66 **1 Introduction**

67 The capability to detect air quality signals – be they meteorological, chemical, or of some 68 other type – is a fundamental component of modern climate science and atmospheric chemistry. The debate over the existence or length of a global warming hiatus (Lewandowski et al., 2015; 69 70 Roberts et al., 2015; Medhaug et al., 2017) and research examining the time of emergence of 71 climatological (Weatherhead et al., 2002; Deser et al., 2012; Hawkins and Sutton, 2012; Elía et 72 al., 2013; Schurer et al., 2013), meteorological (Giorgi and Bi, 2009; King et al., 2015), chemical 73 (Camalier et al., 2007; Strode and Pawson, 2013; Barnes et al., 2016; Garcia-Menendez et al., 74 2017), and other sectoral signals (e.g. Monier et al., 2016) embody an accumulation of 75 techniques and strategies for filtering noise (due to natural variability) and maximizing the 76 capability to detect statistically significant signals and trends in noisy data. It is well established 77 that temporal averaging (e.g. Lewandowski et al., 2015) and spatial averaging (e.g. Frost et al., 78 2006; Hawkins and Sutton, 2012; Barnes et al., 2016) can enhance signal detection capabilities 79 in atmospheric data. Here we extend this research by quantifying the impact of both spatial and 80 temporal averaging – individually and in combination – of surface ozone on the magnitude of the 81 calculated variability, which is largely driven by the influence of meteorological variability on 82 atmospheric chemistry (e.g. Jacob and Winner, 2009). We offer recommendations for 83 strategically averaging in space and time to maximize signal detection capabilities. In particular, 84 we examine estimates of mean ozone and of the ozone variability that results from meteorology, 85 although our approach can be generalized to other air quality applications.

86 For observed ozone data, strategies for reducing spatial and temporal noise are limited: a 87 longer time series is needed, more observations need to be made, or the spatial region over which 88 the ozone observations are being averaged needs to be enlarged. For surface ozone estimates 89 using models, however, there exist a variety of strategies for reducing the noise (due to chemical 90 and meteorological variability) relative to the strength of the signal, although they cluster into 91 three main types. The first strategy is to average or combine multiple runs of structurally 92 different models under the assumption that errors, biases, and uncertainties within the individual 93 models are reduced and the multi-model or multi-dataset mean is a best estimate of the actual, 94 aggregated ozone field. This is most notably done with multi-model ensembles within the 95 Atmospheric Chemistry and Climate Model Intercomparison Project (ACCMIP) framework 96 (Lamarque et al., 2013; Young et al., 2013; Stevenson et al., 2013), and this approach tends to

97 assume that all members in the ensemble are independent and equally skillful. This assumption,

98 however, may result in a loss of some valuable information (Knutti, 2010). Another form of this

99 strategy is to run multiple model runs within a single model, but under different initial conditions

100 or sets of parametric assumptions (e.g. Deser et al, 2012; Monier et al., 2013, 2015; Kay et al.,

101 2015; Garcia-Menendez et al., 2015, 2017). This approach cannot address structural uncertainties

and internal (unforced) variability between models, but is capable of identifying parametric

103 uncertainties within a single model.

104 The second strategy to reduce ozone variability is to expand the temporal averaging window, 105 which can influence the interpretation of the determined ozone value (e.g. Brown-Steiner et al., 2015). The Environmental Protection Agency (EPA) National Ambient Air Quality Standard 106 107 (NAAQS) for ozone (US EPA, 2015) explicitly takes this into account, both in the length of the 108 averaging period (daily maximum 8-hour average) and the selection criteria for the standard 109 (fourth-highest over the previous 3 years). The calculated ozone variability can be further 110 reduced by utilizing even longer averaging periods, such as monthly (e.g. Rasmussen et al., 111 2012), seasonal (e.g. Fiore et al., 2014; Barnes et al., 2016), annual, or decadal mean values (e.g. 112 Garcia-Menendez et al., 2017). This strategy is analogous to the averaging of meteorological 113 data to derive a climate signal, and just as Lewandowsky et al. (2015) recommend averaging 17 114 or more years in order to achieve climatological estimates of temperature trends, there is a 115 growing body of literature recommending averaging short time scale chemical variability (what 116 could be called chemical weather, see Lawrence, 2005) for 15 or more years (e.g. Garcia-117 Menendez et al, 2017) in order to achieve an estimate of the what could be called the chemical 118 climate (see Möller, 2010).

119 The third strategy to reduce ozone variability is to average surface ozone values over larger 120 spatial regions, and while there is a significant body of literature discussing the capability and 121 interpretation of coarse resolution model representations of the sub-grid scale heterogeneity 122 (Pyle and Zavody, 1990; Searle et al., 1998, Wild et al., 2006), there are few that strategically 123 expand the spatial scale over which averaging is applied in order to maximize signal detection 124 capabilities. This strategy has been applied in other fields of the atmospheric sciences as well as 125 for general gridded datasets (e.g. Pogson and Smith, 2015), and spatial averaging has been 126 suggested as a means of reducing temperature variability and smoothing biases at the smallest 127 spatial scales within a single model run (Räisänen and Ylhäsi, 2011). This "scale problem" has

also been noted as an important consideration when analyzing aerosol indirect effects

129 (McComiskey and Feingold, 2012) and for the detection and attribution of extreme weather130 events (Angélil et al., 2017).

131 Our objective in this study is to provide a framework for selecting spatial and temporal 132 averaging scales that reduces the uncertainty in analyzing ozone signals and limits the likelihood 133 of over-confidence in an estimate of surface ozone that arises from meteorological variability. 134 This type of framework can be useful from two different research perspectives. The first research 135 perspective has a priori an ozone estimate (either observed or modeled) at a certain spatial and 136 temporal scale (e.g. a 3-year simulation of surface ozone over the Northeastern US) and wants to 137 quantify the likelihood that this estimate is representative of the long-term ozone behavior (rather 138 than overly sensitive to meteorological variability of that particular 3-year period). Since ozone 139 is strongly influenced by natural fluctuations in meteorology (Jacob and Winner, 2009; Jhun et 140 al., 2015) and since extremes in surface ozone and temperature tend to co-occur (Schnell and 141 Prather, 2017), atypically hot or cold periods can strongly influence ozone behavior over short 142 time scales.

143 The second research perspective is to identify an ozone signal of a certain magnitude (or 144 threshold) and decide what spatial and temporal averaging scales are needed to best identify that 145 signal. The ozone signal could be large (e.g. determining the effectiveness or compliance with a 146 5 ppbv incremental reduction of the EPA NAAQS for ozone (US EPA, 2015)) or small (e.g. 147 identifying annual ozone trends within the US, which Cooper et al. (2012) show can be on the 148 order of 0.10 - 0.45 ppby), and can be highly sensitivity to spatial and temporal heterogeneity 149 and meteorological variability. Barnes et al. (2016) found that surface ozone trends over 20-year 150 periods can vary by  $\pm 2$  ppbv due solely to climate variability, while interannual variability can 151 be on the order of  $\pm$  15 ppbv (Fiore et al., 2003; Tilmes et al., 2012; Lin et al., 2014) and day-to-152 day variability can be even larger, extending regularly from near-background levels of 40-50153 ppbv up to 100 ppbv during the summertime (Fiore et al., 2014).

In this study, we quantify the impact of both temporal and spatial averaging on the calculated
ozone variability – due solely to meteorological variability – in order to maximize the capability
to detect signals. We use simulated ozone (with the Community Atmosphere Model with
Chemistry, CAM-chem) and observational data (with the EPA's Clean Air Status and Trends
Network, CASTNET) within the United States in order to answer the following four questions:

159 (1) Within a given dataset (model or observations), with both spatial and temporal coverage, 160 what is the magnitude of the ozone variability due to meteorology at the smallest scale, and how 161 does spatial and temporal averaging reduce this variability? (2) Are there combinations of 162 temporal and spatial averaging scales that maximize the signal detection capability for surface 163 ozone data? (3) How sensitive are the above strategies to different configurations (i.e. emissions, 164 meteorology, and climate) of the CAM-chem modeling framework? And (4) How could they be 165 applied to other datasets (chemical, meteorological, or climatological)? We limit our focus to 166 spatial scales within the United States as it has high spatial and temporal variability and 167 numerous observations, and since averaging over larger regions (e.g. the Northern Hemisphere, 168 or the globe) would produce a smaller calculated variability. 169 In Section 2, we describe the CAM-chem model and our simulations, as well as the

CASTNET observational database and the regional definitions used throughout this paper. In Section 3 we quantify the temporal and spatial variability of surface ozone, show how temporal and spatial averaging reduces the calculated ozone variability, and demonstrate the spatial heterogeneity of the calculated ozone variability. In Section 4, we discuss the potential strategies that could be used to maximize ozone signal detection due to meteorological variability, explore uncertainties, and make recommendations for future research.

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## 177 **2 Methods**

179 We examine both present-day (one simulation and one observed dataset) and future (two simulations) surface ozone in this study. For present-day analysis, we simulate surface ozone 180 181 using CAM-chem, a component of the Community Earth System Model (CESM) and available 182 observations within the US from the EPA CASTNET database. For future analysis, and in order 183 to examine the potential for patterns of variability to change in the future, we utilize two existing 184 simulations of CAM-chem conducted by Garcia-Menendez et al. (2017). Much of this analysis is 185 conducted using the R language (R-Project, www.r-project.org). Here we summarize each of the 186 three datasets and our approach to our analysis in Section 3.

187

#### 188 **2.1 CAM-chem**

The present-day simulation (MOZ\_2000) was conducted using CAM-chem model
version 1.2.2, with the CAM4 atmospheric component (see Tilmes et al., 2015; 2016 for model

191 description and evaluation). The model has been used extensively for a wide range of 192 atmospheric chemistry research and is included in the ACCMIP (Lamarque et al., 2012; Young 193 et al., 2012 and references therein). We conduct our simulations using the Model for Ozone and 194 Related chemical Tracers, version 4 (MOZART-4) chemical mechanism (Emmons et al., 2010), 195 which is a full tropospheric chemical mechanism integrated into CAM-Chem (e.g. Lamarque et 196 al., 2012; Tilmes et al., 2015; Brown-Steiner et al., in review). Offline forced meteorology is 197 taken from the Modern-Era Retrospective analysis for Research and Applications (MERRA) 198 reanalysis product (Rienecker et al., 2011) for 26 meteorological years (1990 – 2015). Additional 199 model evaluation and comparisons to surface and ozonesonde observations can be found in 200 Brown-Steiner et al. (in review). This simulation has 56 vertical levels - adopted from MERRA 201 meteorology – and 96 latitudinal and 144 longitudinal grid cells. We aim to isolate the variability 202 to the meteorologically-driven impact on atmospheric chemistry so we repeat year-2000 203 anthropogenic emissions from the ACCMIP (Atmospheric Chemistry and Climate Model 204 Intercomparison Project) inventory (Lamarque et al., 2012) and all non-biogenic emissions for 205 all meteorological years, and include specified long-lived stratospheric species (O<sub>3</sub>, NO<sub>x</sub>, HNO<sub>3</sub>, 206 N<sub>2</sub>O, N<sub>2</sub>O<sub>5</sub>) as in MOZART-4 (Emmons et al., 2010), an online biogenic emissions model 207 MEGAN (Guenther et al., 2012), and forced sea ice and sea surface temperatures to year 2000 208 historical conditions. Like many state-of-the-art chemical tracer models, the CAM-chem exhibits 209 some biases, most notably for our purposes a high bias in simulated surface ozone in the Eastern 210 US (e.g. Lamarque et al., 2012; Brown-Steiner et al., 2015; Travis et al., 2016; Barnes et al., 211 2016). Recent efforts have been successful in partially reducing these biases (e.g. Sun et al., 212 2017).

213 We also include two reference simulations of the future climate, MOZ 2050 and 214 MOZ 2100 (simulating the meteorological years 2035 – 2065 and 2085 – 2115, respectively) 215 using the CESM CAM-chem simulations described in detail by Garcia-Menendez et al. (2017) 216 with one set of initial condition data, and a climate sensitivity of 3.0 °C. These simulations do 217 not include projections of any changes in future emissions. Compared to the present-day 218 simulation (MOZ 2000), these future simulations (MOZ 2050 and MOZ 2100) have several 219 parametric differences: the model version is 1.1.2 (see Tilmes et al., 2015 and references for 220 information on model development), the atmospheric component is CAM3, the emissions (which 221 are held constant at year-2000 levels) are from the Precursors of Ozone and their Effects in the

- 222 Troposphere database (see Garcia-Menendez et al., 2017), and the meteorology is derived from a
- 223 linkage between the Massachusetts Institute of Technology Integrated Global System Model
- 224 (MIT IGSM) and the CESM CAM model (Monier et al., 2013), and as such has 26 vertical
- levels. For a full description of these simulations, see Garcia-Menendez et al. (2017).
- 226

#### 227 2.2 CASTNET

228 The observational database comes from the EPA Clean Air Status and Trends Network 229 (CASTNET), which has more than 90 surface observational sites within the United States and 230 has been collecting hourly surface meteorological and chemical data since 1990 (US EPA, 2016 231 and https://www.epa.gov/castnet). We collected data from all sites that reported complete ozone 232 data from each year and removed data that was marked invalid within the downloaded EPA files. 233 The number of sites that matched these criteria varied from year to year, but generally we have 234 between 55 and 94 sites throughout the 1991 – 2014 period. The CASTNET observational 235 network is located primarily in rural sites, and thus is considered to be a reasonable comparison 236 to coarse grid cell model output (e.g. Brown-Steiner et al., 2015; Phalitnonkiat et al., 2016). 237 Since a notable trend in observed ozone data exists, especially in the Northeastern US (Frost et 238 al., 2006), and since the simulations have no change in anthropogenic emissions, and thus no 239 ozone trend, we detrended the CASTNET data for each of the four averaging regions (described 240 below) using a simple linear regression.

241

# 242 **2.3 Telescoping Regional Definitions**

243 In order to isolate the impact of the size of the spatial scale over which ozone data is 244 averaged, we analyze ozone data at different spatial scales. The largest region considered is the 245 entire Continental US, while the smallest regions considered are at the individual grid cell level 246 of the CESM CAM-chem model (1.9°x2.5° latitude/longitude). Data and statistics for the other 247 regions (i.e. the Midwestern and Southeastern US) are included in the Supplemental Material, but do not alter the conclusions we draw from the Northeastern US. For CESM CAM-chem data, 248 249 we averaged all grid cells within each region, while for the CASTNET data we first average sites 250 within each corresponding CESM CAM-chem grid cell, and then averaged these data together. 251 These telescoping regions are shown in Figure 1.

#### 253 2.4 Temporal Averaging Windows

254 To explore the impact of temporal averaging, we examine ozone across a range of 255 temporal averaging windows, from 1 day up to the full 26 years for the CESM data (1990-2015), 256 the full 24 years for the detrended CASTNET data (1991 – 2014), and the 30 years available 257 from the future scenarios of Garcia-Menendez et al. (2017). Each averaging window, therefore, 258 can be considered to be a "sample" of possible realizations of meteorology. For instance, a 259 selection of an averaging window of 1 year has 26 possible slices within the 1990 – 2015 260 MOZ 2000 data, while a selection of an averaging window of 10 years has 17 possible slices 261 within the CESM data (N = # years – length of window +1). In this study, we consider all 262 realizations to be equally likely and compare them to each other and to the long-term trend. 263 However, if we were only able to simulate 5 years, we would not be able to compare to the long-264 term trend, and so be unable to completely quantify the likelihood of error in the context of the 265 long-term behavior.

266

#### 267 **3 Results**

Here we examine the spatial and temporal behavior of MOZ\_2000, MOZ\_2050, and MOZ\_2100 and compare MOZ\_2000 to present-day CASTNET observations. We introduce the moving temporal averaging windows, explore possible thresholds of acceptable error or signal strength, and examine the influence of expanding spatial averaging regions. Finally, we combine these temporal and spatial averaging techniques into a single framework.

273

## 274 **3.1 Spatial and Temporal Comparisons**

275 Figure 2 compares summertime (JJA) maximum daily 8-hour average ozone (MDA8 O<sub>3</sub>) 276 from the present-day model simulation (MOZ 2000, Figure 2a) to the year-2000 CASTNET 277 observations (Figure 2b). Figures 2c and 2d plot the MDA8 O<sub>3</sub> standard deviation and variability 278 for MOZ 2000, while Figures 2d and 2e compare the mean summertime MDA8 O3 for the 279 future simulations (MOZ 2050 and MOZ 2100). Some of the averaging strategies we present 280 can average away the high ozone behavior this MDA8 O<sub>3</sub> metric is intended to quantify, but it is 281 such a well-reported metric that focusing our analysis on it allows for ready comparisons to other 282 studies. The well-known high ozone bias in the Eastern US (e.g. Lamarque et al., 2012; Travis et 283 al., 2016; Barnes et al., 2016) is apparent, but otherwise the spatial variability over the entire

284 Continental US is well captured. While we do examine the magnitude of surface ozone in this 285 paper, most of our analysis is focused on the variability around the mean value (the anomaly), 286 and as we show below, the CASTNET observations and CESM results are largely consistent in 287 their representation of ozone variability (Figure 2, Table 1). The standard deviation of the 288 simulated MDA8  $O_3$  is large over the Eastern US and the Pacific Coast, with peak values of  $\pm 25$ 289 ppbv over the highly populated Atlantic Coast (Figure 2c). The variability (defined as the 290 standard deviation divided by the mean, expressed as a percentage) is lowest over the Western 291 US (~ 15%), only slightly higher over the Eastern US (up to 25%), and highest (up to 50%) over 292 the coastal regions (Figure 2d). We consider both the standard deviation (ppb) and a meannormalized standard deviation (as a percentage). The normalized standard deviation allows for a 293 294 more direct comparison of the shape of the MDA8 O<sub>3</sub> distributions between the simulations and 295 available observations, which accounts for the noted ozone biases (Figures 2b, c and Table 1). 296 The future climate simulations, MOZ 2050 and MOZ 2100 (Figure 2e and 2f, respectively), 297 although run with different parametric settings than MOZ 2000 (see Section 2), simulate a 298 similar spatial distribution of surface ozone, although under the warmer simulated climate of 299 2050 and 2100. These future climate simulations have a similar spatial pattern to the present-day 300 simulation (Figure 2a), with high ozone levels in the Eastern US that increases from 2050 to 301 2100 (see Garcia-Menendez et al. (2017) for more details).

302 Figure 3 compares boxplots over the four telescoping regions (Figure 1) for MOZ 2000, 303 the CASTNET data, the detrended CASTNET data, and for the single year 2000 for the 304 CASTNET data (Figures 3a-d), and Table 1 summarizes relevant statistics. In order to compare 305 CASTNET ozone to the simulated ozone, which do not have a trend over time, we detrend the 306 CASTNET data in order to remove the impact of any temporal trends (e.g. NO<sub>x</sub> emissions 307 reductions) on ozone. The Northeastern US ozone bias is apparent at the smaller spatial scales 308 (Figures 3c,d) and is less apparent when averaging over larger regions (Figures 3a,b). Figure 3e 309 compares the year-to-year boxplots of the JJA MDA8 O<sub>3</sub> for the MOZ 2000 and the detrended 310 CASTNET data, and demonstrates the variability both in the median and spread of the ozone 311 values in both the modeled and simulated data. While the MOZ 2000 ozone is generally higher 312 than the CASTNET data, there are years in which the CASTNET data has higher ozone 313 extremes. The red box plot in Figure 3e, which corresponds to the red box plot in Figure 3b,

indicates that the year 2000 was an anomalously low year for observed ozone, although not thelowest.

316 While all the CESM CAM-chem simulations have high ozone biases in the Northeastern 317 US (Figures 2 and 3, Table 1), their capability to simulate ozone variability is consistent with the 318 available observations (for present day) and for expectations of ozone variability changes in the 319 future climate (for MOZ 2050 and MOZ 2100). It is clear that variability increases when the 320 size of the averaging region decreases, a fact that is well noted in the literature, as in Hawkins 321 and Sutton (2012) for climate variables and Barnes et al. (2016) for ozone. As can be seen in in 322 Table 1, the CASTNET variability increases as the spatial scale decreases (10%, 13%, 16%, and 323 20% for our telescoping regions from continental to a single Northeastern U.S. grid box), and 324 MOZ 2000 largely captures this trend, albeit with lower overall variability (5%, 10%, 15%, and 325 15%). This increase in ozone variability with decreasing spatial scale is maintained in the future 326 climate simulations (6%, 10%, 16%, and 21% for MOZ 2050 and 7%, 12%, 17%, and 20% for 327 MOZ 2100). Table S1 contains statistics for the other telescoping regions.

328 329

#### 330 **3.2 Variability, Averaging Windows, and Thresholds**

331 As we aim to quantify the potential tradeoffs that result from a particular choice of 332 temporal and spatial scales on the assessment of ozone variability within the US, we represent 333 the spatial scale by applying the telescoping regions (see Figure 1 and Section 2.3) and we 334 represent the temporal scale through the use of moving averaging windows (see Section 2.4). We 335 frame much of the following analysis from the perspective of limited simulation length in order 336 to approximate the question that decision-makers and modelers face when constrained by limited 337 computational capabilities or available data: what is the likelihood that a particular estimate (of 338 both the mean and the variability) is not a true representation of the true mean and variability, but 339 rather a product of the underlying variability at the particular choice of spatial and temporal 340 scale?

Figure 4 presents this likelihood by plotting all possible estimates of MDA8 O<sub>3</sub> (as anomalies from the long-term mean) over all possible selections of averaging window (from 1 day up to the complete time series) for our telescoping regions. The semi-cyclical and highly auto-correlated nature of surface ozone is apparent at all spatial scales, with alternating cycles of anomalously high and low ozone. The temporal impact of anomalous ozone events is indicated

346 by the vertical and right-leaning diagonal striations, which show that anomalous ozone events 347 can impact estimates of ozone values within averaging windows up to 15 or 20 years. Figure 4 348 demonstrates how small-scale anomalously high or low ozone values (that come only from 349 meteorological variability) can impact temporal averages of 5, 10, or even 20 years. For instance, 350 a selected 5-year averaging window within the MOZ 2000 simulation averaged over the 351 Northeastern US could be 2.5 ppbv higher or lower than the 25-year mean value of 74 ppbv, a 352 potential error of 7%. Horizontal lines in Figure 4 mark the length of averaging windows that are 353 needed to ensure that ozone anomaly for any selection of averaging window does not exceed a 354 given threshold (5, 1, and 0.5 ppbv for solid, dashed, and dotted lines respectively). This 355 potential error is larger within smaller regions and at the shorter selections of the averaging 356 window. While the high and low ozone anomalies differ in time between CASTNET, MOZ 2000, MOZ 2050, and MOZ 2100 in Figure 4, the impact of spatial and temporal 357

358 averaging is consistent.

359 We also quantify this variability in Supplemental Figures S1 and S2, which plots the 360 likelihood (as a percentage) that a particular selection of spatial (rows) and temporal (x-axis) 361 scale estimates ozone values that exceed a particular threshold (colored lines) away from the true 362 mean value. For instance, if we are interested in characterizing ozone behavior (e.g. estimating a 363 trend, or the mean value) in the Northeastern US, but were limited to a 5-year simulation, there is 364 more than a 50% likelihood that the simulated ozone is 1 ppbv away from the 26-year mean, and 365 an 80% likelihood that the discrepancy is greater than 0.5 ppby. However, these data indicate 366 that there is a virtual certainty that the estimate will be within 2.5 ppbv of the true mean value. 367 We should note that, at the grid-cell level and within a 10-year period, the surface ozone 368 variability can exceed 1 ppbv but is unlikely to exceed 2.5 ppbv (Figure 4), and that a 20-year 369 trend is very likely to be able to identify significant ozone signals among the impact of 370 meteorological variability on atmospheric chemistry. Our results also align with the results from 371 Garcia-Menendez et al. (2017), which recommended that simulations need to be at least 15 years 372 long to identify anthropogenically-forced ozone signals on the order of 1 ppby.

Figures 4 and Supplemental Figures S1 and S2 compare the CASTNET observations to the three CESM CAM-chem simulations, and while there are minor differences, there are broad features that are consistent. First, using longer temporal averaging windows reduces the influence of small-scale ozone variability at all spatial scales, and depending on the acceptable

threshold, one can select a temporal scale that effectively reduces the likelihood of exceeding

that threshold to zero. Second, larger spatial scales also reduce this likelihood of exceeding a

379 given threshold, but not as effectively as longer temporal scales. Finally, the impact of both

380 temporal and spatial averaging on ozone variability is largely consistent for the CASTNET

381 observations and for all three CESM CAM-chem simulations.

382

### **383 3.3 Selection of Temporal Averaging Scales**

384 Figure 5 extends this analysis to examine the spatial heterogeneity of this likelihood of 385 the meteorological variability causing ozone anomalies exceeding particular thresholds at the 386 grid cell level. Here we plot four thresholds (0.5, 1, 2.5, and 5 ppbv) and four averaging windows 387 (1, 5, 10, and 20 years) for the MOZ 2000 simulation. Ozone variability is highest in the Eastern 388 US. At the grid-cell level, there are two strategies for filtering out the noise associated with 389 natural meteorological variability (and thus enhancing signal detection capabilities): either 390 average over longer periods, or acknowledge the level of noise and increase the threshold. For 391 these data, it is virtually certain that any 20-year average will be within 5 ppbv of a full 25-year 392 mean value (which itself may not be an accurate representation of a longer simulation), and 393 virtually certain that any 1-year average will be at least 0.5 ppbv away from the mean.

394 Supplemental Figure S3 extends the analysis of Figure 5 by comparing the MOZ 2000, 395 MOZ 2050, and MOZ 2100 simulations across the four thresholds for the 5-year averaging 396 window. Figure 6 similarly compares the 1 ppbv ozone threshold across the four averaging 397 windows for MOZ 2000, MOZ 2050, and MOZ 2100. Interpreting Figures 6 and Supplemental 398 Figure S3 give largely consistent interpretations than the analysis above (Figure 5). Namely, that 399 at the grid-scale level, increasing the temporal averaging window (Figure 6) or increasing the 400 acceptable ozone threshold (Supplemental Figure S3) are effective at reducing the impact of the 401 meteorological variability on estimates of the ozone signal. Shorter windows (or smaller 402 thresholds) are needed in the Western US (where variability is smaller, see Figure 2d) than in the 403 Eastern US (where variability is larger) as well as over coastal and highly populated regions. 404 Finally, the 1 ppbv threshold and the 5-year averaging window plots (in either Figure 5 or 405 Supplemental Figure S3) indicate that the spatial distribution and location of the peak variability 406 may shift into the future, although this may be due to parametric differences between

407 MOZ\_2000, MOZ\_2050, and MOZ\_2100. Future simulations will be needed to check this shift
408 in peak ozone variability.

409

## 410 **3.4 Selection of Spatial Averaging Scales**

411 We examine the impact of increasing the spatial averaging region (Figure 7) at four 412 different temporal averaging windows (1, 5, 10, and 20 years) and for the smallest ozone 413 threshold from the previous section (0.5 ppbv). It is evident that at all temporal averaging 414 windows, expanding the number of surrounding grid cells that are averaged together consistently 415 decreases the likelihood of exceeding the 0.5 ppbv threshold, although these reductions are 416 relatively small at the 1-year window, especially over the Eastern U.S. While increasing the 417 spatial averaging from a single grid-cell up to include the surrounding 81 grid cells (bottom row 418 in Figure 7) manages to essentially smooth away much of the spatial heterogeneity in surface 419 ozone (by moving down any column in Figure 7), it does not eliminate the likelihood of 420 exceeding the 0.5 ppbv threshold over much of the Eastern U.S. For instance, even at a 20-year 421 averaging window, and by averaging together the surrounding 81 grid-cells over locations in the 422 Eastern U.S., there is still a 20-70% likelihood of exceeding the 0.5 ppbv threshold due to the 423 small-scale impact of the meteorological variability on atmospheric chemistry.

424

# 425 **3.5** Combination of Spatial and Averaging Scales

426 We now examine the combined impact of temporal and spatial averaging on reducing the 427 influence of small-scale ozone variability in order to enhance ozone signal detection capabilities. 428 Table S2 summarizes our analysis by dividing the likelihood of the ozone variability estimates 429 exceeding selected thresholds away from the long-term mean into four categories: (1) the length 430 of the averaging window over which ozone is averaged (columns); (2) the magnitude of the 431 ozone threshold of interest (rows); (3) the observed (CASTNET) and modeled (MOZ 2000, 432 MOZ 2050, and MOZ 2100) ozone data (sub-columns); and (4) the size of the spatial extent 433 over which ozone is averaged (sub-rows). A graphical representation consistent with the data 434 presented in Table S2 is plotted in Figure 8 for the Continental US average and for three grid 435 cells that represent various cases. In each plot in Figure 8, by moving along columns from left to 436 right, we can see the influence of increasing the size of the temporal averaging window, and by 437 moving along rows (from the bottom to the top), we can see the influence of increasing the

spatial averaging scale. By taking in the entire plot as a whole, we can get a feel for the
combined influence of both temporal and spatial averaging. Supplemental Figure S4 contains a
plot for each grid cell in the Continental US.

441 On average within the Continental US, both temporal and spatial averaging are effective 442 at reducing the calculated MDA8 O<sub>3</sub> anomaly, although temporal averaging is more effective 443 (Figure 8a). There are many grid cells in the Eastern and Western US coasts (Figure 8b, 444 Supplemental Figure S4), where both spatial and temporal averaging are effective, but their 445 combined usage is especially effective. There are also many grid cells where temporal averaging 446 is effective, but spatial averaging is barely effective, or not effective at all (Figure 8c and 447 Supplemental Figure S4). Finally, there are some grid cells, particularly in the Central US 448 (Figure 8d and Supplemental Figure S4), where spatial averaging over smaller regions is 449 effective, but spatial averaging of larger regions actually increases the calculated MDA8 O<sub>3</sub> 450 anomaly by including surrounding grid cells that have higher variability.

451

## 452 **4 Discussion**

453 We now return to the original four research questions posed in Section 1. First, what is 454 the magnitude of ozone variability due to meteorology alone at the smallest scale, and what is the 455 impact of increasing the scale of temporal and spatial averaging? In both observed and modeled 456 MDA8 O<sub>3</sub> surface data, the small-scale variability driven solely by the meteorological variability 457 impact on atmospheric chemistry (expressed as the standard deviation as a percentage of the 458 mean) can exceed 20% (Table 1, Figure 2d). The chemical variability examined here is the result 459 of fluctuations in meteorology, which itself results from larger-scale climatological drivers. 460 While variability in emissions also influences atmospheric chemistry, our analysis has removed 461 the influence of emissions variability and isolated the variability due to meteorology. A more 462 comprehensive analysis of chemical variability will need to account for both meteorological and 463 emission variability, which is complicated by temporal trends in both the emissions of ozone 464 precursor species and the climate.

There is high temporal and spatial heterogeneity of surface ozone variability (Figure 2d), with the lowest values found in the Western US (< 10%), higher values found in the Eastern US (up to 20%), and the highest values over coastal or heavily populated regions (up to 30%). Averaging over longer temporal scales (by increasing the averaging window) and over larger

469 spatial scales (by expanding the averaging region) can reduce the magnitude of the calculated 470 variability, with temporal averaging proving to be more effective than spatial averaging in most 471 cases (Figure 8). In this study, we performed simple spatial averaging, but there are other 472 methodologies for smoothing two-dimensional signals (e.g. Räisänen et al., 2011; Pogson and 473 Smith, 2015) that could potentially increase signal detection capabilities.

474 Second, are there combinations of temporal and spatial averaging that maximize the 475 filtration of calculated ozone variability, and thus maximize the potential for signal detection? 476 Figure 8 (and Supplemental Figure S4) demonstrate clearly that there are cases in which the 477 combined usage of temporal and spatial averaging can reduce the calculated variability better 478 than either strategy alone (see Figure 8b), although there are many regions within the Eastern US 479 in which spatial averaging has little to no impact on reducing the calculated variability (Figure 480 8c) or even results in an increase in the calculated variability (Figure 8d). There are no such 481 cases (see Supplemental Figure S4) in which expanding the temporal averaging scale increases 482 the calculated ozone variability. This could potentially enable region-specific averaging 483 strategies that help decision-makers identify and meet regional air quality objectives.

484 Third, are these results dependent on the particular parameterizations of the CESM 485 CAM-chem model, and are they consistent with the available CASTNET observations? The 486 three CESM CAM-chem simulations exhibited consistent representations of ozone variability, 487 consistent with our understanding of future changes to the climate (and meteorology) and the 488 resulting impact on atmospheric chemistry (Table 1, Figures 4, S1, and S2). Compared to the 489 CASTNET observations (which we detrended to remove the influence of changing precursor 490 emissions), the present-day simulation (MOZ 2000) exhibited a high ozone bias in the Eastern 491 US, while the representation of the ozone variability is comparable (Table 1).

492 Fourth, how may these strategies be applied to other datasets, be they chemical, 493 meteorological, or climatological? Much of this analysis could be applied to any dataset that has 494 spatial and temporal coverage, as long as some set of acceptable thresholds is provided. While 495 our time step in this analysis is daily (given the MDA8  $O_3$  metric), and applied only to 496 summertime (JJA) days, any time step (i.e. hourly, monthly, annual, decadal) could be utilized as 497 long as cyclical trends (e.g. diurnal or seasonal cycles) are removed. Indeed, the sliding-scale 498 presentation in Figure 8 and Supplemental Figure S4 can specifically be utilized to identify 499 particular spatial and temporal scales that are sufficient to identify signals at particular thresholds

500 and to identify particular geographic regions that are best suited to identify a given signal. For 501 example, Sofen et al. (2016) identified regions across the globe where additional observations 502 would be particularly suited to improve our understanding of surface ozone behavior, and our 503 analysis could potentially be used to identify particular temporal and spatial averaging scales that 504 could further maximize the capability for trend detection. In particular, Sofen et al. (2016) noted 505 that the peak in the power spectrum of the El Niño-Southern Oscillation (ENSO) on surface 506 ozone is at the 3.8 year time scale, and that within some regions within the US, the amplitude of 507 the ENSO influence on surface ozone approached 0.5 ppbv (and up to 1.1 ppbv globally). Our 508 analysis shows that there are no grid cells within the Continental US where a 0.5 ppbv signal can 509 be identified at the 5-year (or shorter) temporal averaging scale (Supplemental Figure S4), but 510 that there are many regions – especially within the Western US – in which even a modest amount 511 of spatial averaging can identify surface ozone signals below the 1 ppbv level with a 5-year or 512 shorter averaging window. The type of sliding-scale analysis – in which spatial and temporal 513 averaging are utilized individually and in combination – as presented in Figure 8 and 514 Supplemental Figure S4 could readily be applied to a wide range of atmospheric (and other) 515 topics to aid in the capability to identify signals that exist both in space and in time. In particular, 516 low-frequency oscillations (e.g. ENSO, and others) and other forms of internally or externally 517 forced trends (e.g. anthropogenic and natural changes in emissions) are readily adaptable to this 518 type of analysis, which could address signals pertaining to precipitation, biogenic emissions, 519 boundary layer variables, cloud properties, and many others.

520 We did not quantify statistical significance (as in Lewandowski et al., 2015) as our goals 521 were to understand the general nature of ozone variability at all scales and for all signal 522 strengths. Statistical significance testing (and other statistical techniques) can certainly provide 523 additional information as to the strengths of ozone signals within the underlying variability, and 524 can be used to extend these results in a case-by-case manner, but we leave this testing to future 525 studies that can focus on particular air quality objectives at particular temporal and spatial scales. 526 Furthermore, future research examining the impact of spatial and temporal averaging using 527 regional-scale models, models with different resolutions, and the inclusion of urban observations 528 could provide additional insight into understanding chemical variability and averaging 529 techniques.

530 Smaller signals require longer temporal averaging periods to identify. Figure 4 shows that 531 a 0.5 ppb MDA8 O<sub>3</sub> signal will emerge after 15 - 20 years of temporal averaging. The range here 532 reflects different spatial averaging domains, with larger domains requiring shorter temporal 533 averaging windows than smaller domains (i.e. 15 years for averaging over the Continental US 534 and 20 years for averaging over the Northeastern US). This would mean that an average trend of 535 0.25 - 0.33 ppb/year would require a time series of at least 15 years to identify. Similarly, a 1.0 536 ppb MDA8 O<sub>3</sub> signal emerges after 7 - 15 years, which indicates an average trend of 0.14 -0.67 537 ppb/year would take at least 7 years to identify. Finally, a 5 ppb signal can be identified in less 538 than 3 years, which indicates that an average trend of 1.67 ppb/year or greater would only require 539 a 3-year time series. This presents particular difficulties if the ozone signal of interest is a trend 540 spanning a time period on the same order. The 10 - 15 year averaging time scale we propose 541 translates into a length of time beyond which you are likely to not see spurious trends above 0.5 542 ppb, but there are many cases in which the identification of a small trend is desired with less than 543 10 - 15 years of available data. For instance, Jiang et al. (2018) have found that NO<sub>x</sub> emissions 544 reductions since 2005 are not as strong as previously expected, showing a significant slowdown 545 beginning in 2011. This has large implications for ozone and for short-term decisions for air 546 quality managers within the United States, who have to promulgate policies on short-term scales 547 without the luxury of postponing action until longer and more complete data sets become 548 available. As we have shown, spatial and temporal variability due to meteorology is high, and the 549 identification and quantification of trends over 5, 10, or 15 years is difficult, particularly at small 550 spatial scales.

551 However, as we have shown, a consideration of the impact on variability – and how 552 variability changes over time – is often pivotal to understand the nature of the signals being 553 examined. In this paper, we have provided methods for quantifying the spatial and temporal 554 variability and strategies for determining which types of signals are likely detectable at particular 555 temporal and spatial scales. Some signals, especially small signals at small scales, are simply not 556 large enough to emerge from the variability, and thus may not be detectable without additional 557 data or expanding the temporal and spatial averaging scales used for analysis. Quantifying the 558 signal-to-noise ratio at a variety of spatial scales, and determining an acceptable threshold of a 559 particular signal, could be one accessible method for providing this context. The risk in 560 neglecting the quantification and contextualization of the magnitude of the ozone signal relative

to the magnitude of the variability induced by the internal meteorology – and the impact of temporal and spatial averaging – is primarily the risk of drawing conclusions that are more sensitive to a particular peculiarity in the underlying variability rather than the signal itself.

564

## 565 **5 Conclusions**

566 We quantified the impact of spatial and temporal averaging at different scales – both 567 individually and combined – on estimates of summertime surface ozone variability and the 568 resulting likelihood of over-confidence in estimates of chemical signals over the United States 569 using CASTNET observations and the CESM CAM-chem model. We simulate three multi-570 decadal time periods, each with constant surface emissions, and find that this analysis is 571 consistent across our simulated time periods, and that our results are not sensitive to particular 572 configurations and parametric choices within the CESM CAM-chem (i.e. emissions, 573 meteorology, and climate). We also provide a conceptual framework for gaining understanding 574 of the influence of spatial and temporal averaging that may be adapted to a wide range of 575 atmospheric and surface phenomena, provided sufficient spatial and temporal coverage. Here we 576 focus on summertime surface ozone, a highly variable (in both space and time) atmospheric 577 constituent with severe human health impacts and implications for planetary climate, which is 578 the focus of many local, regional, and national policies. However, these ozone signals (e.g. 579 temporal trends or regional averages) are frequently small when compared to the magnitude of 580 the day-to-day ozone variability, and thus detecting these signals can be challenging. In 581 particular, it would be impractical to delay interpreting observations for 10 - 15 years, or 582 alternatively to expand the spatial averaging such that small-scale features are smoothed away. 583 Nonetheless, it is unwise to over-interpret trends and signals based on observations from a 584 limited spatial area and over a short temporal period. Our analysis and conceptual framework 585 presented here cannot solve this tension, but it does demonstrate some strategies which can allow 586 for a selection of spatial and temporal averaging scales, and a consideration of the error 587 threshold, that can aid in this signal detection on a case-by-case basis. Taking into account the 588 complex interactions involving trends and variability between emissions, chemistry, 589 meteorology, and climatology necessitates a variety of strategies. This work quantifies the 590 impact of spatial and temporal averaging in signal detection, which can be used in conjunction

with ensembles of simulations, statistical techniques, and other strategies to further ourunderstanding of the chemical variability in our atmosphere.

593 In order to quantify the impact of spatial and temporal averaging on summertime ozone 594 variability, we start by selecting four telescoping spatial regions (the Continental US, the Eastern 595 US, the Northeastern US, and a single grid cell within the Northeastern US) and examine all 596 possible choices for averaging windows (ranging from daily to multi-decadal windows), 597 although we focused primarily on averaging windows of 1, 5, 10, and 20 years. We find that -598 consistent with previous studies – summertime MDA8 O<sub>3</sub> variability is largest at the smallest 599 spatial and temporal scales, and is frequently on the order of  $\pm 10 - 20$  ppbv, or which is roughly 600 15-20% of the mean ozone signal. In order to minimize the chemical noise that results from 601 meteorological variability – and thus enhance the signal – we find averaging windows of 10-15 602 years (and sometimes longer at the smaller spatial scales) combined with modest (nearest-603 neighbor) spatial averaging substantially improve the capability for signal detection. For signals 604 that are large compared to the underlying meteorological variability (e.g. strong emissions 605 reductions), shorter averaging windows and smaller spatial regions may be used. We recognize 606 that achieving a 10 - 15 year temporal averaging window is difficult, but this recommendation is 607 consistent with recent literature (e.g. Barnes et al., 2015; Garcia-Menendez et al., 2017). For 608 studies where 10 - 15 years of averaging is impractical, we recommend that some spatial and 609 temporal context is provided that demonstrates that the signals being examined are robust and 610 not the result of internal variability or noise. We also recognize that our analysis is just one 611 strategy for enhancing signal detection capabilities, and will ideally be used alongside others, 612 such as perturbed initial condition ensembles, running simulations with either internal or forced 613 meteorology, and examining a region or time period with different models or parameterizations. 614 We show that the largest summertime ozone variability is found in the Eastern US (Figure 5, 615 Figure S4), and subsequently there are many regions within the Eastern US where even a 20-year 616 averaging window has a non-negligible likelihood of estimating ozone variability that is 617 dependent (with possible error in the 1-3 ppby range) on the particular years selected. In 618 addition, over much of the Eastern US, simulations of 5-years or shorter have a substantial 619 likelihood (40 - 90%), Figures S1 and S2) of reflecting the influence of meteorological variability 620 on chemistry rather than the mean state of surface ozone, with the possibility of 5 - 10 ppbv 621 error (Figure S4). While we have detrended the CASTNET observations to compare to the

constant year-2000 cycled emissions in the simulations, the CASTNET time series inherently
includes the compounded variability of both meteorological and emission sources. Future studies
will need to expand this analysis to include trends and variability in the emissions, as well as in
the meteorology.

626 Finally, we demonstrate a conceptual framework that allows for a "sliding-scale" view of 627 surface ozone variability, in which both temporal and spatial averaging is examined at every grid 628 cell within the Continental US. We show that the magnitude of estimates of ozone variability can 629 be reduced with both temporal and spatial averaging, although temporal averaging tends to be 630 more effective. While there are many regions in which both temporal and spatial averaging used 631 in conjunction substantially reduce the estimate of ozone variability, there are some regions 632 where spatial averaging is ineffective, or even counter-effective. In contrast, this is not the case 633 for temporal averaging, which consistently reduces the magnitude of estimated ozone variability. 634 Our analysis could be combined with other studies (e.g. Sofen et al., 2016) to guide 635 observational and modeling strategies and identify regions and scales at which particular signals 636 are most likely to be identified.

# 638 Code Availability

- 639 CESM CAM-Chem code is available through the National Center for Atmospheric Research /
- 640 University Corporation for Atmospheric Research (NCAR/UCAR) website
- 641 (<u>http://www.cesm.ucar.edu/models/cesm1.2/</u>), and this project made no code modifications from
- 642 the released model version.

# 643 Data Availability

- 644 The raw model output is archived on the NCAR servers, and processed data is archived at
- 645 https://dspace.mit.edu/handle/1721.1/114467.

646 Supplemental Link

# 648 Author Contribution

- 649 BBS ran the present-day simulation, analyzed the data, and wrote the manuscript. EM ran the
- 650 future climate simulations, while FGM ran the future atmospheric chemistry simulations and
- 651 made the data available to BBS. NS, RP, EM, ST, and LE guided and reviewed the scientific
- 652 modeling and analysis process. All authors provided feedback throughout the project and
- 653 development of the manuscript.

# **Competing Interests**

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

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#### 669 **References**

- Angélil, O., Stone, D., Perkins-Kirkpatrick, S., Alexander, L.V., Wehner, M., Shiogama, H.,
  Wolski, P., Ciavarella, A., and Christidis, N.: On the nonlinearity of spatial scales in
  extreme weather attribution statements, Clim. Dyn., 2017.
- Barnes, E. A., Fiore, A. M., and Horowitz, L. W.: Detection of trends in surface ozone in the
  presence of climate variability, J. Geophys. Res. Atmos., 121, 6112–6129, 2016.
- Brown-Steiner, B., Hess, P. G., and Lin, M. Y.: On the capabilities and limitations of GCCM
  simulations of summertime regional air quality: A diagnostic analysis of ozone and
  temperature simulations in the US using CESM CAM-chem, Atmos. Environ., 101, 134–
  148, 2015.
- Brown-Steiner, B., Selin, N. E., Prinn, R., Tilmes, S., Emmons, L., Lamarque, J.-F., and
  Cameron-Smith, P.: Evaluating Simplified Chemical Mechanisms within CESM Version
  1.2 CAM-chem (CAM4): MOZART-4 vs. Reduced Hydrocarbon vs. Super-Fast Chemistry,
  Geosci. Model Dev. Discuss., in review, 2018.
- 683 Camalier, L., Cox, W., and Dolwick, P.: The effects of meteorology on ozone in urban areaas
  684 and their use in assessing ozone trends, Atmos. Environ., 41, 7127-7137, 2007.
- Cooper, O. R., Gao, R. S., Tarasick, D., Leblanc, T., and Sweeney, C.: Long-term ozone trends
  at rural ozone monitoring sites across the United States, 1990-2010, J. Geophys. Res., 117,
  D22307, 2012.
- de Elía, R., Biner, S., and Frigon, A.: Interannual variability and expected regional climate
  change over North America, Clim. Dyn., 41, 1245, 2013.
- 690 Deser, C., Phillips, A., Bourdette, V., and Teng, H.: Uncertainty in climate change projections:
  691 the role of internal variability, Clim. Dyn., 38, 527, 2012.
- Diem, J. E., and Comrie, A. C.: Predictive mapping of air pollution involving sparse spatial
   observations, Environmental Pollution, 119, 1, 99–117, 2002.
- Emmons, L. K., Walters, S., Hess, P. G., Lamarque, J.-F., Pfister, G. G., Fillmore, D., Granier,
  C., Guenther, A., Kinnison, D., Laepple, T., Orlando, J., Tie, X., Tyndall, G., Wiedinmyer,
  C., Baughcum, S. L., and Kloster, S.: Description and evaluation of the Model for Ozone
  and Related chemical Tracers, version 4 (MOZART-4), Geosci. Model Dev., 3, 43-67,
  2010.
- Fiore, A. M., Oberman, J. T., Lin, M. Y., Zhang, L., Clifton, O. E., Jacob, D. J., Naik, V.,
  Horowitz, L. W., Pinto, J. P., and Milly, G. P.: Estimating North American background
  ozone in U.S. surface air with two independent global models: Variability, uncertainties,
  and recommendations, Atmos. Environ., 96, 284–300, 2014.
- Fiore, A. M., Jacob, D. J., Liu, H., Yantosca, R. M., Fairlie, T. D., and Li, Q.: Variability in
  surface ozone background over the United States: Implications for air quality policy, J.
  Geophys. Res. Atmos., 108, D24, 1787, 2003.
- Garcia-Menendez, F., Saari, R. K., Monier, E., and Selin, N. E.: U.S. Air Quality and Health
   Benefits from Avoided Climate Change under Greenhouse Gas Mitigation, Environ. Sci.
   Technol., 49, 7580–7588, 2015.

- Garcia-Menendez, F., Monier, E., and Selin, N. E.: The role of natural variability in projections
  of climate change impacts on U.S. ozone pollution, Geophys. Res. Lett., 44, 2911–2921,
  2017.
- Giorgi, F., and Bi, X.: Time of emergence (TOE) of GHG-forced precipitation change hot-spots,
   Geophys. Res. Lett., 36, L06709, 2009.
- Guenther, A. B., Jiang, X., Heald, C. L., Sakulyanontvittaya, T., Duhl, T., Emmons, L. K., and
  Wang, X.: The Model of Emissions of Gases and Aerosols from Nature version 2.1
  (MEGAN2.1): An extended and updated framework for modeling biogenic emissions,
  Geosci. Model Dev., 5, 1471–1492, 2012.
- Hawkins, E., and Sutton, R.: Time of emergence of climate signals, Geophys. Res. Lett., 39,
   L01702, 2012.
- Jacob, D. J., and Winner, D. A.: Effect of climate change on air quality, Atmos. Environ, 43, 51–
   63, 2009.
- Jiang, Z., McDonald, B. C., Worden, H., Worden, J. R., Miyazaki, K., Qu, Z., Henze, D. K.,
  Jones, D. B. A., Arellano, A. F., Fischer, E. V., Zhu, K., and Boersma, F.: (2018).
  Unexpected slowdown of US pollutant emission reduction in the past decade. Proceedings
  of the National Academy of Sciences, 201801191.
- Jhun, I., Coull, B. A., Schwartz, J., Hubbell, B., and Koutrakis, P.: The impact of weather
  changes on air quality and health in the United States in 1994–2012, Environ. Res. Lett., 10,
  084009, 2015.
- Kay, J. E., Deser, C., Phillips, A., Mai, A., Hannay, C., Strand, G., Arblaster, J. M., Bates, S. C.,
  Danabasoglu, G., Edwards, J., Holland, M., Kushner, P., Lamarque, J.-F., Lawrence, D.,
  Lindsay, K., Middleton, A., Munoz, E., Neale, R., Oleson, K., Polvani, L., and Vertenstein,
  M.: The Community Earth System Model (CESM) large ensemble project: A community
  resource for studying climate change in the presence of internal climate variability, Bull.
  Amer. Meteor. Soc., 96, 1333-1349, 2015.
- King, A. D., Donat, M. G., Fischer, E. M., Hawkins, E., Alexander, L. V, Karoly, D. J., Dittus,
  A. J., Lweis, S. C., and Perkins, S. E.: The timing of anthropogenic emergence in simulated
  climate extremes, Environ. Res. Lett., 10, 094015, 2015.
- Knote, C., Tuccella, P., Curci, G., Emmons, L., Orlando, J. J., Madronich, S., Baró, R., JoménezGuerrero, P., Luecken, D., Hogrefe, C., Forkel, R., Werhahn, J., Hirtl, M., Pérez, J. L., José,
  R. S., Giordano, L., Brunner, D., Yahya, K., and Zhang, Y.: Influence of the choice of gasphase mechanism on predictions of key gaseous pollutants during the AQMEII phase-2
  intercomparison, Atmos. Environ., 115, 553–568, 2015.
- 743 Knutti, R.: The end of model democracy?, Clim. Change, 102, 395–404, 2010.
- Lamarque, J.-F., Bond, T. C., Eyring, V., Granier, C., Heil, A., Klimont, Z., Lee, D., Liousse, C.,
  Mieville, A., Owen, B., Schultz, M. G., Shindell, D., Smith, S. J., Stehfest, E., Van
- Aardenne, J., Cooper, O. R., Kainuma, M., Mahowald, N., McConnell, J. R., Naik, V.,
- Riahi, K., and Van Vuuren, D. P.: Historical (1850-2000) gridded anthropogenic and
- 747 Riam, K., and Van Vullen, D. F., Instonear (1830-2000) gridded antihopogenic and 748 biomass burning emissions of reactive gases and aerosols: Methodology and application,
- Atmos. Chem. Phys., 10, 7017–7039, 2010.

- Lamarque, J.-F., Dentener, F., McConnell, J., Ro, C.-U., Shaw, M., Vet, R., Bergmann, D.,
- 751 Cameron-Smith, P., Dalsoren, S., Doherty, R., Faluvegi, G., Ghan, S. J., Josse, B.,
- MacKenzie, I. A., Plummer, D., Shindell, D. T., Skeie, R. B., Stevenson, D. S., Strode, S.,
  Zeng, G., Curran, M., Dahl-Jensen, D., Das, S., Fritzsche, D., and Nolan, M.: Multi-model
  mean nitrogen and sulfur deposition from the atmospheric chemistry and climate model
  intercomparison project (ACCMIP): Evaluation of historical and projected future changes,
- 756 Atmos. Chem. Phys., 13, 7997–8018, 2013.
- Lamarque, J.-F., Emmons, L. K., Hess, P. G., Kinnison, D. E., Tilmes, S., Vitt, F., Heald, C. L.,
  Holland, E. A., Lauritzen, P. H., Neu, J., Orlando, J. J., Rasch, P. J., and Tyndall, G. K.:
  CAM-chem: Description and evaluation of interactive atmospheric chemistry in the
- Community Earth System Model, Geosci. Model Dev., 5, 369–411, 2012.
- Lawrence, M. G., Hov, Ø., Beekmann, M., Brandt, J., Elbern, H., Eskes, H., Feichter, H., and
   Takigawa, M.: The chemical weather, Environ. Chem, 2, 6–8, 2005.
- Lewandowsky, S., Risbey, J. S., and Oreskes, N.: On the definition and identifiability of the
   alleged "hiatus" in global warming, Sci. Rep., 5, 16784, 2015.
- Lin, M., Horowitz, L. W., Oltmans, S. J., Fiore, A. M., and Fan, S.: Tropospheric ozone trends at
   Mauna Loa Observatory tied to decadal climate variability, Nat. Geosci., 7, 136–143, 2014.
- McComiskey, A., and Feingold, G.: The scale problem in quantifying aerosol indirect effects,
   Atmos. Chem. Phys., 12, 1031–1049, 2012.
- Medhaug, I., Stolpe, M. B., Fischer, E. M., and Knutti, R.: Reconciling controversies about the
  'global warming hiatus,' Nature, 545, 41–47, 2017.
- Möller, D. Chemistry of the Climate System, pp. 331-334, Walter de Gruyter GmbH and Co.,
   KG, Berlin/New York, 2010.
- Monier, E., Scott, J. R., Sokolov, A. P., Forest, C. E., and Schlosser, C. A.: An integrated
  assessment modeling framework for uncertainty studies in global and regional climate
  change: The MIT IGSM-CAM (version 1.0), Geosci. Mod. Dev., 6, 2063–2085, 2013.
- Monier, E., Gao, X., Scott, J. R., Sokolov, A. P., and Schlosser, C. A.: A framework for
   modeling uncertainty in regional climate change, Clim. Change, 131, 51–66, 2015.
- Monier, E., Xu, L., and Snyder, R.: Uncertainty in future agro-climate projections in the United
   States and benefits of greenhouse gas mitigation, Environ. Res. Lett., 11, 055001, 2016.
- Paltsev, S., Reilly, J. M., Jacoby, H. D., Eckaus, R. S., McFarland, J. R., Sarofim, M. C.,
  Asadoorian, M. O., and Babiker, M. H.: The MIT emissions prediction and policy analysis
  (EPPA) model: Version 4, Rep. 125, MIT Joint Program on the Sci. and Policy of Global
  Change, 2005.
- Pogson, M., and Smith, P.: Effect of spatial data resolution on uncertainty, Environ. Model.
   Softw., 63, 87–96, 2015.
- Pyle, J. A., and Zavody, A. M.: The modelling problems associated with spatial averaging, Q. J.
  R. Meteorol. Soc., 116, 753–766, 1990.
- Räisänen, J., and Ylhäisi, J. S.: How much should climate model output be smoothed in space?,
  J. Climate, 24, 867–880, 2011.

- Rasmussen, D. J., Fiore, A. M., Naik, V., Horowitz, L. W., McGinnis, S. J., and Schultz, M. G.:
  Surface ozone-temperature relationships in the eastern US: A monthly climatology for
  evaluating chemistry-climate models, Atmos. Environ., 47, 142–153, 2012.
- Rienecker, M. M., Suarez, M. J., Gelaro, R., Todling, R., Bacmeister, J., Liu, R., Bosilovich, M.
  G., Schubert, S. D., Takacs, L., Kim, G-K, Bloom, S., Chen, J., Collins, D., Conaty, A., da
  Silva, A., Gu, W., Joiner, J., Koster, R. D., Lucchesi, R., Molod, A., Owens, T., Pawson, S.,
  Pegion, P., Redder, C. R., Reichle, R., Robertson, F. R., Ruddick, A. G., Sienkiewicz, M.,
  and Woollen, J.: MERRA: NASA's Modern-Era Retrospective analysis for Research and
  Applications, J. Climate, 24, 3624–3648, 2011.
- Roberts, C. D., Palmer, M. D., McNeall, D., and Collins, M.: Quantifying the likelihood of a
   continued hiatus in global warming, Nat. Clim. Change, 5, 337–342, 2015.
- Schnell, J. L., and Prather, M. J.: Co-occurrence of extremes in surface ozone, particulate matter,
  and temperature over eastern North America, Proc. Natl. Acad. Sci. U.S.A., 114, 11, 28542859, 2017.
- Schurer, A. P., Hegerl, G. C., Mann, M. E., Tett, S. F. B., and Phipps, S. J.: Separating forced
   from chaotic climate variability over the past millennium, J. Climate, 26, 6954–6973, 2013.
- Searle, K. R., Chipperfield, M. P., Bekki, S., and Pyle, J. A.: The impact of spatial averaging on
  calculated polar ozone loss: 2. Theoretical analysis, J. Geophys. Res, 103, D19, 25409–
  25416, 1998.
- Sofen, E. D., Bowdalo, D., and Evans, M. J.: How to most effectively expand the global surface
  ozone observing network, Atmos. Chem. Phys., 16, 1445–1457, 2016.
- Stevenson, D. S., Young, P. J., Naik, V., Lamarque, J. F., Shindell, D. T., Voulgarakis, A., Skeie,
  R. B., Dalsøren, S. B., Myhre, G., Berntsen, T. K., Folberth, G. A., Rumbold, S. T., Collins,
  W. J., MacKenzie, I. A., Doherty, R. M., Zeng, G., van Noije, T. P. C., Strunk, A.,
- 814 Bergmann, D., Cameron-Smith, P., Plummer, D. A., Strode, S. A., Horowitz, L., Lee, Y. H.,
- Szopa, S., Sudo, K., Nagashima, T., Josse, B., Cionni, I., Righi, M., Eyring, V., Conley, A.,
- Bowman, K. W., and Wild, O.: Tropospheric ozone changes, radiative forcing and
   attribution to emissions in the Atmospheric Chemistry and Climate Model Intercomparison
- 818 Project (ACCMIP), Atmos. Chem. Phys., 13, 3063–3085, 2013.
- Strode, S. A., and Pawson, S.: Detection of carbon monoxide trends in the presence of
  interannual variability, J. Geophys. Res. Atmos., 118, 12257-12273, 2013.
- Sun, J., Fu, J. S., Drake, J., Lamarque, J.-F., Tilmes, S., and Vitt, F.: Improvement of the
  prediction of surface ozone concentration over conterminous U.S. by a computationally
  efficient second-order Rosenbrock solver in CAM4-Chem, J. Adv. Model Earth. Sy., 9,
  482–500, 2017.
- Tilmes, S., Lamarque, J.-F., Emmons, L. K., Conley, A., Schultz, M. G., Saunois, M., Thouret,
  V., Thompson, A. M., Oltmans, S. J., Johnson, B., and Tarasick, D.: Technical Note:
  Ozonesonde climatology between 1995 and 2011 : description, evaluation and applications,
  Atmos. Chem. Phys., 12, 7475-7497, 2012.
- Tilmes, S., Lamarque, J. F., Emmons, L. K., Kinnison, D. E., Ma, P. L., Liu, X., Ghan, S.,
- 830 Bardeen, C., Arnold, S., Deeter, M., Vitt, F., Ryerson, T., Elkins, J. W., Moore, F.,
- 831 Spackman, J. R., and Val Martin, M.: Description and evaluation of tropospheric chemistry

- and aerosols in the Community Earth System Model (CESM1.2), Geosci. Model Dev., 8,
  1395–1426, 2015.
- Tilmes, S., Lamarque, J. F., Emmons, L. K., Kinnison, D. E., Marsh, D., Garcia, R. R., Smith, A.
  K., Neely, R. R., Conley, A., Vitt, F., Val Martin, M., Tanimoto, h., Simpson, I., Blake, D.
  R., and Blake, N.: Representation of the Community Earth System Model (CESM1) CAM4chem within the Chemistry-Climate Model Initiative (CCMI), Geosci. Model Dev., 9,
  1853–1890, 2016.
- Travis, K. R., Jacob, D. J., Fisher, J. A., Kim, P. S., Marais, E. A., Zhu, L., Yu, K., Miller, C. C.,
  Yantosca, R. M., Sulprizio, M. P., Thompson, A. M., Wennberg, P. O., Crounse, J. D., St.
  Clair, J. M., Cohen, R. C., Laughner, J. L., Dibb, J. E., Hall, S. R., Ullmann, K., Wolfe, G.
  M., Pollack, I. B., Peischl, J., Neuman, J. A., and Zhou, X.: Why do models overestimate
  surface ozone in the Southeast United States?, Atmos. Chem. Phys., 16, 13561–13577,
  2016.
- US EPA: National Ambient Air Quality Standards for Ozone: Final Rule. Fed. Regist. 80 (206),
  65292-65468. 2015.
- US EPA: CASTNET 2014 Annual Report Prepared by Environmental Engineering and
   Measurement Services, Inc. for the U.S. Environmental Protection Agency, 2016.
- Weatherhead, E. C., Stevermer, A. J., and Schwartz, B. E., Detecting environmental changes and
   trends, Physics and Chemistry of the Earth, 27, 399-403, 2002.
- Wild, O., and Prather, M. J.: Global tropospheric ozone modeling: Quantifying errors due to grid
   resolution, J. Geophys. Res., 111, D11305, 2006.
- 853 Young, P. J., Archibald, A. T., Bowman, K. W., Lamarque, J.-F., Naik, V., Stevenson, D. S.,
- Tilmes, S., Voulgarakis, A., Wild, O., Bergmann, D., Cameron-Smith, P., Cionni, I.,
- 855 Collins, W. J., Dalsøren, S. B., Doherty, R. M., Eyring, V., Faluvegi, G., Horowitz, L. W.,
- Josse, B., Lee, Y. H., MacKenzie, I. A., Nagashima, T., Plummer, D. A., Righi, M.,
- 857 Rumbold, S. T., Skeie, R. B., Shindell, D. T., Strode, S. A., Sudo, K., Szopa, S., and Zeng,
- 6.: Pre-industrial to end 21st century projections of tropospheric ozone from the
- 859 Atmospheric Chemistry and Climate Model Intercomparison Project (ACCMIP), Atmos.
- 860 Chem. Phys., 13, 2063–2090, 2013.

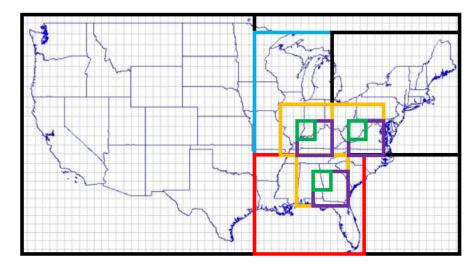
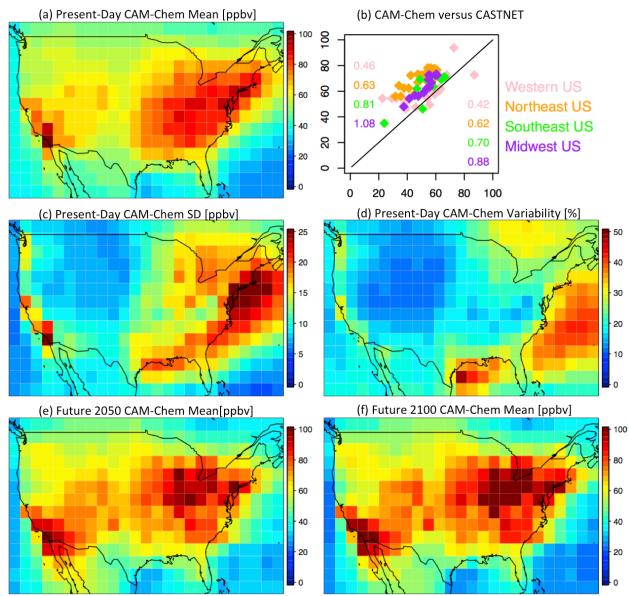




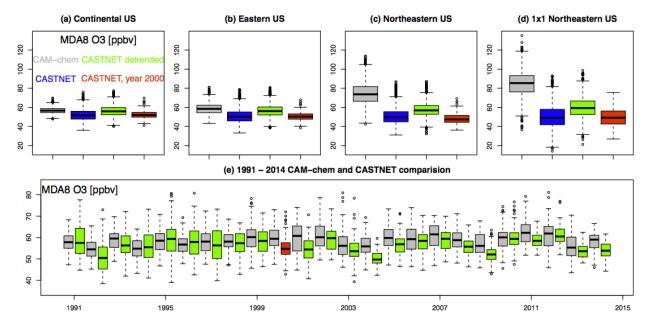
Figure 1: Telescoping Spatial Regions included in this study. The largest scale we consider is the Continental
US (outer border). We focus on the Eastern US, by subdividing into three subregions: the Midwest (blue),
Northeast (black), and Southeast (red). Within each subregion we telescope into a 3x3 grid cell (yellow), 2x2
grid cell (purple), and a 1x1 grid cell (green). In the paper, we only show a subset of these telescoping regions,

- 866 and we include the rest in the Supplemental Material.
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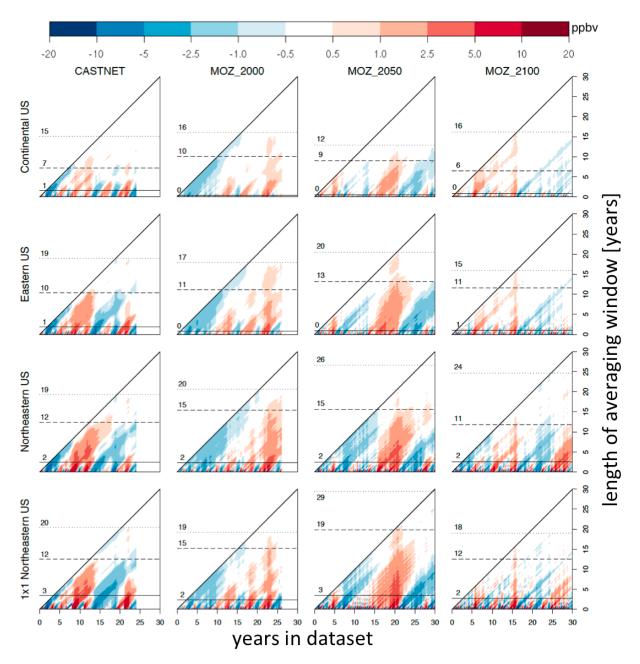
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Figure 2: Continental US surface maps of (a) present-day CAM-chem mean MDA8 O<sub>3</sub>; (b) CAM-Chem (y-871 axis) comparison to CASTNET observations (x-axis) for the year 2000 (see Brown-Steiner et al. (in review) 872 for additional comparisons); (c) present-day CAM- chem standard deviation of MDA8 O<sub>3</sub>; (d) present-day 873 CAM- chem variability (standard deviation divided by mean, as a percent); (e) future CAM- chem year 2050 874 mean MDA8 O3; and (f) future CAM- chem year-2100 mean MDA8 O3. All model results are averaged over 875 every JJA day in the time series, while the CASTNET results are only for the year 2000. The numbers in 876 Figure 2b are slopes (left) and R<sup>2</sup> values (right).



878 879

880 Figure 3: (a-d): Boxplots for surface MDA8 O<sub>3</sub> for every summertime (JJA) day from 1991 – 2014 averaged 881 over the Continental US, the Eastern US, the Northeastern US, and a single grid cell in the Northeastern US 882 from CAM-chem (grey), CASTNET observations (blue), detrended CASTNET observations centered at the 883 year 2000 (green), and since the CAM-chem simulations have cycled year-2000 emissions and boundary 884 conditions, the CASTNET values for the year 2000 only (red). (e) Comparison of the yearly JJA MDA8 O3 885 estimates averaged over the Eastern US for CAM-chem (grey) and the detrended CASTNET (green) from 886 1991 – 2014. The single red boxplot coincides with the red boxplot in (b). The units are in ppby, and for each 887 boxplot the box contains the Inter Quartile Range (IQR), the horizontal line within the box is the median, and 888 the whiskers extend out to the farthest point which is within 1.5 times the IQR with circles indicating any 889 outliers.





894 Figure 4: Comparisons of the variability represented by the summertime MDA8 O<sub>3</sub> anomaly (from the 895 long-term summertime mean) for the four datasets in this study (CASTNET, MOZ 2000, MOZ 2050, 896 MOZ 2100, columns) averaged over the four telescoping regions (CUS, EUS, NEUS, NEUS 1x1, rows). 897 In each panel, the horizontal axis is the number of years in the dataset (24 years (1991-2014) for 898 CASTNET, 26 years (1990-2015) for MOZ 2000, and 30 years (2036-2065 and 2086-2115) for 899 MOZ 2050 and MOZ 2100), and the vertical axis represents the length of the averaging window 900 (ranging from 1 day (bottom row) up to the entire time series (top pixel)). Each pixel represents the 901 estimate of the ozone anomaly for a given averaging window (vertical axis) ending at a given time 902 (horizontal axis). Horizontal lines indicate the length of averaging window required to guarantee that 903 the variability drops below thresholds of 5 ppbv (solid), 1 ppbv (dashed), and 0.5 ppbv (dotted). 904

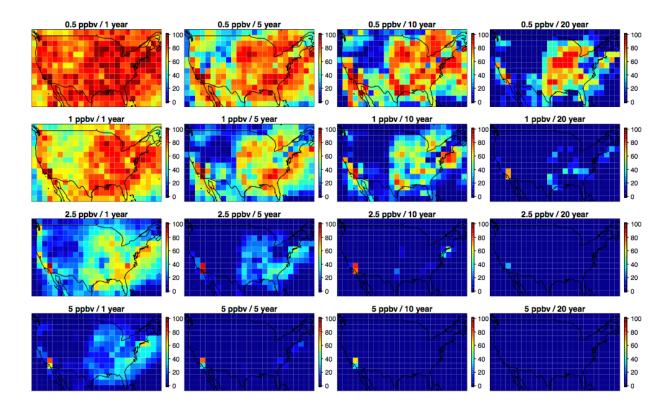
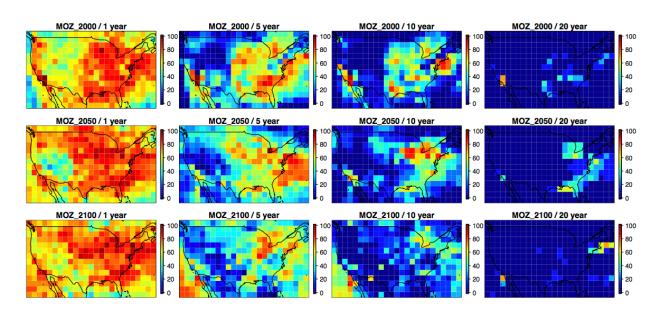


Figure 5: Spatial Plots over the Continental US plotting the likelihood (%) that an estimate of ozone

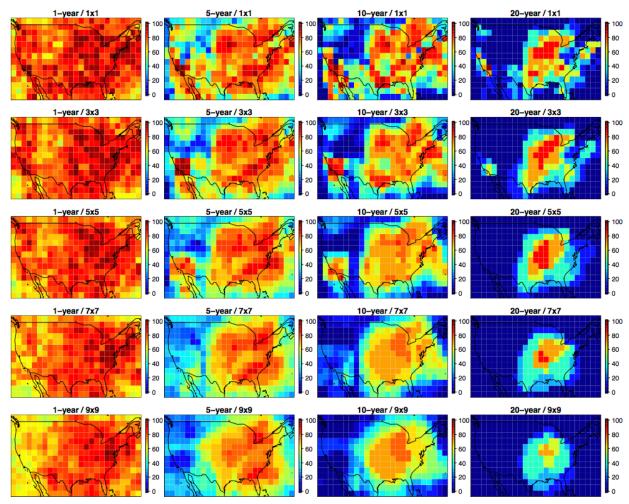
exceeds a given threshold due to meteorological variability (rows) at the grid-cell level when using

different lengths of averaging windows (columns) for the present-day CESM simulation (MOZ 2000).



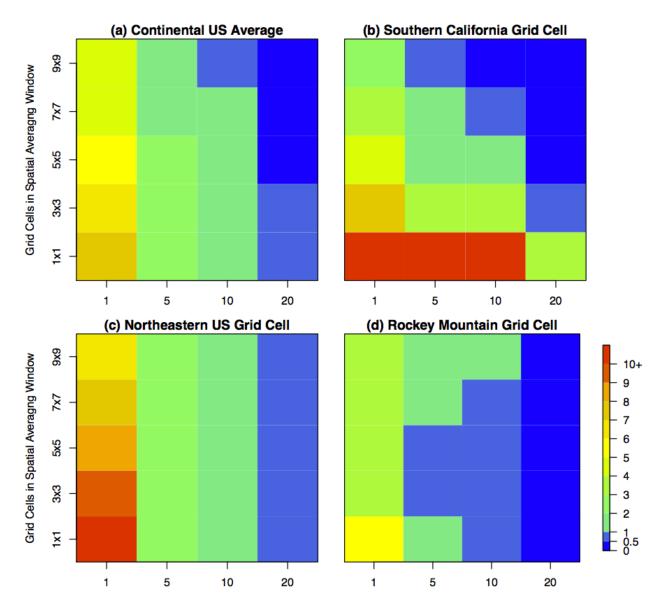
914 Figure 6: As in Figure 5, but only the second row (1 ppbv threshold), for present-day CAM-chem

- 915 (MOZ\_2000), future CAM-chem 2050 (MOZ\_2050), and future CAM-chem 2100 (MOZ\_2100).



918 919 Figure 7: Combined impact of temporal and spatial averaging on reducing ozone variability on the 920 likelihood (%) of exceeding the 0.5 ppbv threshold (as in Figures 5, 6, and Supplemental Figure S3) for 921 the present-day MOZ 2000 simulation. The top row is the same as in Figure 6, while the lower rows

- 922 have averaged the values within a 3x3, 5x5, 7x7, and 9x9 grid box surrounding each individual grid cell.
- 923



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926 Figure 8: The maximum potential calculated MDA8 O<sub>3</sub> anomaly [ppbv] from the long-term mean for (a) 927 the Continental US average and three individual grid cells taken from (b) Southern California, 928 demonstrating effective temporal and spatial averaging, (c) the Northeast, where spatial averaging is 929 ineffective, and (d) the Rocky Mountains, where spatial averaging initially reduces the anomaly, but 930 then increases the anomaly as surrounding regions get included in the spatial average. The number of 931 years included in the temporal averaging window increase along the x-axis and the number of grid cells 932 included in the spatial averaging window increase along the y-axis. A full map of the Continental US can 933 be found in the Supplemental Material (Figure S4). Note that the color scale is non-linear, and the color 934 transitions are selected to match the thresholds established throughout this paper.

			CASTNET	MOZ_2000	MOZ_2050	MOZ_2100	
	Mean	ppbv	52.4	56.7	56.8	57.4	
Continental US	Standard Deviation	ppbv	5.04	3.08	3.54	3.73	
Continental US	Variability	%	10%	5%	6%	7%	
	Bias	ppbv		4.31			
	Mean	ppbv	50.7	58.6	55.5	56.5	
Eastern US	Standard Deviation	ppbv	5.78	5.77	5.80	6.50	
Lastern US	Variability	%	11%	10%	10%	12%	
	Bias	ppbv		7.91			
	Mean	ppbv	48.3	74.4	68.4	73.0	
Northeastern US	Standard Deviation	ppbv	6.89	11.4	11.1	12.7	
Northeastern US	Variability	%	14%	15%	16%	17%	
	Bias	ppbv		26.1			
	Mean	ppbv	49.6	84.9	81.1	85.1	
1x1 Northestern US	Standard Deviation	ppbv	10.2	12.8	16.7	17.3	
1x1 Normestern US	Variability	%	21%	15%	21%	20%	
	Bias	ppbv		35.3			

936 937 938 Table 1: Statistical Summary of the CASTNET observations and the three CAM-chem simulations for different spatial averaging regions within the US. Variability is defined as the standard deviation divided by the mean value (in percent). Biases are only included for the present-day CAM-chem 939 940 simulation compared to the CASTNET data. Similar tables for the other regions in this study are 941 included in the Supplemental Material.