

1 **Maximizing Ozone Signals Among Chemical, Meteorological, and Climatological**
2 **Variability**

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21

22 **Abstract**

23
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_x 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. We recommend temporal averaging of at least 10 - 15 years combined with
43 regional spatial averaging over several hundred kilometer spatial scales. If this level of averaging
44 is not practical (e.g. the signal being examined is at a local scale), we recommend some
45 exploration of the spatial and temporal variability to provide context and confidence in the
46 robustness of the result. These results are consistent between simulated and observed data, and
47 within a single model with different sets of parameters. The strategies selected in this study are
48 not limited to surface ozone data, and could potentially maximize signal detection capabilities
49 within a broad array of climate and chemical observations or model output.

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62

63 **1 Introduction**

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

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

94 assume that all members in the ensemble are independent and equally skillful. This assumption,
95 however, may result in a loss of some valuable information (Knutti, 2010). Another form of this
96 strategy is to run multiple model runs within a single model, but under different initial conditions
97 or sets of parametric assumptions (e.g. Deser et al, 2012; Monier et al., 2013, 2015; Kay et al.,
98 2015; Garcia-Menendez et al., 2015, 2017). This approach cannot address structural uncertainties
99 and internal (unforced) variability between models, but is capable of identifying parametric
100 uncertainties within a single model.

101 The second strategy to reduce ozone variability is to expand the temporal averaging window,
102 which can influence the interpretation of the determined ozone value (e.g. Brown-Steiner et al.,
103 2015). The Environmental Protection Agency (EPA) National Ambient Air Quality Standard
104 (NAAQS) for ozone (US EPA, 2015) explicitly takes this into account, both in the length of the
105 averaging period (daily maximum 8-hour average) and the selection criteria for the standard
106 (fourth-highest over the previous 3 years). The calculated ozone variability can be further
107 reduced by utilizing even longer averaging periods, such as monthly (e.g. Rasmussen et al.,
108 2012), seasonal (e.g. Fiore et al., 2014; Barnes et al., 2016), annual, or decadal mean values (e.g.
109 Garcia-Menendez et al., 2017). This strategy is analogous to the averaging of meteorological
110 data to derive a climate signal, and just as Lewandowsky et al. (2015) recommend averaging 17
111 or more years in order to achieve climatological estimates of temperature trends, there is a
112 growing body of literature recommending averaging short time scale chemical variability (what
113 could be called chemical weather, see Lawrence, 2005) for 15 or more years (e.g. Garcia-
114 Menendez et al, 2017) in order to achieve an estimate of the what could be called the chemical
115 climate (see Möller, 2010).

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

125 also been noted as an important consideration when analyzing aerosol indirect effects
126 (McComiskey and Feingold, 2012) and for the detection and attribution of extreme weather
127 events (Angélil et al., 2017).

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

140 The second research perspective is to identify an ozone signal of a certain magnitude (or
141 threshold) and decide what spatial and temporal averaging scales are needed to best identify that
142 signal. The ozone signal could be large (e.g. determining the effectiveness or compliance with a
143 5 ppbv incremental reduction of the EPA NAAQS for ozone (US EPA, 2015)) or small (e.g.
144 identifying annual ozone trends within the US, which Cooper et al. (2012) show can be on the
145 order of 0.10 – 0.45 ppbv), and can be highly sensitivity to spatial and temporal heterogeneity
146 and meteorological variability. Barnes et al. (2016) found that surface ozone trends over 20-year
147 periods can vary by ± 2 ppbv due solely to climate variability, while interannual variability can
148 be on the order of ± 15 ppbv (Fiore et al., 2003; Tilmes et al., 2012; Lin et al., 2014) and day-to-
149 day variability can be even larger, extending regularly from near-background levels of 40 – 50
150 ppbv up to 100 ppbv during the summertime (Fiore et al., 2014).

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

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

166 In Section 2, we describe the CAM-chem model and our simulations, as well as the
167 CASTNET observational database and the regional definitions used throughout this paper. In
168 Section 3 we quantify the temporal and spatial variability of surface ozone, show how temporal
169 and spatial averaging reduces the calculated ozone variability, and demonstrate the spatial
170 heterogeneity of the calculated ozone variability. In Section 4, we discuss the potential strategies
171 that could be used to maximize ozone signal detection due to meteorological variability, explore
172 uncertainties, and make recommendations for future research.

173

174 **2 Methods**

175

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

184

185 **2.1 CAM-chem**

186 The present-day simulation (MOZ_2000) was conducted using CAM-chem model
187 version 1.2.2, with the CAM4 atmospheric component (see Tilmes et al., 2015; 2016 for model

188 description and evaluation). The model has been used extensively for a wide range of
189 atmospheric chemistry research and is included in the ACCMIP (Lamarque et al., 2012; Young
190 et al., 2012 and references therein). We conduct our simulations using the Model for Ozone and
191 Related chemical Tracers, version 4 (MOZART-4) chemical mechanism (Emmons et al., 2010),
192 which is a full tropospheric chemical mechanism integrated into CAM-Chem (e.g. Lamarque et
193 al., 2012; Tilmes et al., 2015; Brown-Steiner et al., in review). Offline forced meteorology from
194 the Modern-Era Retrospective analysis for Research and Applications (MERRA) reanalysis
195 product (Rienecker et al., 2011) for 26 meteorological years (1990 – 2015). Additional model
196 evaluation and comparisons to surface and ozonesonde observations can be found in Brown-
197 Steiner et al. (in review). This simulation has 56 vertical levels – adopted from MERRA
198 meteorology – and 96 latitudinal and 144 longitudinal grid cells. We aim to isolate the variability
199 to the meteorologically-driven impact on atmospheric chemistry so we repeat year-2000
200 anthropogenic emissions from the ACCMIP (Atmospheric Chemistry and Climate Model
201 Intercomparison Project) inventory (Lamarque et al., 2012) and all non-biogenic emissions for
202 all meteorological years, and include specified long-lived stratospheric species (O_3 , NO_x , HNO_3 ,
203 N_2O , N_2O_5) as in MOZART-4 (Emmons et al., 2010), an online biogenic emissions model
204 MEGAN (Guenther et al., 2012), and forced sea ice and sea surface temperatures to year 2000
205 historical conditions. Like many state-of-the-art chemical tracer models, the CAM-chem exhibits
206 some biases, most notably for our purposes a high bias in simulated surface ozone in the Eastern
207 US (e.g. Lamarque et al., 2012; Brown-Steiner et al., 2015; Travis et al., 2016; Barnes et al.,
208 2016). Recent efforts have been successful in partially reducing these biases (e.g. Sun et al.,
209 2017).

210 We also include two reference simulations of the future climate, MOZ_2050 and
211 MOZ_2100 (simulating the meteorological years 2035 – 2065 and 2085 – 2115, respectively)
212 using the CESM CAM-chem simulations described in detail by Garcia-Menendez et al. (2017)
213 with one set of initial condition data, and a climate sensitivity of 3.0 °C. These simulations do
214 not include projections of any changes in future emissions. Compared to the present-day
215 simulation (MOZ_2000), these future simulations (MOZ_2050 and MOZ_2100) have several
216 parametric differences: the model version is 1.1.2 (see Tilmes et al., 2015 and references for
217 information on model development), the atmospheric component is CAM3, the emissions (which
218 are held constant at year-2000 levels) are from the Precursors of Ozone and their Effects in the

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

223

224 **2.2 CASTNET**

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

238

239 **2.3 Telescoping Regional Definitions**

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

249

250 **2.4 Temporal Averaging Windows**

251 To explore the impact of temporal averaging, we examine ozone across a range of
252 temporal averaging windows, from 1 day up to the full 26 years for the CESM data (1990-2015),
253 the full 24 years for the detrended CASTNET data (1991 – 2014), and the 30 years available
254 from the future scenarios of Garcia-Menendez et al. (2017). Each averaging window, therefore,
255 can be considered to be a “sample” of possible realizations of meteorology. For instance, a
256 selection of an averaging window of 1 year has 26 possible slices within the 1990 – 2015
257 MOZ_2000 data, while a selection of an averaging window of 10 years has 17 possible slices
258 within the CESM data ($N = \# \text{ years} - \text{length of window} + 1$). In this study, we consider all
259 realizations to be equally likely and compare them to each other and to the long-term trend.
260 However, if we were only able to simulate 5 years, we would not be able to compare to the long-
261 term trend, and so be unable to completely quantify the likelihood of error in the context of the
262 long-term behavior.

263

264 **3 Results**

265 Here we examine the spatial and temporal behavior of MOZ_2000, MOZ_2050, and
266 MOZ_2100 and compare MOZ_2000 to present-day CASTNET observations. We introduce the
267 moving temporal averaging windows, explore possible thresholds of acceptable error or signal
268 strength, and examine the influence of expanding spatial averaging regions. Finally, we combine
269 these temporal and spatial averaging techniques into a single framework.

270

271 **3.1 Spatial and Temporal Comparisons**

272 Figure 2 plots the averaged spatial distribution of the maximum daily 8-hour average
273 ozone (MDA8 O₃) for summertime (JJA) days for 1990-2015 for the present-day CAM-chem
274 simulation, MOZ_2000 (Figure 2a) and compares to the year 2000 for CASTNET data (Figure
275 2b). Some of the averaging strategies we present can average away the high ozone behavior this
276 MDA8 O₃ metric is intended to quantify, but it is such a well-reported metric that focusing our
277 analysis on it allows for ready comparisons to other studies. The well-known high ozone bias in
278 the Eastern US (e.g. Lamarque et al., 2012; Travis et al., 2016; Barnes et al., 2016) is apparent,
279 but otherwise the spatial variability over the entire Continental US is well captured. While we do
280 examine the magnitude of surface ozone in this paper, most of our analysis is focused on the

281 variability around the mean value (the anomaly), and as we show below, the CASTNET
282 observations and CESM results are largely consistent in their representation of ozone variability
283 (Figure 2, Table 1). The standard deviation of the simulated MDA8 O₃ is large over the Eastern
284 US and the Pacific Coast, with peak values of ± 25 ppbv over the highly populated Atlantic
285 Coast (Figure 2c). The variability (defined as the standard deviation divided by the mean,
286 expressed as a percentage) is lowest over the Western US ($\sim 15\%$), only slightly higher over the
287 Eastern US (up to 25%), and highest (up to 50%) over the coastal regions (Figure 2d). We
288 include this relative standard deviation metric since the CAM-chem biases make it difficult to
289 compare standard deviations directly. The future climate simulations, MOZ_2050 and
290 MOZ_2100 (Figure 2e and 2f, respectively), although run with different parametric settings than
291 MOZ_2000 (see Section 2), simulate a similar spatial distribution of surface ozone, although
292 under the warmer simulated climate of 2050 and 2100. These future climate simulations have a
293 similar spatial pattern to the present-day simulation (Figure 2a), with high ozone levels in the
294 Eastern US that increases from 2050 to 2100 (see Garcia-Menendez et al. (2017) for more
295 details).

296 Figure 3 compares boxplots over the four telescoping regions (Figure 1) for MOZ_2000,
297 the CASTNET data, the detrended CASTNET data, and for the single year 2000 for the
298 CASTNET data (Figures 3a-d), and Table 1 summarizes relevant statistics. In order to compare
299 CASTNET ozone to the simulated ozone, which do not have a trend over time, we detrend the
300 CASTNET data in order to remove the impact of any temporal trends (e.g. NO_x emissions
301 reductions) on ozone. The Northeastern US ozone bias is apparent at the smaller spatial scales
302 (Figures 3c,d) and is less apparent when averaging over larger regions (Figures 3a,b). Figure 3e
303 compares the year-to-year boxplots of the JJA MDA8 O₃ for the MOZ_2000 and the detrended
304 CASTNET data, and demonstrates the variability both in the median and spread of the ozone
305 values in both the modeled and simulated data. While the MOZ_2000 ozone is generally higher
306 than the CASTNET data, there are years in which the CASTNET data has higher ozone
307 extremes. The red box plot in Figure 3e, which corresponds to the red box plot in Figure 3b,
308 indicates that the year 2000 was an anomalously low year for observed ozone, although not the
309 lowest.

310 While all the CESM CAM-chem simulations have high ozone biases in the Northeastern
311 US (Figures 2 and 3, Table 1), their capability to simulate ozone variability is consistent with the

312 available observations (for present day) and for expectations of ozone variability changes in the
313 future climate (for MOZ_2050 and MOZ_2100). It is clear that variability increases when the
314 size of the averaging region decreases, a fact that is well noted in the literature, as in Hawkins
315 and Sutton (2012) for climate variables and Barnes et al. (2016) for ozone. As can be seen in in
316 Table 1, the CASTNET variability increases as the spatial scale decreases (10%, 13%, 16%, and
317 20% for our telescoping regions from continental to a single Northeastern U.S. grid box), and
318 MOZ_2000 largely captures this trend, albeit with lower overall variability (5%, 10%, 15%, and
319 15%). This increase in ozone variability with decreasing spatial scale is maintained in the future
320 climate simulations (6%, 10%, 16%, and 21% for MOZ_2050 and 7%, 12%, 17%, and 20% for
321 MOZ_2100). Table S1 contains statistics for the other telescoping regions.

322
323

324 **3.2 Variability, Averaging Windows, and Thresholds**

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

335 Figure 4 presents this likelihood by plotting all possible estimates of MDA8 O₃ (as
336 anomalies from the long-term mean) over all possible selections of averaging window (from 1
337 day up to the complete time series) for our telescoping regions. The semi-cyclical and highly
338 auto-correlated nature of surface ozone is apparent at all spatial scales, with alternating cycles of
339 anomalously high and low ozone. The temporal impact of anomalous ozone events is indicated
340 by the vertical and right-leaning diagonal striations, which show that anomalous ozone events
341 can impact estimates of ozone values within averaging windows up to 15 or 20 years. Figure 4
342 demonstrates how small-scale anomalously high or low ozone values (that come only from
343 meteorological variability) can impact temporal averages of 5, 10, or even 20 years. For instance,

344 a selected 5-year averaging window within the MOZ_2000 simulation averaged over the
345 Northeastern US could be 2.5 ppbv higher or lower than the 25-year mean value of 74 ppbv, a
346 potential error of 7%. Horizontal lines in Figure 4 mark the length of averaging windows that are
347 needed to ensure that ozone anomaly for any selection of averaging window does not exceed a
348 given threshold (5, 1, and 0.5 ppbv for solid, dashed, and dotted lines respectively). This
349 potential error is larger within smaller regions and at the shorter selections of the averaging
350 window. While the high and low ozone anomalies differ in time between CASTNET,
351 MOZ_2000, MOZ_2050, and MOZ_2100 in Figure 4, the impact of spatial and temporal
352 averaging is consistent.

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

367 Figures 4 and Supplemental Figures S1 and S2 compare the CASTNET observations to
368 the three CESM CAM-chem simulations, and while there are minor differences, there are broad
369 features that are consistent. First, using longer temporal averaging windows reduces the
370 influence of small-scale ozone variability at all spatial scales, and depending on the acceptable
371 threshold, one can select a temporal scale that effectively reduces the likelihood of exceeding
372 that threshold to zero. Second, larger spatial scales also reduce this likelihood of exceeding a
373 given threshold, but not as effectively as longer temporal scales. Finally, the impact of both

374 temporal and spatial averaging on ozone variability is largely consistent for the CASTNET
375 observations and for all three CESM CAM-chem simulations.

376

377 **3.3 Selection of Temporal Averaging Scales**

378 Figure 5 extends this analysis to examine the spatial heterogeneity of this likelihood of
379 the meteorological variability causing ozone anomalies exceeding particular thresholds at the
380 grid cell level. Here we plot four thresholds (0.5, 1, 2.5, and 5 ppbv) and four averaging windows
381 (1, 5, 10, and 20 years) for the MOZ_2000 simulation. Ozone variability is highest in the Eastern
382 US. At the grid-cell level, there are two strategies for filtering out the noise associated with
383 natural meteorological variability (and thus enhancing signal detection capabilities): either
384 average over longer periods, or acknowledge the level of noise and increase the threshold. For
385 these data, it is virtually certain that any 20-year average will be within 5 ppbv of a full 25-year
386 mean value (which itself may not be an accurate representation of a longer simulation), and
387 virtually certain that any 1-year average will be at least 0.5 ppbv away from the mean.

388 Supplemental Figure S3 extends the analysis of Figure 5 by comparing the MOZ_2000,
389 MOZ_2050, and MOZ_2100 simulations across the four thresholds for the 5-year averaging
390 window. Figure 6 similarly compares the 1 ppbv ozone threshold across the four averaging
391 windows for MOZ_2000, MOZ_2050, and MOZ_2100. Interpreting Figures 6 and Supplemental
392 Figure S3 give largely consistent interpretations than the analysis above (Figure 5). Namely, that
393 at the grid-scale level, increasing the temporal averaging window (Figure 6) or increasing the
394 acceptable ozone threshold (Supplemental Figure S3) are effective at reducing the impact of the
395 meteorological variability on estimates of the ozone signal. Shorter windows (or smaller
396 thresholds) are needed in the Western US (where variability is smaller, see Figure 2d) than in the
397 Eastern US (where variability is larger) as well as over coastal and highly populated regions.
398 Finally, the 1 ppbv threshold and the 5-year averaging window plots (in either Figure 5 or
399 Supplemental Figure S3) indicate that the spatial distribution and location of the peak variability
400 may shift into the future, although this may be due to parametric differences between
401 MOZ_2000, MOZ_2050, and MOZ_2100. Future simulations will be needed to check this shift
402 in peak ozone variability.

403

404 **3.4 Selection of Spatial Averaging Scales**

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

418

419 **3.5 Combination of Spatial and Averaging Scales**

420 We now examine the combined impact of temporal and spatial averaging on reducing the
421 influence of small-scale ozone variability in order to enhance ozone signal detection capabilities.
422 Table S2 summarizes our analysis by dividing the likelihood of the ozone variability estimates
423 exceeding selected thresholds away from the long-term mean into four categories: (1) the length
424 of the averaging window over which ozone is averaged (columns); (2) the magnitude of the
425 ozone threshold of interest (rows); (3) the observed (CASTNET) and modeled (MOZ_2000,
426 MOZ_2050, and MOZ_2100) ozone data (sub-columns); and (4) the size of the spatial extent
427 over which ozone is averaged (sub-rows). A graphical representation consistent with the data
428 presented in Table S2 is plotted in Figure 8 for the Continental US average and for three grid
429 cells that represent various cases. In each plot in Figure 8, by moving along columns from left to
430 right, we can see the influence of increasing the size of the temporal averaging window, and by
431 moving along rows (from the bottom to the top), we can see the influence of increasing the
432 spatial averaging scale. By taking in the entire plot as a whole, we can get a feel for the
433 combined influence of both temporal and spatial averaging. Supplemental Figure S4 contains a
434 plot for each grid cell in the Continental US.

435 On average within the Continental US, both temporal and spatial averaging are effective
436 at reducing the calculated MDA8 O₃ anomaly, although temporal averaging is more effective
437 (Figure 8a). There are many grid cells in the Eastern and Western US coasts (Figure 8b,
438 Supplemental Figure S4), where both spatial and temporal averaging are effective, but their
439 combined usage is especially effective. There are also many grid cells where temporal averaging
440 is effective, but spatial averaging is barely effective, or not effective at all (Figure 8c and
441 Supplemental Figure S4). Finally, there are some grid cells, particularly in the Central US
442 (Figure 8d and Supplemental Figure S4), where spatial averaging over smaller regions is
443 effective, but spatial averaging of larger regions actually increases the calculated MDA8 O₃
444 anomaly by including surrounding grid cells that have higher variability.

445

446 **4 Discussion**

447 We now return to the original three research questions posed in Section 1. First, what is
448 the magnitude of ozone variability due to meteorology alone at the smallest scale, and what is the
449 impact of increasing the scale of temporal and spatial averaging? In both observed and modeled
450 MDA8 O₃ surface data, the small-scale variability driven solely by the meteorological variability
451 impact on atmospheric chemistry (expressed as the standard deviation as a percentage of the
452 mean) can exceed 20% (Table 1, Figure 2d). The chemical variability examined here is the result
453 of fluctuations in meteorology, which itself results from larger-scale climatological drivers.
454 While variability in emissions also influences atmospheric chemistry, our analysis has removed
455 the influence of emissions variability and isolated the variability due to meteorology. A more
456 comprehensive analysis of chemical variability will need to account for both meteorological and
457 emission variability, which is complicated by temporal trends in both the emissions of ozone
458 precursor species and the climate.

459 There is high temporal and spatial heterogeneity of surface ozone variability (Figure 2d),
460 with the lowest values found in the Western US (< 10%), higher values found in the Eastern US
461 (up to 20%), and the highest values over coastal or heavily populated regions (up to 30%).
462 Averaging over longer temporal scales (by increasing the averaging window) and over larger
463 spatial scales (by expanding the averaging region) can reduce the magnitude of the calculated
464 variability, with temporal averaging proving to be more effective than spatial averaging in most
465 cases (Figure 8). In this study, we performed simple spatial averaging, but there are other

466 methodologies for smoothing two-dimensional signals (e.g. Räisänen et al., 2011; Pogson and
467 Smith, 2015) that could potentially increase signal detection capabilities.

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

478 Third, are these results dependent on the particular parameterizations of the CESM
479 CAM-chem model, and are they consistent with the available CASTNET observations? The
480 three CESM CAM-chem simulations exhibited consistent representations of ozone variability,
481 consistent with our understanding of future changes to the climate (and meteorology) and the
482 resulting impact on atmospheric chemistry (Table 1, Figures 4, S1, and S2). Compared to the
483 CASTNET observations (which we detrended to remove the influence of changing precursor
484 emissions), the present-day simulation (MOZ_2000) exhibited a high ozone bias in the Eastern
485 US, while the representation of the ozone variability is comparable (Table 1).

486 Finally, how may these strategies be applied to other datasets, be they chemical,
487 meteorological, or climatological? Much of this analysis could be applied to any dataset that has
488 spatial and temporal coverage, as long as some set of acceptable thresholds is provided. While
489 our time step in this analysis is daily (given the MDA8 O₃ metric), and applied only to
490 summertime (JJA) days, any time step (i.e. hourly, monthly, annual, decadal) could be utilized as
491 long as cyclical trends (e.g. diurnal or seasonal cycles) are removed. Indeed, the sliding-scale
492 presentation in Figure 8 and Supplemental Figure S4 can specifically be utilized to identify
493 particular spatial and temporal scales that are sufficient to identify signals at particular thresholds
494 and to identify particular geographic regions that are best suited to identify a given signal. For
495 example, Sofen et al. (2016) identified regions across the globe where additional observations
496 would be particularly suited to improve our understanding of surface ozone behavior, and our

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

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

524

525 **5 Conclusions**

526 We quantified the impact of spatial and temporal averaging at different scales – both
527 individually and combined – on estimates of summertime surface ozone variability and the

528 resulting likelihood of over-confidence in estimates of chemical signals over the United States
529 using CASTNET observations and the CESM CAM-chem model. We simulate three multi-
530 decadal time periods, each with constant surface emissions, and find that this analysis is
531 consistent across our simulated time periods, and that our results are not sensitive to particular
532 configurations and parametric choices within the CESM CAM-chem (i.e. emissions,
533 meteorology, and climate). We also provide a conceptual framework for gaining understanding
534 of the influence of spatial and temporal averaging that may be adapted to a wide range of
535 atmospheric and surface phenomena, provided sufficient spatial and temporal coverage. Here we
536 focus on summertime surface ozone, a highly variable (in both space and time) atmospheric
537 constituent with severe human health impacts and implications for planetary climate, which is
538 the focus of many local, regional, and national policies. However, the resultant magnitude of
539 these changes and signals are small compared to the magnitude of the day-to-day ozone
540 variability, and detecting these changes and signals can be challenging. In particular, it would be
541 impractical to delay interpreting observations for 10 – 15 years, or alternatively to expand the
542 spatial averaging such that small-scale features are smoothed away. Nonetheless, it is unwise to
543 over-interpret trends and signals based on observations from a limited spatial area and over a
544 short temporal period. Our analysis and conceptual framework presented here cannot solve this
545 tension, but it does demonstrate some strategies which can allow for a selection of spatial and
546 temporal averaging scales, and a consideration of the error threshold, that can aid in this signal
547 detection on a case-by-case basis. Taking into account the complex interactions involving trends
548 and variability between emissions, chemistry, meteorology, and climatology necessitates a
549 variety of strategies. This work quantifies the impact of spatial and temporal averaging in signal
550 detection, which can be used in conjunction with ensembles of simulations, statistical techniques,
551 and other strategies to further out understanding of the chemical variability in our atmosphere.

552 In order to quantify the impact of spatial and temporal averaging on summertime ozone
553 variability, we start by selecting four telescoping spatial regions (the Continental US, the Eastern
554 US, the Northeastern US, and a single grid cell within the Northeastern US) and examine all
555 possible choices for averaging windows (ranging from daily to multi-decadal windows),
556 although we focused primarily on averaging windows of 1, 5, 10, and 20 years. We find that –
557 consistent with previous studies – summertime ozone variability is largest at the smallest scales,
558 and is frequently on the order of $\pm 10 - 20$ ppbv, or which is roughly 15-20% of the mean ozone

559 signal. In order to minimize the chemical noise that results from meteorological variability – and
560 thus enhance the signal – we find averaging windows of 10-15 years (and sometimes longer at
561 the smaller spatial scales) combined with modest (nearest-neighbor) spatial averaging
562 substantially improve the capability for signal detection. We recognize that achieving a 10 – 15
563 year temporal averaging window is difficult, but this recommendation is consistent with recent
564 literature (e.g. Barnes et al., 2015; Garcia-Menendez et al., 2017). For studies where 10 – 15
565 years of averaging is impractical, we recommend that some spatial and temporal context is
566 provided that demonstrates that the signals being examined are robust and not the result of
567 internal variability or noise. We also recognize that our analysis is just one strategy for
568 enhancing signal detection capabilities, and will ideally be used alongside others, such as
569 perturbed initial condition ensembles, running simulations with either internal or forced
570 meteorology, and examining a region or time period with different models or parameterizations.

571 We show that the largest summertime ozone variability is found in the Eastern US (Figure 5,
572 Figure S4), and subsequently there are many regions within the Eastern US where even a 20-year
573 averaging window has a non-negligible likelihood of estimating ozone variability that is
574 dependent (with possible error in the 1 – 3 ppbv range) on the particular years selected. In
575 addition, over much of the Eastern US, simulations of 5-years or shorter have a substantial
576 likelihood (40 – 90%, Figures S1 and S2) of reflecting the influence of meteorological variability
577 on chemistry rather than the mean state of surface ozone, with the possibility of 5 – 10 ppbv
578 error (Figure S4). While we have detrended the CASTNET observations to compare to the
579 constant year-2000 cycled emissions in the simulations, the CASTNET time series inherently
580 includes the compounded variability of both meteorological and emission sources. Future studies
581 will need to expand this analysis to include trends and variability in the emissions, as well as in
582 the meteorology.

583 Finally, we demonstrate a conceptual framework that allows for a “sliding-scale” view of
584 surface ozone variability, in which both temporal and spatial averaging is examined at every grid
585 cell within the Continental US. We show that the magnitude of estimates of ozone variability can
586 be reduced with both temporal and spatial averaging, although temporal averaging tends to be
587 more effective. While there are many regions in which both temporal and spatial averaging used
588 in conjunction substantially reduce the estimate of ozone variability, there are some regions
589 where spatial averaging is ineffective, or even counter-effective. In contrast, this is not the case

590 for temporal averaging, which consistently reduces the magnitude of estimated ozone variability.
591 Our analysis could be combined with other studies (e.g. Sofen et al., 2016) to guide
592 observational and modeling strategies and identify regions and scales at which particular signals
593 are most likely to be identified.
594

595 **Code Availability**

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

600 **Data Availability**

601 The raw model output is archived on the NCAR servers, and processed data is archived at
602 <https://dspace.mit.edu/handle/1721.1/114467>.

603 **Supplemental Link**

604

605 **Author Contribution**

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

611

612 **Competing Interests**

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

614

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625

626 **References**

- 627 Angéilil, O., Stone, D., Perkins-Kirkpatrick, S., Alexander, L.V., Wehner, M., Shiogama, H.,
628 Wolski, P., Ciavarella, A., and Christidis, N.: On the nonlinearity of spatial scales in
629 extreme weather attribution statements, *Clim. Dyn.*, 2017.
- 630 Barnes, E. A., Fiore, A. M., and Horowitz, L. W.: Detection of trends in surface ozone in the
631 presence of climate variability, *J. Geophys. Res. Atmos.*, 121, 6112–6129, 2016.
- 632 Brown-Steiner, B., Hess, P. G., and Lin, M. Y.: On the capabilities and limitations of GCCM
633 simulations of summertime regional air quality: A diagnostic analysis of ozone and
634 temperature simulations in the US using CESM CAM-chem, *Atmos. Environ.*, 101, 134–
635 148, 2015.
- 636 Brown-Steiner, B., Selin, N. E., Prinn, R., Tilmes, S., Emmons, L., Lamarque, J.-F., and
637 Cameron-Smith, P.: Evaluating Simplified Chemical Mechanisms within CESM Version
638 1.2 CAM-chem (CAM4): MOZART-4 vs. Reduced Hydrocarbon vs. Super-Fast Chemistry,
639 *Geosci. Model Dev. Discuss.*, in review, 2018.
- 640 Camalier, L., Cox, W., and Dolwick, P.: The effects of meteorology on ozone in urban areas
641 and their use in assessing ozone trends, *Atmos. Environ.*, 41, 7127-7137, 2007.
- 642 Cooper, O. R., Gao, R. S., Tarasick, D., Leblanc, T., and Sweeney, C.: Long-term ozone trends
643 at rural ozone monitoring sites across the United States, 1990-2010, *J. Geophys. Res.*, 117,
644 D22307, 2012.
- 645 de Elía, R., Biner, S., and Frigon, A.: Interannual variability and expected regional climate
646 change over North America, *Clim. Dyn.*, 41, 1245, 2013.
- 647 Deser, C., Phillips, A., Bourdette, V., and Teng, H.: Uncertainty in climate change projections:
648 the role of internal variability, *Clim. Dyn.*, 38, 527, 2012.
- 649 Diem, J. E., and Comrie, A. C.: Predictive mapping of air pollution involving sparse spatial
650 observations, *Environmental Pollution*, 119, 1, 99–117, 2002.
- 651 Emmons, L. K., Walters, S., Hess, P. G., Lamarque, J.-F., Pfister, G. G., Fillmore, D., Granier,
652 C., Guenther, A., Kinnison, D., Laepple, T., Orlando, J., Tie, X., Tyndall, G., Wiedinmyer,
653 C., Baughcum, S. L., and Kloster, S.: Description and evaluation of the Model for Ozone
654 and Related chemical Tracers, version 4 (MOZART-4), *Geosci. Model Dev.*, 3, 43-67,
655 2010.
- 656 Fiore, A. M., Oberman, J. T., Lin, M. Y., Zhang, L., Clifton, O. E., Jacob, D. J., Naik, V.,
657 Horowitz, L. W., Pinto, J. P., and Milly, G. P.: Estimating North American background
658 ozone in U.S. surface air with two independent global models: Variability, uncertainties,
659 and recommendations, *Atmos. Environ.*, 96, 284–300, 2014.
- 660 Fiore, A. M., Jacob, D. J., Liu, H., Yantosca, R. M., Fairlie, T. D., and Li, Q.: Variability in
661 surface ozone background over the United States: Implications for air quality policy, *J.*
662 *Geophys. Res. Atmos.*, 108, D24, 1787, 2003.
- 663 Garcia-Menendez, F., Saari, R. K., Monier, E., and Selin, N. E.: U.S. Air Quality and Health
664 Benefits from Avoided Climate Change under Greenhouse Gas Mitigation, *Environ. Sci.*
665 *Technol.*, 49, 7580–7588, 2015.

- 666 Garcia-Menendez, F., Monier, E., and Selin, N. E.: The role of natural variability in projections
667 of climate change impacts on U.S. ozone pollution, *Geophys. Res. Lett.*, 44, 2911–2921,
668 2017.
- 669 Giorgi, F., and Bi, X.: Time of emergence (TOE) of GHG-forced precipitation change hot-spots,
670 *Geophys. Res. Lett.*, 36, L06709, 2009.
- 671 Guenther, A. B., Jiang, X., Heald, C. L., Sakulyanontvittaya, T., Duhl, T., Emmons, L. K., and
672 Wang, X.: The Model of Emissions of Gases and Aerosols from Nature version 2.1
673 (MEGAN2.1): An extended and updated framework for modeling biogenic emissions,
674 *Geosci. Model Dev.*, 5, 1471–1492, 2012.
- 675 Hawkins, E., and Sutton, R.: Time of emergence of climate signals, *Geophys. Res. Lett.*, 39,
676 L01702, 2012.
- 677 Jacob, D. J., and Winner, D. A.: (2009). Effect of climate change on air quality, *Atmos. Environ.*,
678 43, 51–63, 2009.
- 679 Jhun, I., Coull, B. A., Schwartz, J., Hubbell, B., and Koutrakis, P.: The impact of weather
680 changes on air quality and health in the United States in 1994–2012, *Environ. Res. Lett.*, 10,
681 084009, 2015.
- 682 Kay, J. E., Deser, C., Phillips, A., Mai, A., Hannay, C., Strand, G., Arblaster, J. M., Bates, S. C.,
683 Danabasoglu, G., Edwards, J., Holland, M., Kushner, P., Lamarque, J.-F., Lawrence, D.,
684 Lindsay, K., Middleton, A., Munoz, E., Neale, R., Oleson, K., Polvani, L., and Vertenstein,
685 M.: The Community Earth System Model (CESM) large ensemble project: A community
686 resource for studying climate change in the presence of internal climate variability, *Bull.*
687 *Amer. Meteor. Soc.*, 96, 1333–1349, 2015.
- 688 King, A. D., Donat, M. G., Fischer, E. M., Hawkins, E., Alexander, L. V., Karoly, D. J., Dittus,
689 A. J., Lweis, S. C., and Perkins, S. E.: The timing of anthropogenic emergence in simulated
690 climate extremes, *Environ. Res. Lett.*, 10, 094015, 2015.
- 691 Knote, C., Tuccella, P., Curci, G., Emmons, L., Orlando, J. J., Madronich, S., Baró, R., Joménez-
692 Guerrero, P., Luecken, D., Hogrefe, C., Forkel, R., Werhahn, J., Hirtl, M., Pérez, J. L., José,
693 R. S., Giordano, L., Brunner, D., Yahya, K., and Zhang, Y.: Influence of the choice of gas-
694 phase mechanism on predictions of key gaseous pollutants during the AQMEII phase-2
695 intercomparison, *Atmos. Environ.*, 115, 553–568, 2015.
- 696 Knutti, R.: The end of model democracy?, *Clim. Change*, 102, 395–404, 2010.
- 697 Lamarque, J.-F., Bond, T. C., Eyring, V., Granier, C., Heil, A., Klimont, Z., Lee, D., Liousse, C.,
698 Mieville, A., Owen, B., Schultz, M. G., Shindell, D., Smith, S. J., Stehfest, E., Van
699 Aardenne, J., Cooper, O. R., Kainuma, M., Mahowald, N., McConnell, J. R., Naik, V.,
700 Riahi, K., and Van Vuuren, D. P.: Historical (1850–2000) gridded anthropogenic and
701 biomass burning emissions of reactive gases and aerosols: Methodology and application,
702 *Atmos. Chem. Phys.*, 10, 7017–7039, 2010.
- 703 Lamarque, J.-F., Dentener, F., McConnell, J., Ro, C.-U., Shaw, M., Vet, R., Bergmann, D.,
704 Cameron-Smith, P., Dalsoren, S., Doherty, R., Faluvegi, G., Ghan, S. J., Josse, B.,
705 MacKenzie, I. A., Plummer, D., Shindell, D. T., Skeie, R. B., Stevenson, D. S., Strode, S.,
706 Zeng, G., Curran, M., Dahl-Jensen, D., Das, S., Fritzsche, D., and Nolan, M.: Multi-model
707 mean nitrogen and sulfur deposition from the atmospheric chemistry and climate model

708 intercomparison project (ACCMIP): Evaluation of historical and projected future changes,
709 *Atmos. Chem. Phys.*, 13, 7997–8018, 2013.

710 Lamarque, J.-F., Emmons, L. K., Hess, P. G., Kinnison, D. E., Tilmes, S., Vitt, F., Heald, C. L.,
711 Holland, E. A., Lauritzen, P. H., Neu, J., Orlando, J. J., Rasch, P. J., and Tyndall, G. K.:
712 CAM-chem: Description and evaluation of interactive atmospheric chemistry in the
713 Community Earth System Model, *Geosci. Model Dev.*, 5, 369–411, 2012.

714 Lawrence, M. G., Hov, Ø., Beekmann, M., Brandt, J., Elbern, H., Eskes, H., Feichter, H., and
715 Takigawa, M.: The chemical weather, *Environ. Chem*, 2, 6–8, 2005.

716 Lewandowsky, S., Risbey, J. S., and Oreskes, N.: On the definition and identifiability of the
717 alleged “hiatus” in global warming, *Sci. Rep.*, 5, 16784, 2015.

718 Lin, M., Horowitz, L. W., Oltmans, S. J., Fiore, A. M., and Fan, S.: Tropospheric ozone trends at
719 Mauna Loa Observatory tied to decadal climate variability, *Nat. Geosci.*, 7, 136–143, 2014.

720 McComiskey, A., and Feingold, G.: The scale problem in quantifying aerosol indirect effects,
721 *Atmos. Chem. Phys.*, 12, 1031–1049, 2012.

722 Medhaug, I., Stolpe, M. B., Fischer, E. M., and Knutti, R.: Reconciling controversies about the
723 ‘global warming hiatus,’ *Nature*, 545, 41–47, 2017.

724 Möller, D. *Chemistry of the Climate System*, pp. 331-334, Walter de Gruyter GmbH and Co.,
725 KG, Berlin/New York, 2010.

726 Monier, E., Scott, J. R., Sokolov, A. P., Forest, C. E., and Schlosser, C. A.: An integrated
727 assessment modeling framework for uncertainty studies in global and regional climate
728 change: The MIT IGSM-CAM (version 1.0), *Geosci. Mod. Dev.*, 6, 2063–2085, 2013.

729 Monier, E., Gao, X., Scott, J. R., Sokolov, A. P., and Schlosser, C. A.: A framework for
730 modeling uncertainty in regional climate change, *Clim. Change*, 131, 51–66, 2015.

731 Monier, E., Xu, L., and Snyder, R.: Uncertainty in future agro-climate projections in the United
732 States and benefits of greenhouse gas mitigation, *Environ. Res. Lett.*, 11, 055001, 2016.

733 Paltsev, S., Reilly, J. M., Jacoby, H. D., Eckaus, R. S., McFarland, J. R., Sarofim, M. C.,
734 Asadoorian, M. O., and Babiker, M. H.: The MIT emissions prediction and policy analysis
735 (EPPA) model: Version 4, Rep. 125, MIT Joint Program on the Sci. and Policy of Global
736 Change, 2005.

737 Pogson, M., and Smith, P.: Effect of spatial data resolution on uncertainty, *Environ. Model.*
738 *Softw.*, 63, 87–96, 2015.

739 Pyle, J. A., and Zavody, A. M.: The modelling problems associated with spatial averaging, *Q. J.*
740 *R. Meteorol. Soc.*, 116, 753–766, 1990.

741 Räisänen, J., and Ylhäisi, J. S.: How much should climate model output be smoothed in space?,
742 *J. Climate*, 24, 867–880, 2011.

743 Rasmussen, D. J., Fiore, A. M., Naik, V., Horowitz, L. W., McGinnis, S. J., and Schultz, M. G.:
744 Surface ozone-temperature relationships in the eastern US: A monthly climatology for
745 evaluating chemistry-climate models, *Atmos. Environ.*, 47, 142–153, 2012.

746 Rienecker, M. M., Suarez, M. J., Gelaro, R., Todling, R., Bacmeister, J., Liu, R., Bosilovich, M.
747 G., Schubert, S. D., Takacs, L., Kim, G-K, Bloom, S., Chen, J., Collins, D., Conaty, A., da

748 Silva, A., Gu, W., Joiner, J., Koster, R. D., Lucchesi, R., Molod, A., Owens, T., Pawson, S.,
749 Pegion, P., Redder, C. R., Reichle, R., Robertson, F. R., Ruddick, A. G., Sienkiewicz, M.,
750 and Woollen, J.: MERRA: NASA's Modern-Era Retrospective analysis for Research and
751 Applications, *J. Climate*, 24, 3624–3648, 2011.

752 Roberts, C. D., Palmer, M. D., McNeall, D., and Collins, M.: Quantifying the likelihood of a
753 continued hiatus in global warming, *Nat. Clim. Change*, 5, 337–342, 2015.

754 Schnell, J. L., and Prather, M. J.: Co-occurrence of extremes in surface ozone, particulate matter,
755 and temperature over eastern North America, *Proc. Natl. Acad. Sci. U.S.A.*, 114, 11, 2854-
756 2859, 2017.

757 Schurer, A. P., Hegerl, G. C., Mann, M. E., Tett, S. F. B., and Phipps, S. J.: Separating forced
758 from chaotic climate variability over the past millennium, *J. Climate*, 26, 6954–6973, 2013.

759 Searle, K. R., Chipperfield, M. P., Bekki, S., and Pyle, J. A.: The impact of spatial averaging on
760 calculated polar ozone loss: 2. Theoretical analysis, *J. Geophys. Res.*, 103, D19, 25409–
761 25416, 1998.

762 Sofen, E. D., Bowdalo, D., and Evans, M. J.: How to most effectively expand the global surface
763 ozone observing network, *Atmos. Chem. Phys.*, 16, 1445–1457, 2016.

764 Stevenson, D. S., Young, P. J., Naik, V., Lamarque, J. F., Shindell, D. T., Voulgarakis, A., Skeie,
765 R. B., Dalsøren, S. B., Myhre, G., Bernsten, T. K., Folberth, G. A., Rumbold, S. T., Collins,
766 W. J., MacKenzie, I. A., Doherty, R. M., Zeng, G., van Noije, T. P. C., Strunk, A.,
767 Bergmann, D., Cameron-Smith, P., Plummer, D. A., Strode, S. A., Horowitz, L., Lee, Y. H.,
768 Szopa, S., Sudo, K., Nagashima, T., Josse, B., Cionni, I., Righi, M., Eyring, V., Conley, A.,
769 Bowman, K. W., and Wild, O.: Tropospheric ozone changes, radiative forcing and
770 attribution to emissions in the Atmospheric Chemistry and Climate Model Intercomparison
771 Project (ACCMIP), *Atmos. Chem. Phys.*, 13, 3063–3085, 2013.

772 Strode, S. A., and Pawson, S.: Detection of carbon monoxide trends in the presence of
773 interannual variability, *J. Geophys. Res. Atmos.*, 118, 12257-12273, 2013.

774 Sun, J., Fu, J. S., Drake, J., Lamarque, J.-F., Tilmes, S., and Vitt, F.: Improvement of the
775 prediction of surface ozone concentration over conterminous U.S. by a computationally
776 efficient second-order Rosenbrock solver in CAM4-Chem, *J. Adv. Model Earth. Sy.*, 9,
777 482–500, 2017.

778 Tilmes, S., Lamarque, J.-F., Emmons, L. K., Conley, A., Schultz, M. G., Saunoy, M., Thouret,
779 V., Thompson, A. M., Oltmans, S. J., Johnson, B., and Tarasick, D.: Technical Note:
780 Ozonesonde climatology between 1995 and 2011 : description, evaluation and applications,
781 *Atmos. Chem. Phys.*, 12, 7475-7497, 2012.

782 Tilmes, S., Lamarque, J. F., Emmons, L. K., Kinnison, D. E., Ma, P. L., Liu, X., Ghan, S.,
783 Bardeen, C., Arnold, S., Deeter, M., Vitt, F., Ryerson, T., Elkins, J. W., Moore, F.,
784 Spackman, J. R., and Val Martin, M.: Description and evaluation of tropospheric chemistry
785 and aerosols in the Community Earth System Model (CESM1.2), *Geosci. Model Dev.*, 8,
786 1395–1426, 2015.

787 Tilmes, S., Lamarque, J. F., Emmons, L. K., Kinnison, D. E., Marsh, D., Garcia, R. R., Smith, A.
788 K., Neely, R. R., Conley, A., Vitt, F., Val Martin, M., Tanimoto, h., Simpson, I., Blake, D.
789 R., and Blake, N.: Representation of the Community Earth System Model (CESM1) CAM4-

790 chem within the Chemistry-Climate Model Initiative (CCMI), *Geosci. Model Dev.*, 9,
791 1853–1890, 2016.

792 Travis, K. R., Jacob, D. J., Fisher, J. A., Kim, P. S., Marais, E. A., Zhu, L., Yu, K., Miller, C. C.,
793 Yantosca, R. M., Sulprizio, M. P., Thompson, A. M., Wennberg, P. O., Crouse, J. D., St.
794 Clair, J. M., Cohen, R. C., Laughner, J. L., Dibb, J. E., Hall, S. R., Ullmann, K., Wolfe, G.
795 M., Pollack, I. B., Peischl, J., Neuman, J. A., and Zhou, X.: Why do models overestimate
796 surface ozone in the Southeast United States?, *Atmos. Chem. Phys.*, 16, 13561–13577,
797 2016.

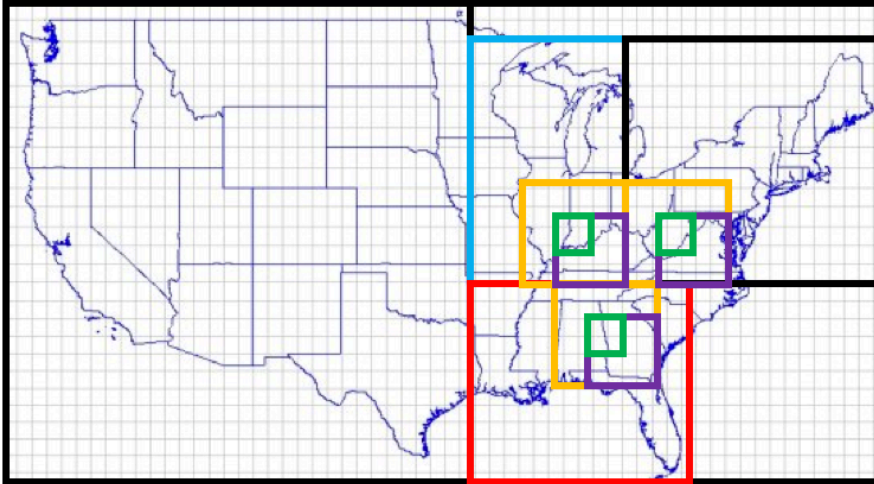
798 US EPA: National Ambient Air Quality Standards for Ozone: Final Rule. *Fed. Regist.* 80 (206),
799 65292-65468. 2015.

800 US EPA: CASTNET 2014 Annual Report Prepared by Environmental Engineering and
801 Measurement Services, Inc. for the U.S. Environmental Protection Agency, 2016.

802 Weatherhead, E. C., Stevermer, A. J., and Schwartz, B. E., Detecting environmental changes and
803 trends, *Physics and Chemistry of the Earth*, 27, 399-403, 2002.

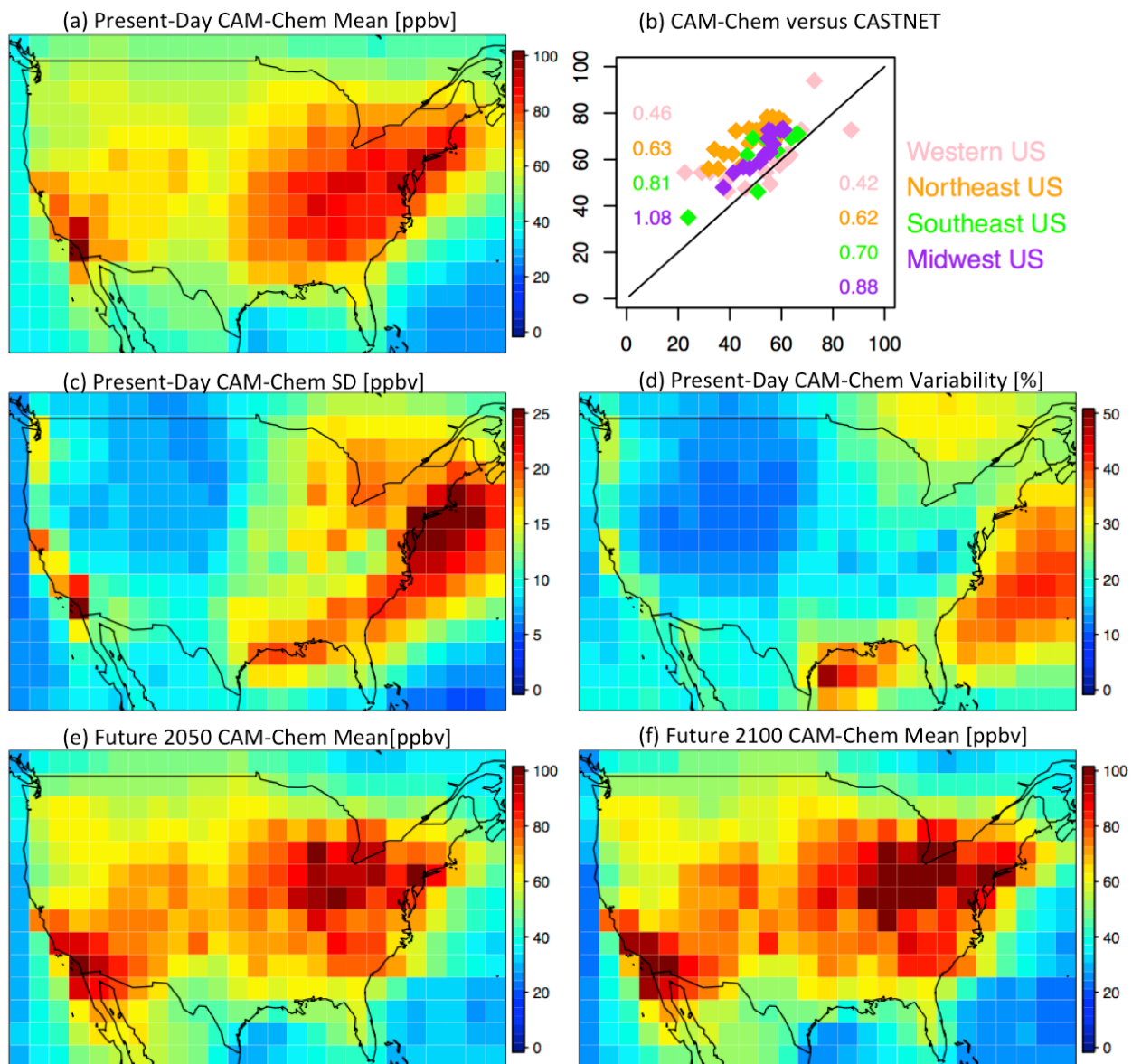
804 Wild, O., and Prather, M. J.: Global tropospheric ozone modeling: Quantifying errors due to grid
805 resolution, *J. Geophys. Res.*, 111, D11305, 2006.

806 Young, P. J., Archibald, A. T., Bowman, K. W., Lamarque, J.-F., Naik, V., Stevenson, D. S.,
807 Tilmes, S., Voulgarakis, A., Wild, O., Bergmann, D., Cameron-Smith, P., Cionni, I.,
808 Collins, W. J., Dalsøren, S. B., Doherty, R. M., Eyring, V., Faluvegi, G., Horowitz, L. W.,
809 Josse, B., Lee, Y. H., MacKenzie, I. A., Nagashima, T., Plummer, D. A., Righi, M.,
810 Rumbold, S. T., Skeie, R. B., Shindell, D. T., Strode, S. A., Sudo, K., Szopa, S., and Zeng,
811 G.: Pre-industrial to end 21st century projections of tropospheric ozone from the
812 Atmospheric Chemistry and Climate Model Intercomparison Project (ACCMIP), *Atmos.*
813 *Chem. Phys.*, 13, 2063–2090, 2013.

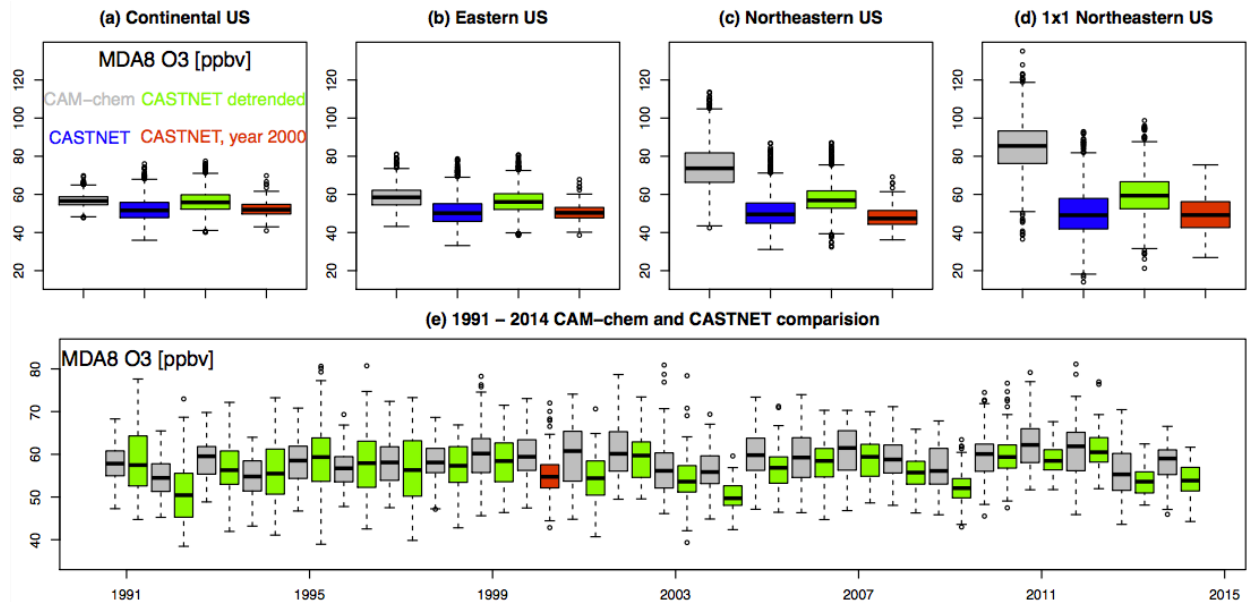


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Figure 1: Telescoping Spatial Regions included in this study. The largest scale we consider is the Continental US (outer border). We focus on the Eastern US, by subdividing into three subregions: the Midwest (blue), Northeast (black), and Southeast (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, and we include the rest in the Supplemental Material.

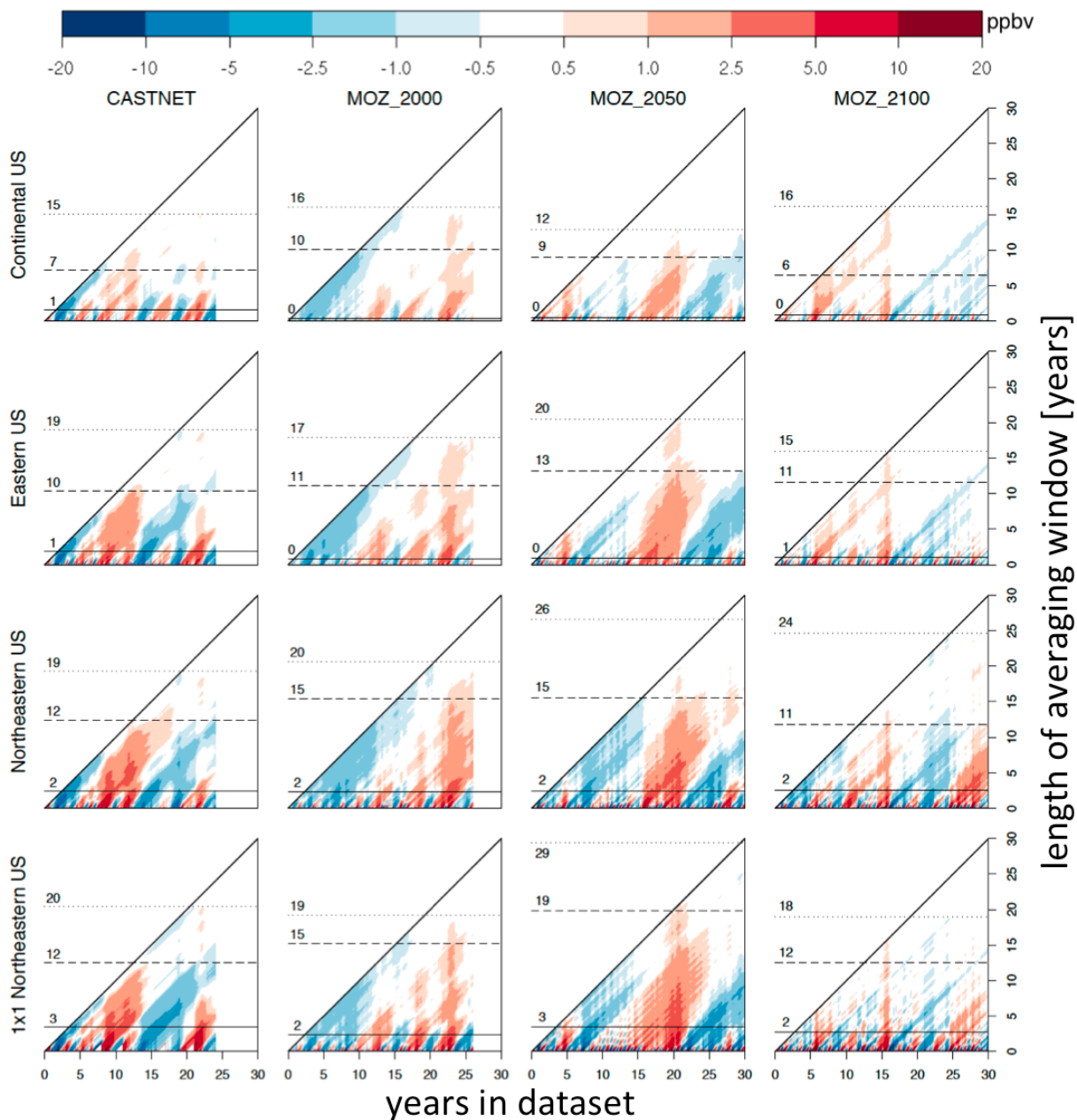


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 823 **Figure 2: Continental US surface maps of (a) present-day CAM-chem mean MDA8 O₃; (b) CAM-Chem (y-**
 824 **axis) comparison to CASTNET observations (x-axis) for the year 2000 (see Brown-Steiner et al. (in review)**
 825 **for additional comparisons); (c) present-day CAM- chem standard deviation of MDA8 O₃; (d) present-day**
 826 **CAM- chem variability (standard deviation divided by mean, as a percent); (e) future CAM- chem year 2050**
 827 **mean MDA8 O₃; and (f) future CAM- chem year-2100 mean MDA8 O₃. All model results are averaged over**
 828 **every JJA day in the time series, while the CASTNET results are only for the year 2000. The numbers in**
 829 **Figure 2b are slopes (left) and R² values (right).**
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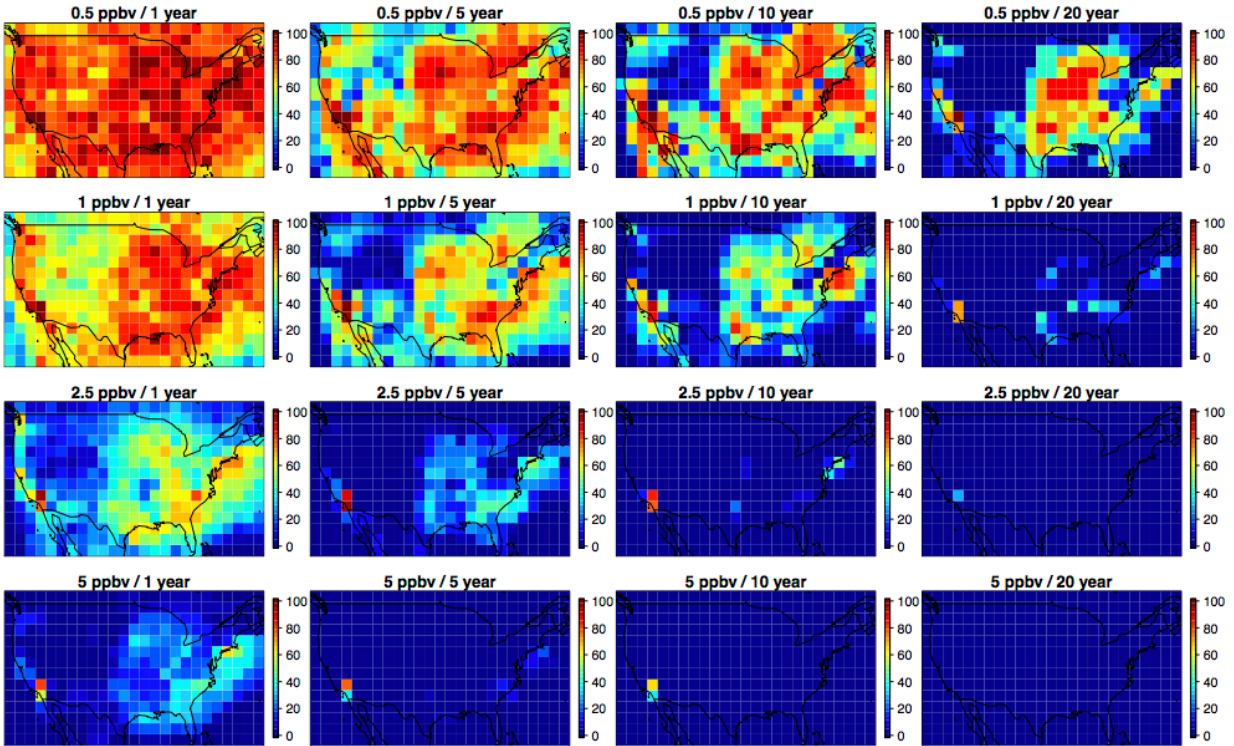


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Figure 3: (a-d): Boxplots for surface MDA8 O₃ for every summertime (JJA) day from 1991 – 2014 averaged over the Continental US, the Eastern US, the Northeastern US, and a single grid cell in the Northeastern US from CAM-chem (grey), CASTNET observations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

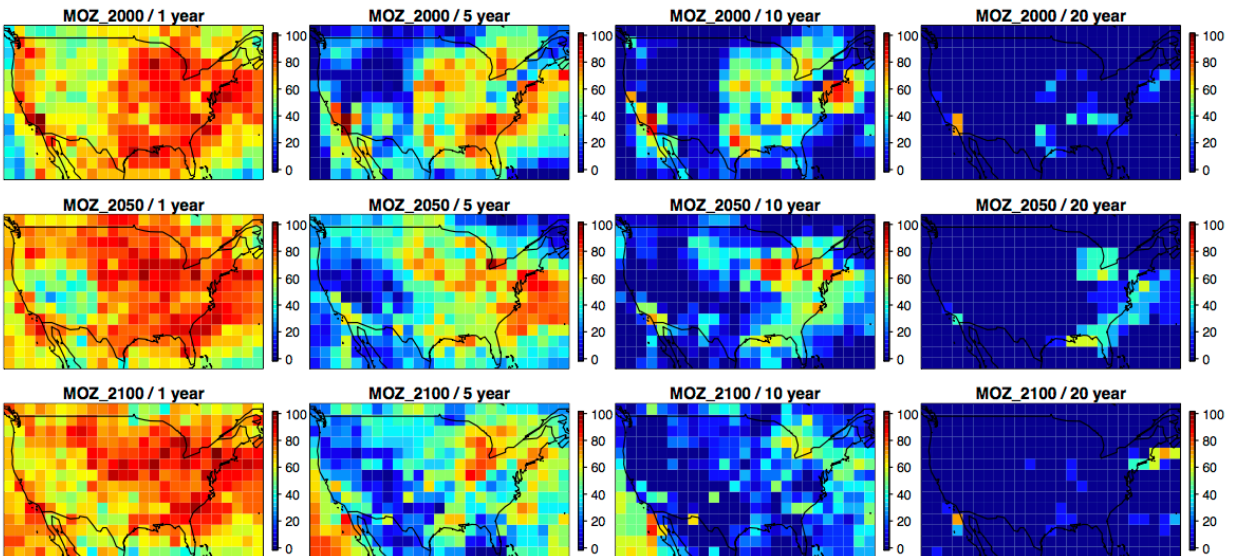


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 847 **Figure 4: Comparisons of the variability represented by the summertime MDA8 O₃ anomaly (from the**
 848 **long-term summertime mean) for the four datasets in this study (CASTNET, MOZ_2000, MOZ_2050,**
 849 **MOZ_2100, columns) averaged over the four telescoping regions (CUS, EUS, NEUS, NEUS 1x1, rows).**
 850 **In each panel, the horizontal axis is the number of years in the dataset (24 years (1991-2014) for**
 851 **CASTNET, 26 years (1990-2015) for MOZ_2000, and 30 years (2036-2065 and 2086-2115) for**
 852 **MOZ_2050 and MOZ_2100), and the vertical axis represents the length of the averaging window**
 853 **(ranging from 1 day (bottom row) up to the entire time series (top pixel)). Each pixel represents the**
 854 **estimate of the ozone anomaly for a given averaging window (vertical axis) ending at a given time**
 855 **(horizontal axis). Horizontal lines indicate the length of averaging window required to guarantee that**
 856 **the variability drops below thresholds of 5 ppbv (solid), 1 ppbv (dashed), and 0.5 ppbv (dotted).**
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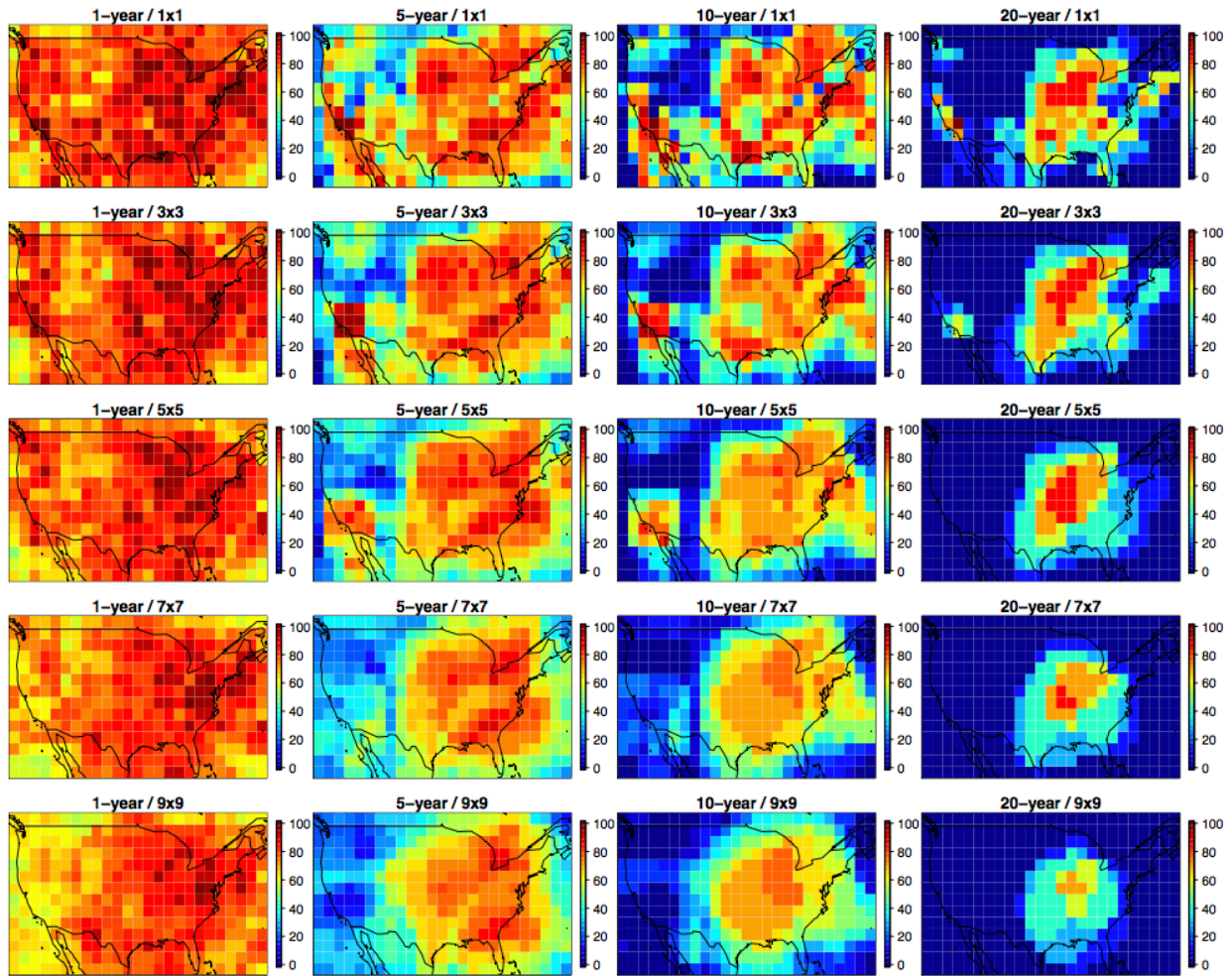
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Figure 5: Spatial Plots over the Continental US plotting the likelihood (%) that an estimate of ozone exceeds a given threshold due to meteorological variability (rows) at the grid-cell level when using different lengths of averaging windows (columns) for the present-day CESM simulation (MOZ_2000).

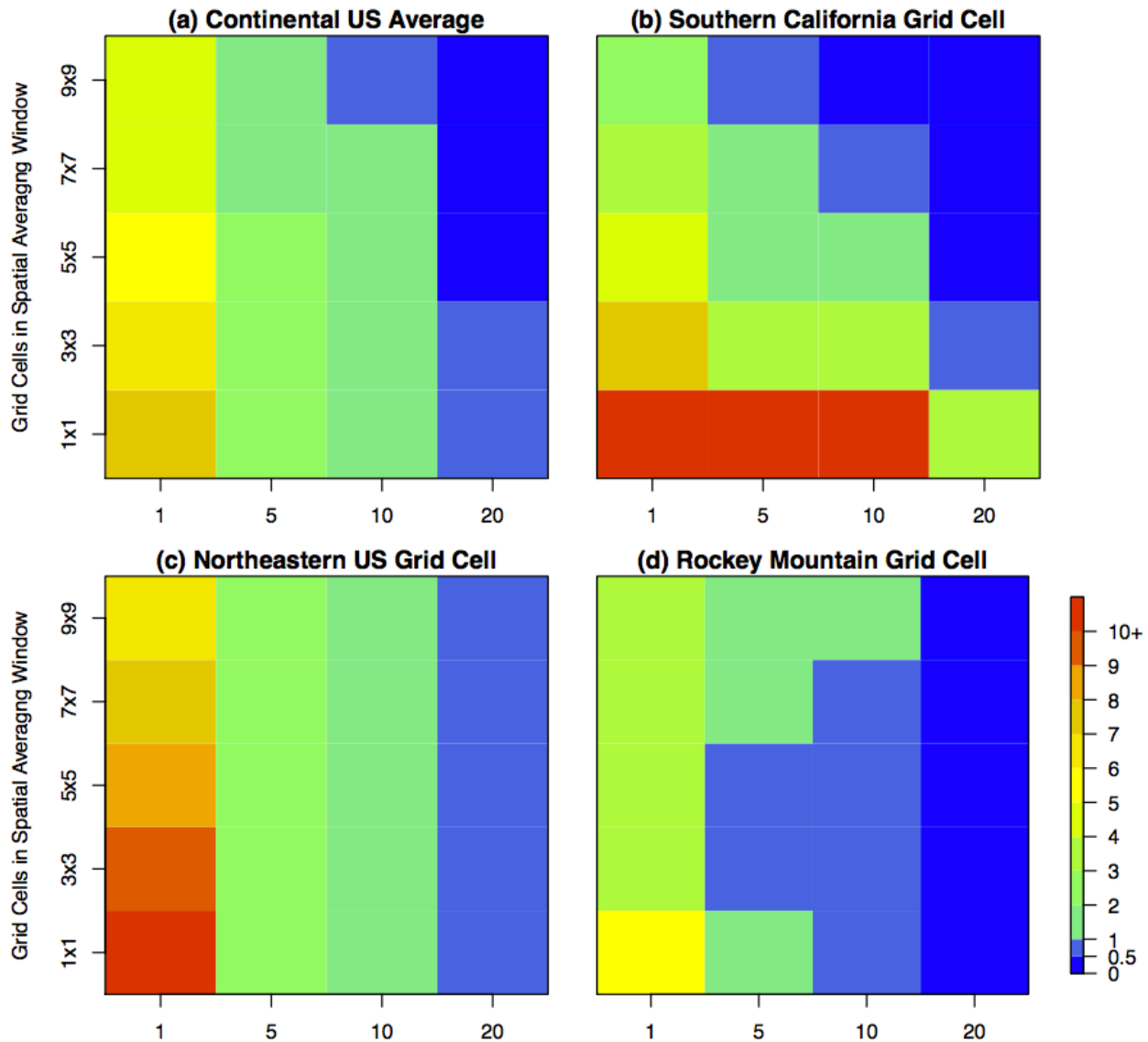


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Figure 6: As in Figure 5, but only the second row (1 ppbv threshold), for present-day CAM-chem (MOZ_2000), future CAM-chem 2050 (MOZ_2050), and future CAM-chem 2100 (MOZ_2100).



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 872 **Figure 7: Combined impact of temporal and spatial averaging on reducing ozone variability on the**
 873 **likelihood (%) of exceeding the 0.5 ppbv threshold (as in Figures 5, 6, and Supplemental Figure S3)**
 874 **for the present-day MOZ_2000 simulation. The top row is the same as in Figure 6, while the lower rows**
 875 **have averaged the values within a 3x3, 5x5, 7x7, and 9x9 grid box surrounding each individual grid cell.**
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879 **Figure 8: The maximum potential calculated MDA8 O₃ anomaly [ppbv] from the long-term mean for (a)**
 880 **the Continental US average and three individual grid cells taken from (b) Southern California,**
 881 **demonstrating effective temporal and spatial averaging, (c) the Northeast, where spatial averaging is**
 882 **ineffective, and (d) the Rocky Mountains, where spatial averaging initially reduces the anomaly, but**
 883 **then increases the anomaly as surrounding regions get included in the spatial average. The number of**
 884 **years included in the temporal averaging window increase along the x-axis and the number of grid cells**
 885 **included in the spatial averaging window increase along the y-axis. A full map of the Continental US can**
 886 **be found in the Supplemental Material (Figure S4). Note that the color scale is non-linear, and the color**
 887 **transitions are selected to match the thresholds established throughout this paper.**
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			CASTNET	MOZ_2000	MOZ_2050	MOZ_2100
Continental US	Mean	ppbv	52.4	56.7	56.8	57.4
	Standard Deviation	ppbv	5.04	3.08	3.54	3.73
	Variability	%	10%	5%	6%	7%
	Bias	ppbv		4.31		
Eastern US	Mean	ppbv	50.7	58.6	55.5	56.5
	Standard Deviation	ppbv	5.78	5.77	5.80	6.50
	Variability	%	11%	10%	10%	12%
	Bias	ppbv		7.91		
Northeastern US	Mean	ppbv	48.3	74.4	68.4	73.0
	Standard Deviation	ppbv	6.89	11.4	11.1	12.7
	Variability	%	14%	15%	16%	17%
	Bias	ppbv		26.1		
1x1 Northeastern US	Mean	ppbv	49.6	84.9	81.1	85.1
	Standard Deviation	ppbv	10.2	12.8	16.7	17.3
	Variability	%	21%	15%	21%	20%
	Bias	ppbv		35.3		

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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 simulation compared to the CASTNET data. Similar tables for the other regions in this study are included in the Supplemental Material.