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Complete List of Authors:	Jaffe, Daniel; University of Washington, School of Science, Technology, Engineering and Mathematics Wigder, Nicole; University of Washington Seattle, Atmospheric Sciences Downey, Nicole; Earth System Sciences, LLC, Pfister, Gabriel; National Center for Atmospheric Research, Boynard, Anne; National Center for Atmospheric Research, Reid, Stephen; Sonoma Technology, Inc,

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Impact of Wildfires on Ozone Exceptional Events in the Western U.S.

Daniel A. Jaffe^{1,2}, Nicole Wigder^{1,2}, Nicole Downey³, Gabriele Pfister⁴, Anne Boynard^{4,6},
Stephen B. Reid⁵

1. University of Washington-Bothell, School of Science and Technology, Bothell, WA 98011
2. University of Washington-Seattle, Department of Atmospheric Sciences, Seattle WA 98195
3. Earth System Sciences, LLC, Houston, TX 77005
4. National Center for Atmospheric Research, Boulder, CO 80307
5. Sonoma Technology, Inc, Petaluma, CA 94954
6. Now at UPMC Univ. Paris 06; Université Versailles St-Quentin; CNRS/INSU, LATMOS-IPSL, Paris, France

Abstract

Wildfires generate substantial emissions of nitrogen oxides (NO_x) and volatile organic compounds (VOCs). As such, wildfires contribute to elevated ozone (O₃) in the atmosphere. However, there is a large amount of variability in the emissions of O₃ precursors and the amount of O₃ produced between fires. There is also significant interannual variability as seen in median O₃, organic carbon and satellite derived carbon monoxide mixing ratios in the western U.S. To better understand O₃ produced from wildfires, we developed a statistical model that estimates the maximum daily 8-hour average (MDA8) O₃ as a function of several meteorological and temporal variables for three urban areas in the western U.S.: Salt Lake City, UT; Boise, ID; and Reno, NV. The model is developed using data from June-September 2000-2012. For these three locations, the statistical model can explain 60, 52 and 27% of the variability in daily MDA8. The Statistical Model Residual (SMR) can give information on additional sources of O₃ that are not explained by the usual meteorological pattern. Several possible O₃ sources can explain high SMR values on any given day.

We examine several cases with high SMR that are due to wildfire influence. The first case considered is for Reno in June 2008 when the MDA8 reached 82 ppbv. The wildfire influence for this episode is supported by PM concentrations, the known location of wildfires at the time and simulations with the Weather and Research Forecasting Model with Chemistry (WRF-Chem) which indicates transport to Reno from large fires burning in California. The contribution to the MDA8 in Reno from the California wildfires is estimated to be 26 ppbv, based on the SMR, and 60 ppbv, based on WRF-Chem. The WRF-Chem model also indicates an important role for peroxyacetyl nitrate (PAN) in producing O₃ during transport from the California wildfires. We hypothesize that enhancements in PAN due to wildfire emissions may lead to regional enhancements in O₃ during high fire years. The second case is for the Salt Lake City (SLC) region for August 2012. During this period the MDA8 reached 83 ppbv and the SMR suggests a wildfire contribution of 19 ppbv to the MDA8. The wildfire influence is supported by PM_{2.5} data, the known location of wildfires at the time, HYSPLIT dispersion modeling that indicates transport from fires in Idaho, and results from the CMAQ model that confirm the fire impacts. Concentrations of PM_{2.5} and O₃ are enhanced during this period, but overall there is a poor relationship between them, which is consistent with the complexities in the secondary production of O₃. A third case looks at high MDA8 in Boise ID during July 2012 and reaches similar conclusions. These results support the use of statistical modeling as a tool to quantify the influence from wildfires on urban O₃ concentrations.

I. Introduction

Wildfires generate substantial emissions of particulate matter (PM) and ozone (O₃) precursors (1, 2). They are also a major driver for interannual variations in summer air quality in the western U.S. for O₃ (3), PM (4) and black carbon aerosol (5). However, O₃ production from wildfires is highly variable. In a recent review on O₃ production, the majority of published studies identified a positive relationship between carbon monoxide (CO) and O₃ in wildfire plumes, which

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3 60 is a good indicator of O₃ production (6). The $\Delta\text{O}_3/\Delta\text{CO}$ ratio were on average 0.018, 0.15 and 0.22
4 61 ppbv ppbv⁻¹ for plumes aged 1-2 days, 2-5 days and ≥ 5 days, respectively, showing that O₃
5 62 production generally increased with age of the plume, but with large plume-to-plume variability.

6 63 O₃ production is complicated by a number of factors including highly variable emissions (1, 7),
7 64 aerosol effects (8, 9); complex meteorology and emissions of oxygenated VOCs (1, 10, 11) that can
8 65 result in rapid conversion of NO_x to PAN. If a wildfire plume mixes with urban emissions, more
9 66 rapid O₃ production than either the fire or urban emissions would generate by themselves is likely
10 67 (12, 13). The complexity of emissions, meteorology, radiation and aerosol effects make it very
11 68 difficult to accurately model O₃ photochemistry using standard Eulerian chemical models.

12 69 For O₃, the U.S. EPA uses the one-hour daily maximum and the maximum daily 8-hour average
13 70 O₃ (MDA8) for its regulatory standards. Across most of the western U.S., background O₃ is already
14 71 elevated due to the high elevations and exposure to the free troposphere (14-17). During high fire
15 72 years the distribution of MDA8 values across the western U.S. shifts by 5-7 ppbv, making
16 73 compliance with the O₃ standard much more challenging (14). For this reason, the EPA has
17 74 developed a policy on “exceptional events,” which can be defined as “Unusual or natural events that
18 75 affect air quality but are not reasonably controlled...” (See:

19 76 <http://www.epa.gov/ttn/analysis/exevents.htm>). Exceptional events can include natural dust
20 77 storms, transport from sources outside of North America, transport of air from the upper
21 78 troposphere or lower stratosphere or pollution impacts due to wildfires. To exclude data from
22 79 consideration, a region must submit a request to EPA that demonstrates, quantitatively, that the air
23 80 quality would have met the appropriate standard but for the exceptional event.

24 81 One approach to identify exceptional events is to use a regression model that calculates O₃
25 82 mixing ratios as a function of various meteorological variables. In most cases, daily maximum
26 83 temperature has been found to be the best predictor for peak or MDA8 O₃ at most sites (18). Other
27 84 studies have included a wide array of meteorological variables, such as temperature, cloud cover, or
28 85 humidity to improve the model fit (19, 20). In one study of 74 regions in the eastern U.S. (21), 10
29 86 different variables were considered in the model. Daily maximum temperature and daily average
30 87 relative humidity (RH) were found to be the most important predictors for MDA8 O₃. Using a
31 88 generalized additive model, the authors were able to predict the MDA8 values with an R² of 0.5-0.7
32 89 for most sites.

33 90 In this paper we demonstrate a statistical model that can predict the MDA8 O₃ for three
34 91 metropolitan regions in the western U.S.: Salt Lake City, Utah; Boise, Idaho; and Reno, Nevada.
35 92 For three cases we show that the high residuals from the statistical model are due to wildfire
36 93 influence and this can provide quantitative information on the O₃ contribution. The wildfire
37 94 influence is supported by a variety of indicators including the CMAQ and WRF-CHEM Eulerian
38 95 models. This work can help guide future analyses to quantify the influence of wildfires on O₃ in
39 96 urban areas, in support of exceptional event designations.

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II. Methods

We use data from a variety of sources for this analysis including the EPA's AQS sites, CASTNET O₃ measurements, NPS O₃ measurements, IMPROVE aerosol measurements, meteorological data from NCDC and satellite observations from instruments onboard the NASA A-train constellation. Gridded meteorological data are from the NCEP/NCAR Reanalysis dataset. For the Salt Lake City (SLC) urban area, we use the daily average MDA8 from all AQS sites in Salt Lake and Davis Counties, Utah. For Boise and Reno, we use the daily average of all MDA8 values in each metropolitan statistical area as defined by EPA. We also used surface meteorological data from the SLC, Boise and Reno airports. We used satellite data for CO mixing ratios derived for 800 mb from the AIRS instrument onboard the Aqua satellite and aerosol optical depth (AOD) from the MODIS instruments on board the Aqua and Terra satellites. When daily data from both instruments was available, we averaged the AOD values. For all analyses, we use data for the primary fire season, June-September 2000-2012, except there is no data for Boise in 2000. Further details and data sources are given in Table S1 in the Supplementary Information.

We used two different Eulerian models to help quantify the contribution to the MDA8 from wildfires, since neither model had results for the full time period of our study. For the 2012 wildfires, we used the Community Multiscale Air Quality (CMAQ) model (22) to quantify the influence of wildfires on O₃ concentrations. CMAQ is a peer-reviewed, state-of-the-science Eulerian photochemical model that is run daily as part of the experimental BlueSky Gateway air quality modeling system, which quantifies air pollutant concentrations resulting from wildfires and other emissions sources on a national scale (23). BlueSky Gateway combines meteorological predictions from the Pennsylvania State University/National Center for Atmospheric Research Mesoscale Model (MM5) version 3.7 with air quality predictions from CMAQ version 4.5.1 at a coarse (36 km) grid resolution, and aerosol tracers have been implemented in CMAQ to track primary PM_{2.5} generated by fires. For the summer of 2012, CMAQ was run with and without fire emissions to evaluate fire impacts on O₃ concentrations across the U.S.

For 2008, we performed simulations with the regional Weather Research and Forecasting Model with Chemistry (WRF-Chem version 3.2) (24) to quantify the influence from wildfires. The model covers the contiguous U.S. at a horizontal resolution of 24 km x 24 km and is run for the time period from 10 June – 10 July 2008. The anthropogenic emissions are obtained from the U.S. EPA 2005 National Emissions Inventory (NEI-2005). Biomass burning emissions are obtained from the Fire INventory from NCAR (FINN V1) (25) and are distributed in the model vertically following the online plume-rise module (26). The model is configured for the MOZART gas phase chemical scheme linked to the GOCART aerosol model (27). A more detailed description of the model configuration can be found in (28).

In addition to standard chemical tracers we include in these model runs a synthetic tracer that keeps track of O₃ that is due to NO_x emissions from fires. The O₃ tracer method or "XNO_x" method is described in detail in (29) and has been used in global models for identifying source contributions, such as for quantifying the O₃ budget (30-34) and here for the first time it has been applied in a regional model. The method tags emissions of NO and its resulting nitrogen-containing

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3 142 products (e.g. HNO₃, PAN, HNO₄, etc.) and follows them to the production of O₃. In addition to the
4 143 standard tagging method, we further conducted a simulation where we did not allow O₃ to be
5 144 produced through PAN decomposition from fires, in order to provide an estimate of the role of PAN
6 145 on O₃ production in fire plumes.
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8 146 9 147 **III. Results** 10 148

11 148
12 149 As noted above, there are large interannual variations in MDA8. To examine the
13 150 relationship between seasonal MDA8 and other parameters that are likely associated with wildfires,
14 151 we compared the SLC and regional MDA8 with organic carbon, AIRS CO and AOD. Figure 1
15 152 shows this comparison using summer median values for 2000-2012. In all cases, the median values
16 153 are significantly correlated ($p \leq 0.05$) with SLC median MDA8. Thus we conclude that wildfires in
17 154 the western U.S. are the primary driver to explain these large interannual variations and may cause a
18 155 significant shift in median MDA8, up to +8 ppbv in SLC. While this broad seasonal comparison
19 156 does not help identify wildfire impacts on individual days, it does demonstrate the challenge regions
20 157 have in meeting the O₃ air quality standard during high fire years. Also apparent in Figure 1 is a
21 158 downward trend in median MDA8 values in the SLC region. This is likely due to regional
22 159 emissions controls, as demonstrated by the downward trend in urban NO_x concentrations for the
23 160 SLC region. NO_x concentrations in the region have decreased approximately 5% per year since
24 161 2000 (see <http://www.epa.gov/airtrends/nitrogen.html>).
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26 159 **Statistical model development** 27 160 28 161 29 162 30 162 31 163

32 163
33 164 We used PASW Statistics software, version 18.0.3, to develop the statistical model. For
34 165 each location, we examined the multi-linear relationship (MLR) between the indicated variable and
35 166 the regional averaged MDA8 value for June-September. A large number of variables were
36 167 considered to identify the best model including surface variables (daily maximum temperature,
37 168 daily average wind speed) and upper air parameters (see Table S1). We identified the variables that
38 169 gave the strongest explanatory power (correlation) with the least multicollinearity, as this would
39 170 confuse the interpretation of the model results (35). For all locations, we found that daily
40 171 maximum temperature was the strongest predictor for MDA8. Figure S1 (SI section) shows a
41 172 scatter plot of MDA8 versus daily maximum temperature for SLC. However, other variables also
42 173 show a significant relationship with MDA8 and should be included in the model. The MLR model
43 174 fits an equation of the form:
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$$45 173 \text{MDA8} = a + bX_1 + cX_2 \dots + \text{residual}$$

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50 177 Where the MDA8 is the dependent variable, X₁, X₂, etc. are the independent predictors, a,b,c, etc.
51 178 are the coefficients and the residual is the unfit portion of the model. In this analysis we refer to
52 179 the residual as the statistical model residual (SMR). We examined the best form for each predictor
53 180 including linear, squared, quadratic, log, etc. For most variables a linear fit gave the best
54 181 performance, except for day of year, where a squared term yielded an improved fit.
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3 182 We also examined the performance of Generalized Linear Models (GLM) as a tool to build
4 183 the statistical models, but did not find a significant improvement in predictive ability. Therefore we
5 184 used the simpler MLR approach, which makes interpretation of the predictors more straightforward.

6
7 185 Table 1 summarizes the model predictors and Table 2 provides a summary of the MDA8 and
8 186 model residuals for each location. For SLC and Boise, the best fit model explained 60 and 52% of
9 187 the variance in MDA8 values. For these two regions we found that daily maximum temperature,
10 188 daily average surface wind speed, day of year², year and the 700 mb zonal wind component gave
11 189 the best fit with minimal multicollinearity. Inclusion of additional meteorological variables made
12 190 virtually no difference to the model fit. Expanding the analysis to include May data generally
13 191 reduced the performance of the model in all three locations. For Reno, the model explained less of
14 192 the total variance (27%) and the variables included only daily maximum temperature, day of year²
15 193 and the 700 mb zonal wind component. The most likely explanation for this is that SLC and Boise,
16 194 are more isolated cities, whereas Reno is adjacent to larger emissions sources in California. We
17 195 expect that a parameter that included a better measure of transport from California might explain
18 196 more of the variance in MDA8 values.

19 197 Because the MDA8 values have significant autocorrelation, inclusion of the previous day's
20 198 MDA8 value will improve the model fit. For SLC and Boise, this improvement was relatively
21 199 minor (R^2 values increased by 0.05 for each), whereas for Reno the model fit was improved
22 200 substantially (R^2 increased by 0.20). For most days, this had relatively little impact on the
23 201 calculated residual, except in a multi-day pollution event. For such an event, the residual for the
24 202 second and succeeding days was reduced if the previous day's MDA8 was included as a model
25 203 predictor. However, since our goal is to identify exceptional events, we feel that these should be
26 204 predicted from meteorological parameters alone, not the previous day's MDA8 values. Thus the
27 205 final statistical model used in this analysis does not include the previous day's MDA8 as a
28 206 predictor. As a result, the SMRs show significant autocorrelation. We examined the influence
29 207 that this autocorrelation has on the SLC model results as follows. We removed 80% of the data
30 208 points by selecting data on only every 5th day and then reran the model with the same predictors.
31 209 We found that this made very little difference in the overall R^2 of the model and all predictor
32 210 variables remained statistically significant. The residuals from this reduced data model (every 5th
33 211 day) no longer show significant autocorrelation. With the reduced dataset, the Durbin-Watson test
34 212 confirmed no autocorrelation at a significance level of $p=0.05$ (test statistic =1.93), demonstrating
35 213 that autocorrelation has minimal influence in the reduced data model. So, while autocorrelation is
36 214 present in the residuals for the full data model, we found it has little influence on the predictors or
37 215 the form of the model.

38 216 It is important to check the distribution of the residuals and the relationship with respect to
39 217 the original independent variables. A histogram of the SMR values for SLC is shown in Figure S2
40 218 and a plot of SMR versus daily maximum temperature is shown in Figure S3. Mean and standard
41 219 deviation (SD) for the SMRs for each location is shown in Table 2. At all locations, the SMR has a
42 220 mean of 0 and a SD that is smaller than the SD of the MDA8. The residuals are normally
43 221 distributed and show no pattern with respect to temperature. Similar results are found when the
44 222 residuals are plotted against the other independent variables in the model for all locations.

223 Interpretation of the Statistical Model Residual (SMR)

224 The SMR can yield information about days that have higher MDA8 than predicted from the
225 meteorological conditions. These days are then candidates to consider as exceptional events, but
226 further evidence is needed to identify the cause of high O₃. Possible causes for high SMR might
227 include:

- 228 1) Additional precursors from unusual sources within the region;
- 229 2) Additional O₃ produced from precursors emitted by wildfires;
- 230 3) Unusually large contribution of O₃ from the upper troposphere/lower stratosphere (UTLS);
- 231 4) Unusually large contribution from transport of O₃ and/or precursors from distant sources.

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233 Both transport from Asian sources and transport from the UTLS have been previously identified as
234 important sources of O₃ in the western U.S. (14, 15, 36-38). Wildfires have also been suggested as
235 important O₃ sources (6), especially in the western U.S. (3,34). To examine the utility of the SMR
236 as a tool to quantify the influence on specific days, we will focus here on three cases with high
237 SMR.

238 While 2008 was not an exceptional year over the entire western U.S., wildfires in California
239 burned approximately 1.5 million acres in 2008 compared with 0.7 million acres on average for the
240 state between 1997-2012 (data from the National Interagency Fire Center www.nifc.gov).

241 Exceedances of the hourly O₃ standard were reported at a number of sites in California and several
242 of these were considered “exceptional events” by the state (39). Figure 2 shows the modeled O₃ for
243 local afternoon on 24 June 2008. While the fires were located mainly in California, westerly winds
244 carried plumes into Nevada, with O₃ reaching up to ~100 ppbv. A tagged WRF-CHEM model run
245 indicates that between 40-60 ppbv of O₃ was contributed by the wildfires across a large section of
246 western Nevada. Figure 3 shows a time series of the measured MDA8 and modeled MDA8 in
247 Reno, NV for 20-30 June 2008. The figure also shows the WRF-CHEM fire contribution to the
248 MDA8 and the SMR. Both the measured and modeled MDA8 values peak on 24 June, as does the
249 wildfire contribution and the SMR. The MDA8 in Reno on 24 June was 80 ppbv and the SMR
250 suggests that 24 ppbv was due to the wildfire contribution. The calculated fire contribution using
251 WRF-CHEM is 60 ppbv, a much higher value than the SMR; however, we expect this to differ as it
252 shows the tagged contribution under the chemical regime of the fire plume, whereas the SMR is the
253 residual from what is typical for the meteorological conditions. The consistency in timing between
254 the WRF-CHEM and observed values and the WRF-CHEM wildfire contribution supports the use
255 of the SMR as an indicator of the magnitude of the wildfire contribution.

256 Figure 2c also shows that a large fraction of the O₃ that is produced from wildfire precursors
257 is due to NO_x that has been cycled through peroxyacetyl nitrate (PAN). Substantial production of
258 PAN in wildfires has been noted previously (10), and the WRF-CHEM results demonstrate the
259 importance of PAN in generating O₃ far downwind of the fire region. This may also explain part of
260 the reason O₃ and PM enhancements from some wildfires show little relationship (13) and suggests
261 that fire influences on O₃ can occur far downwind of the emission source, driven by PAN transport
262 and decomposition back to NO_x.

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3 263 The 2012 wildfire season was unusually strong across most of the western U.S. In total
4 264 more than 7 million acres burned in the western U.S. compared to approximately 4 million acres in
5 265 an average year. Unusually large areas burned in California, Oregon, Idaho, Nevada and Montana
6 266 in 2012. In late July and early August a large number of fires burned across northern California,
7 267 southeastern Oregon, northern Nevada and southern Idaho. Figure 4 shows the contribution to the
8 268 MDA8 due to wildfires for 12 August 2012 as calculated by the CMAQ model. Figure 5 shows the
9 269 observed and CMAQ-modeled MDA8 for 1-17 August 2012, as well as the CMAQ fire contribution
10 270 and the SMR. Over the period between 7-18 August, the average CMAQ wildfire contribution of 8
11 271 ppbv is very close to the average SMR of 9 ppbv. However, the CMAQ model tends to
12 272 underpredict peak MDA8 values in excess of 75 ppbv observed on 7, 8, and 12 August. On these
13 273 days, the CMAQ modeled wildfire contribution are also lower than the SMR values. Figure S4
14 274 shows the relationship between observed MDA8 and daily average PM_{2.5} for the SLC region during
15 275 the fire influenced period. For this period, the R² for this relationship is 0.17, whereas in the
16 276 CMAQ model it is 0.51.

17 277 For Boise Idaho, results from the statistical model are shown in Tables 1 and 2. Figure S5
18 278 shows the SMR and CMAQ fire contribution for Boise in July 2012. During this time period, large
19 279 fires burning in northern California, Oregon and Idaho were influencing air quality across the
20 280 western states. The MDA8 values in Boise peaked at 74 ppbv on 11 July 2012. During this time
21 281 period the CMAQ results showed a relatively weak correlation with the observed MDA8 values in
22 282 Boise (R² of 0.12) and the calculated fire contributions were much smaller than the SMR values (see
23 283 Figure S5).

24 284 These findings highlight sources of error associated with the BlueSky Gateway CMAQ
25 285 modeling, which include parameterizations used to solve the atmospheric momentum equations,
26 286 spatial grid cell resolution, and uncertainties associated with fire emissions estimates (23). As in
27 287 the Reno case, the timing and magnitude of the observations and the Eulerian model results support
28 288 the use of the SMR to quantitatively characterize the O₃ production due to wildfires.
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30 290 **IV. Discussion and summary**

31 291
32 292 Figure 1 demonstrates that wildfires can have a significant influence on MDA8 levels in
33 293 urban areas of the western U.S., but quantifying the daily impact is a challenge. Development of a
34 294 statistical model for O₃ is an important and useful exercise that can indicate the types of
35 295 meteorological conditions that are conducive to O₃ formation in a specific region. Outliers from
36 296 this model, called the SMR, can then indicate unusual sources of O₃ or unusual conditions that may
37 297 qualify as exceptional events per the EPA definitions. For cases where corroborating analyses point
38 298 to the influence of fire emissions on elevated O₃ concentrations, the SMR can provide an estimate
39 299 of this impact. This may then satisfy EPA's requirement of a quantitative demonstration that O₃
40 300 levels would not have exceeded the standard "but for the unusual" event. The statistical modeling
41 301 technique described in this paper is able to provide this estimate without requiring the resources and
42 302 expertise needed for complex Eulerian photochemical modeling of wildfire impacts.
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3 303 We have estimated the magnitude of the wildfire impact on MDA8 for three cases using the
4 304 SMR and two different Eulerian models (WRF-Chem and CMAQ). None of these calculations can
5 305 be considered exact, nor can they be considered identical. Each method gives an estimate of the
6 306 true wildfire contribution for the given case. The SMR value likely underestimates the true
7 307 impact. This is because the value is calculated as the outlier, and thus ignores any background or
8 308 average wildfire contribution that is embedded in the seasonal cycle or relationship with
9 309 temperature. For the CMAQ calculation, the wildfire contribution is calculated as the difference in
10 310 model runs with all wildfires emissions turned on/off. Using WRF-Chem, the contribution is
11 311 calculated by tagging each emission source and using this to quantify the O₃ production. The
12 312 WRF-Chem results also demonstrate an important role for PAN in redistributing primary wildfire
13 313 emissions and enhancing O₃ over a larger region. One consequence of PAN chemistry and complex
14 314 aerosol affects is that O₃ and aerosol enhancements associated with wildfires show minimal
15 315 correlation.
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Figure 1. Time series of summer (June-September) median MDA8 (ppbv, left axis) from Salt Lake City urban area. Also shown are summer median organic carbon (OC, $\mu\text{g}/\text{m}^3$, right axis) from three background IMPROVE sites (CRM, PND, GRB) and two satellite observations: CO and AOD. Units are ppbv (left axis) for SLC MDA8, $\text{ppbv} \times 0.5$ (left axis) for the AIRS CO data, $\mu\text{g m}^{-3}$ (right axis) for OC. For AOD, the value is multiplied by 4 and shown on the right axis. The correlation coefficient between the annual median SLC MDA8 and OC, AIRS CO and AOD are 0.75, 0.86 and 0.58 respectively.

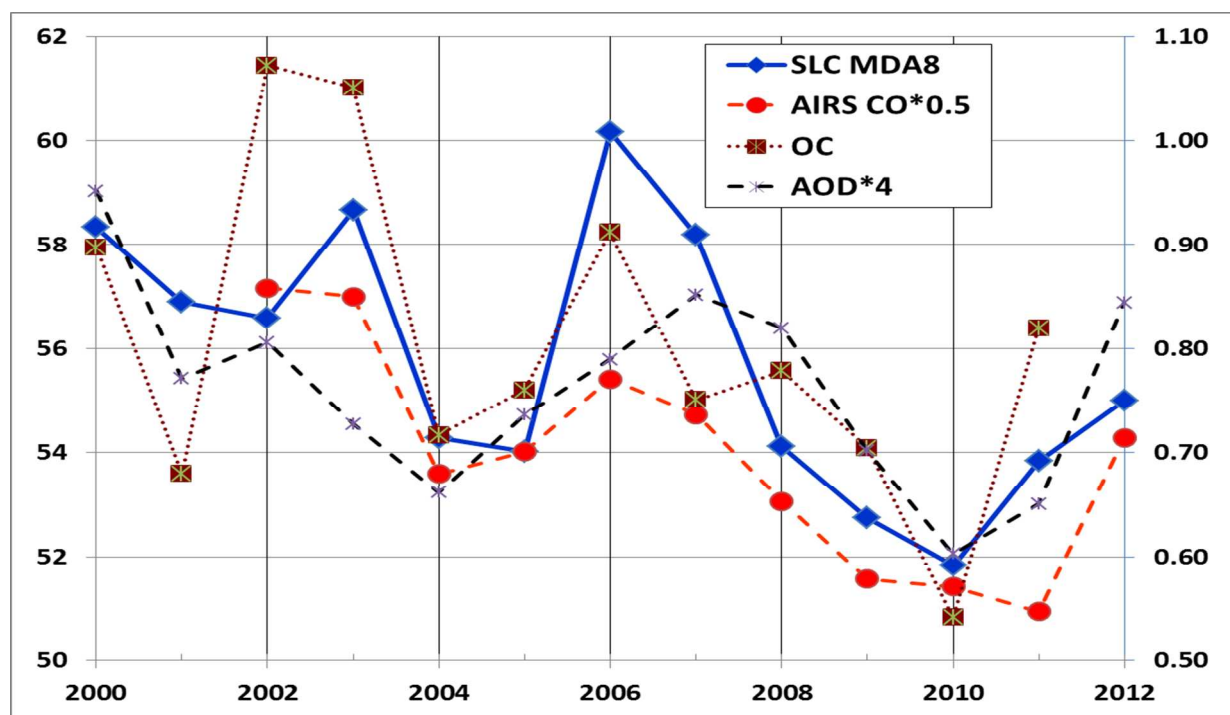


Figure 2: Surface O₃ from AQS sites for 24 June 2008 (circles) and WRF-Chem model results (ppbv, left panel), O₃ due to fires in WRF-Chem (ppbv, middle panel) and % of O₃ from fires where the NO_x cycled through PAN decomposition (% , right panel). Observations are an average for hours 15-17 local time, model results for 0 UTC (~16 local time).

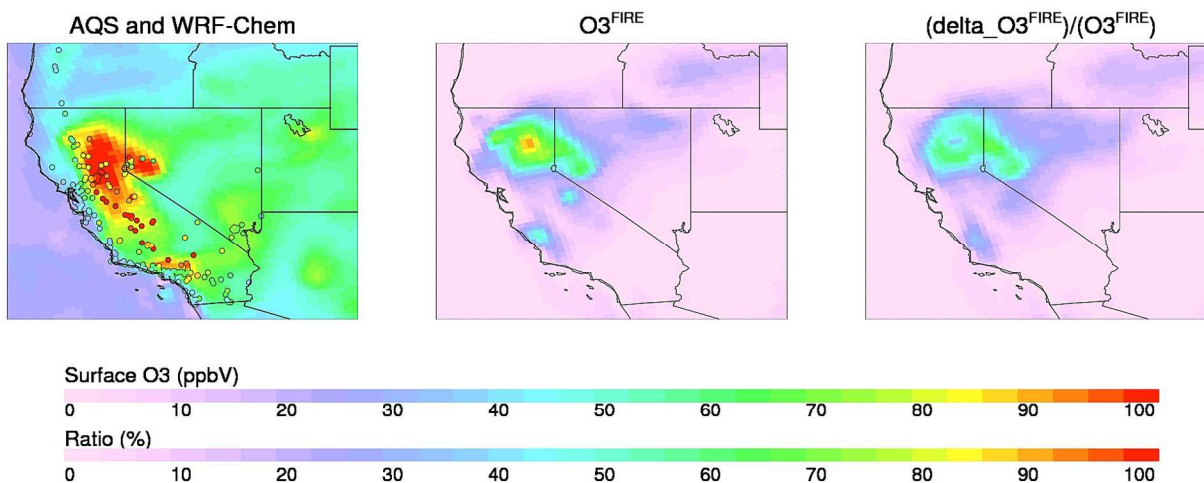
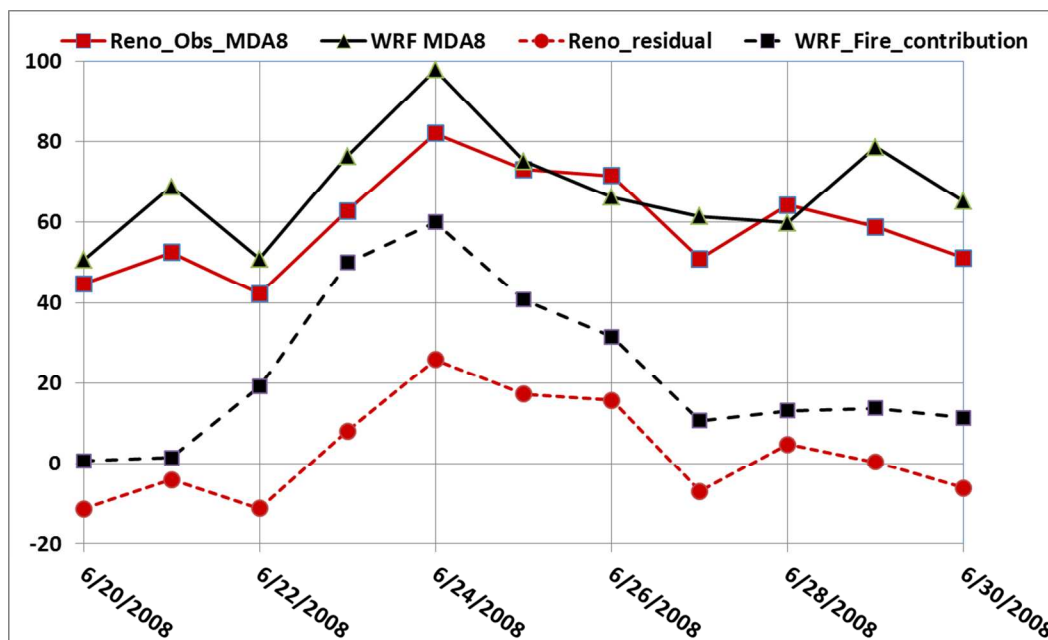
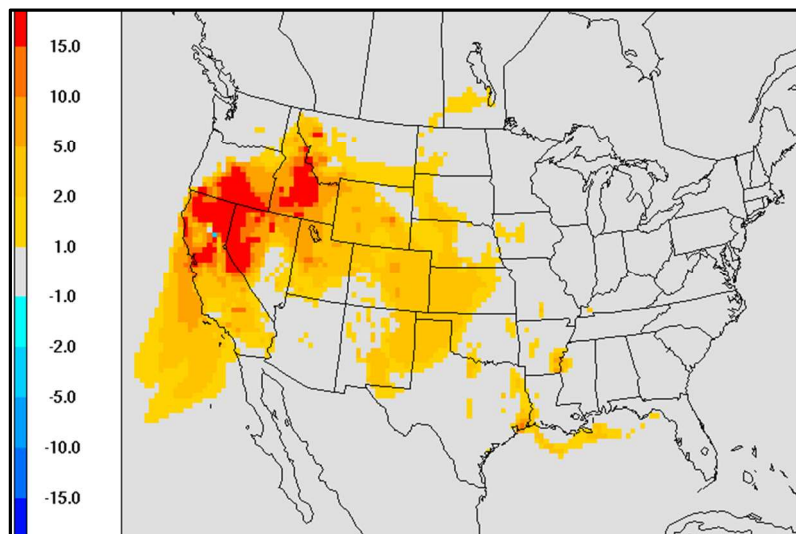


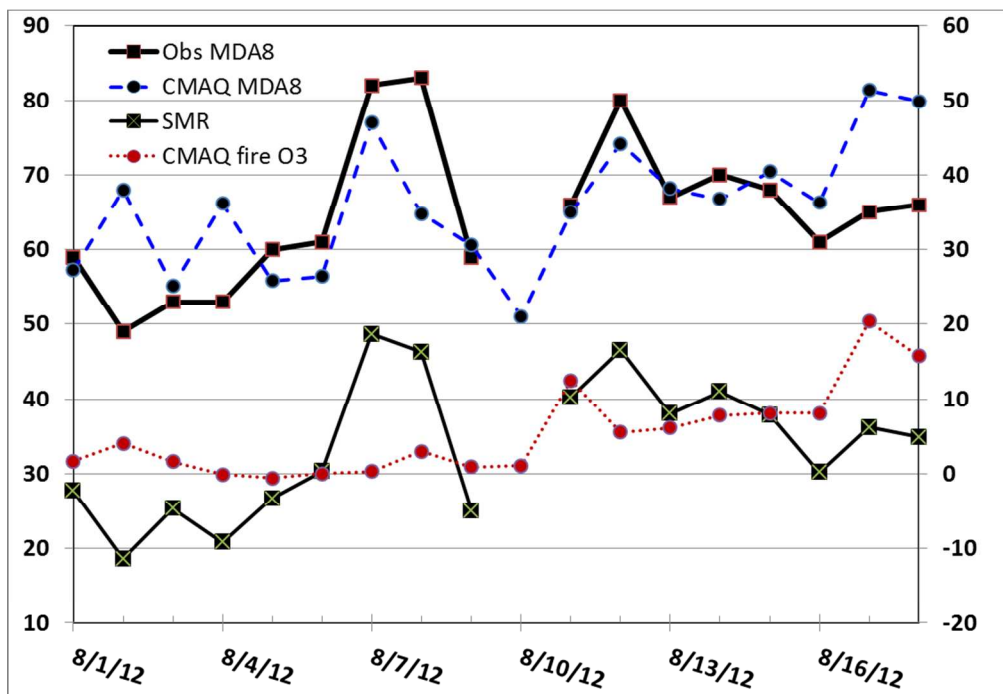
Figure 3. Observed MDA8 and residual from the statistical model for Reno. Calculated MDA8 from the WRF model and the model calculated contribution from fires.



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3 511 **Figure 4: Calculated contribution to MDA8 (ppbv) on 12 August 2012 due to wildfires in the**
4 512 **western U.S. This estimate is produced by running CMAQ with and without wildfire**
5 513 **emissions. The difference is assumed to be the wildfire contribution. For SLC, the observed**
6 514 **MDA8 is 80 ppbv. The modeled MDA8 is 74 ppbv, with a wildfire contribution of 5.5 ppbv.**
7 515 **By comparison, the SMR for this day is 19.6 ppbv.**
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520 **Figure 5. Observed and modeled (CMAQ) MDA8 for SLC for 1-18 Aug. 2012. Also shown is**
 521 **the contribution due to wildfires from the CMAQ model and the SMR. Observations and the**
 522 **SMR are not available for August 10, 2012.**
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3 528 **Table 1. Regression models for SLC, Boise and Reno MDA8 (ppbv). Overall R² for the SLC,**
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5 529 **Boise and Reno models are 0.60, 0.52 and 0.27, respectively. Units for MDA8, Tmax and wind**
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7 530 **speeds are ppbv, °C and m sec⁻¹, respectively. Inclusion of previous day's MDA8 increases**
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9 531 **the R² values to 0.65, 0.57 and 0.47, respectively. Note that while DOY is initially included as a**
10 532 **quadratic term, there is no difference in the final model fit by including it as a squared term.**
11 533

SLC Model		Unstandardized Coefficients		Standardized Coefficients	t	Sig.
		B	Std. Error	Beta		
	Constant	814	95.8		8.50	.000
	Daily max temp	1.29	.035	.634	36.7	.000
	Daily avg. wind spd.	-.197	.014	-.237	-13.7	.000
	Yr	-.388	.048	-.132	-8.14	.000
	DOY ²	-1.95E-4	1.24E-5	-.256	-15.6	.000
	700 mb zonal wind	-.615	.076	-.140	-8.06	.000

Boise Model		Unstandardized Coefficients		Standardized Coefficients	t	Sig.
		B	Std. Error	Beta		
	Constant	781	111		7.03	.000
	Daily max temp	1.02	.036	.562	28.1	.000
	Daily avg. wind spd.	-.087	.019	-.087	-4.59	.000
	Yr	-.372	.055	-.123	-6.74	.000
	DOY ²	-1.85E-4	1.34E-5	-.260	-13.8	.000
	700 mb zonal wind	-.521	.068	-.151	-7.66	.000

Reno Model		Unstandardized Coefficients		Standardized Coefficients	t	Sig.
		B	Std. Error	Beta		
	Constant	32.6	1.49		21.9	.000
	Daily max temp	.813	.039	.456	20.7	.000
	DOY²	-1.11E-4	1.27E-5	-.193	-8.76	.000
	700 mb zonal wind	-.189	.059	.071	-3.20	.001

Table 2. Statistical summary of SLC, Boise and Reno MDA8 and model residuals (June-Sept.)

	N	Minimum	Maximum	Mean	Std. Deviation
SLC MDA8	1585	19.7	101.5	55.8	11.0
SLC Resid.	1582	-24.7	34.3	0.0	7.0
Boise MDA8	1449	17.0	86.0	50.9	10.3
Boise Resid.	1449	-30.8	26.1	0.0	7.2
Reno MDA8	1586	24.8	82.0	52.8	8.4
Reno Resid.	1586	-27.7	25.9	0.0	7.2

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1 **Supporting information to accompany:**

2 Impact of Wildfires on Ozone Exceptional Events in the Western U.S.

3 by Daniel A. Jaffe, Nicole Wigder, Nicole Downey, Gabriele Pfister, Anne Boynard, and
4 Stephen B. Reid.

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8 **Table S1: Data sources and variables included in analysis**

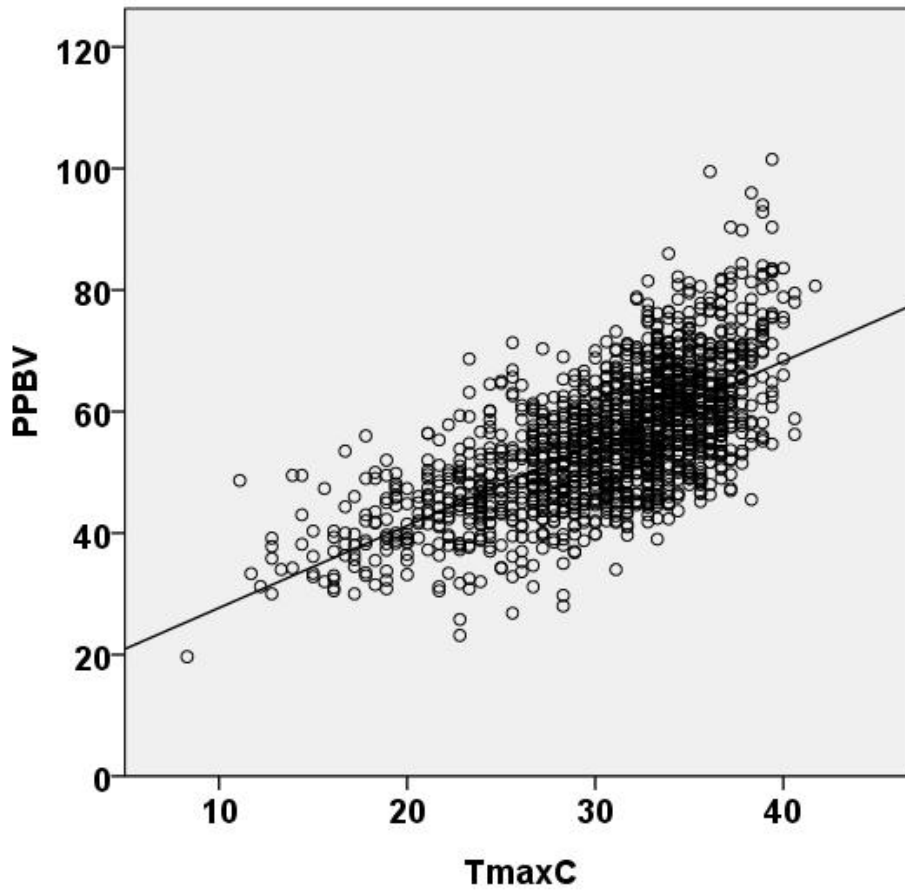
Data source	Data type/variables	Reference and/or data source
O ₃ data from EPA AQS sites	Daily MDA8, 2000-2012	http://www.epa.gov/airdata/
O ₃ data from CASTNET sites	Daily MDA8, 2000-2012	http://epa.gov/castnet
O ₃ data from one NPS site (Craters of the Moon)	Daily MDA8, 2000-2012	http://ard-request.air-resource.com/
IMPROVE aerosol data	PM _{2.5} organic carbon	Malm et al (2007) http://vista.cira.colostate.edu/improve/Default.htm
Satellite observations of CO from AIRS instrument ¹	800 mb retrieved value for 38-43° N -110-115° W	http://disc.sci.gsfc.nasa.gov/giovanni
Satellite observations of AOD from Aqua and Terra MODIS instrument	Average AOD from both instruments for 38-43° N, 110-115° W	http://disc.sci.gsfc.nasa.gov/giovanni
NOAA, National Climatic Data Center: Climate Data Online	Daily GHCND data: Daily maximum temperature and average daily wind speed (AWND). ²	http://www.ncdc.noaa.gov/cdo-web/
NCEP/NCAR Reanalysis dataset	Gridded time series of 700 mb zonal winds, geopotential heights and temperatures at 500mb, 700mb and surface, specific humidity at 850 mb.	http://www.esrl.noaa.gov/psd/data/timeseries/daily/

9 ¹Analyses and visualizations used in this study were produced with the Giovanni online data
10 system, developed and maintained by the NASA GES DISC.

11 ²For about 11% of the data record, the AWND data was not available. For these days we used a
12 derived a value based on the daily max 2-minute wind speed (WSF2) and a non-linear fitting
13 procedure. We found a good correlation between AWND and WSF2 (R=0.82 and 0.85) which
14 allowed us to use this procedure.

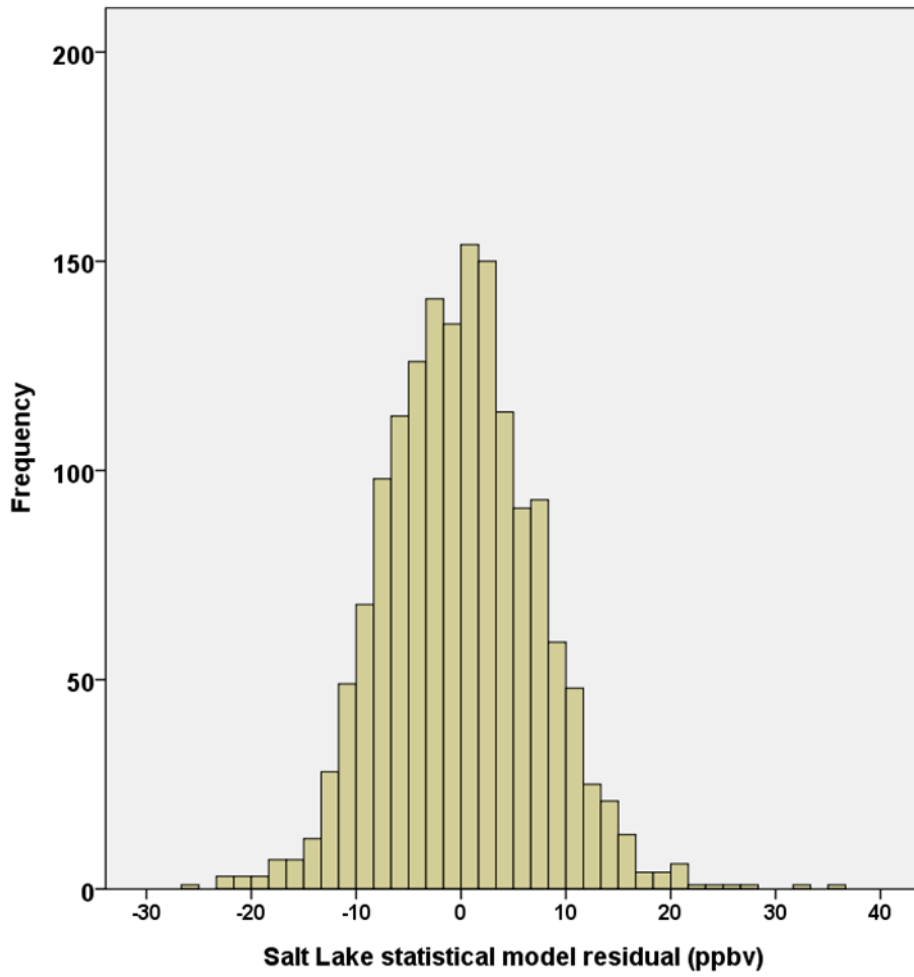
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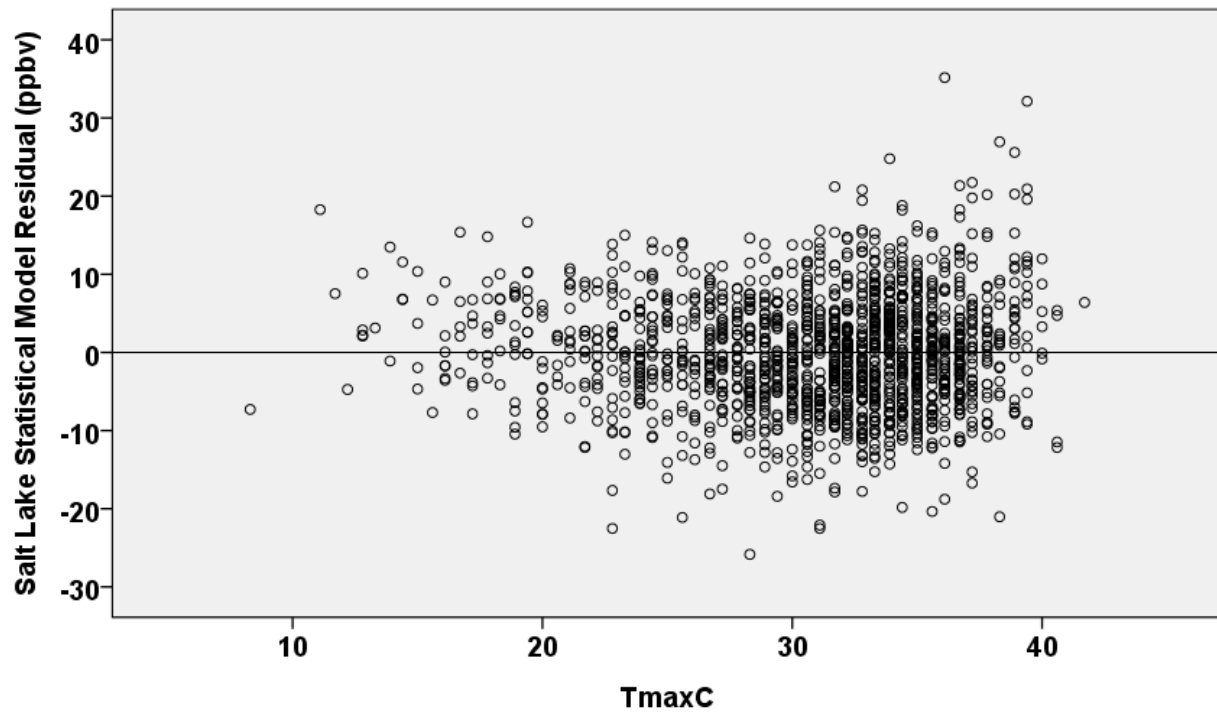


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23 **Figure S1: Plot of MDA8 O₃ vs. daily maximum temperature for AQS sites in Salt Lake and**
24 **Davis counties, Utah. The least squares regression has an R² of 0.44.**

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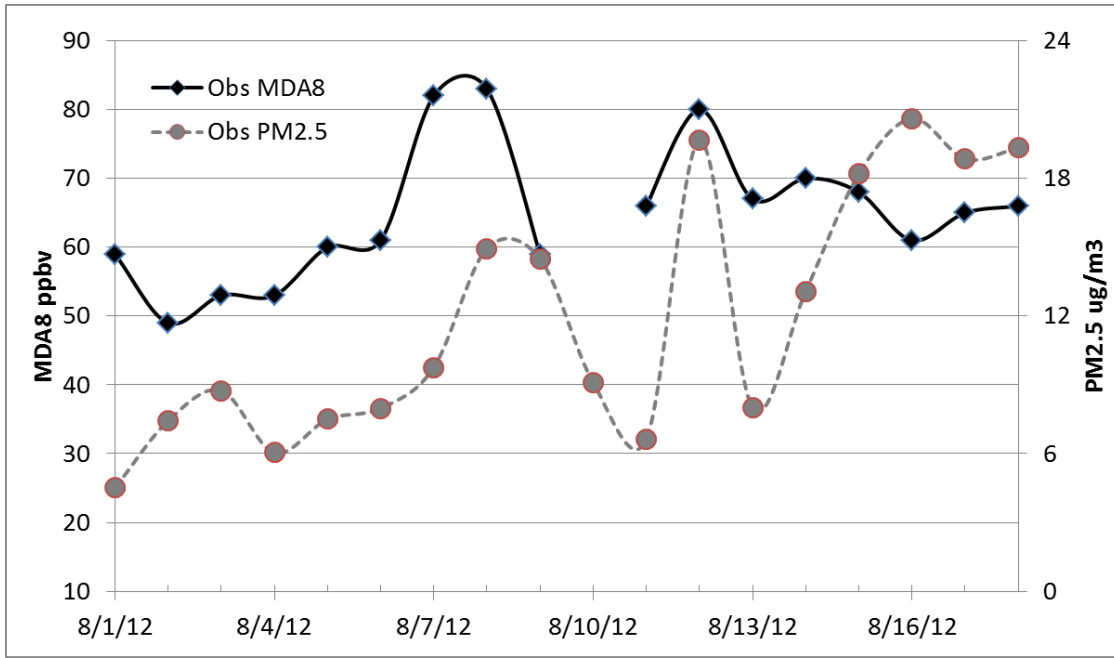


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34 **Figure S2: Histogram of the statistical model residuals for the Salt Lake City model. The**
35 **mean and standard deviation are 0.0 and 7.2 ppbv, respectively.**
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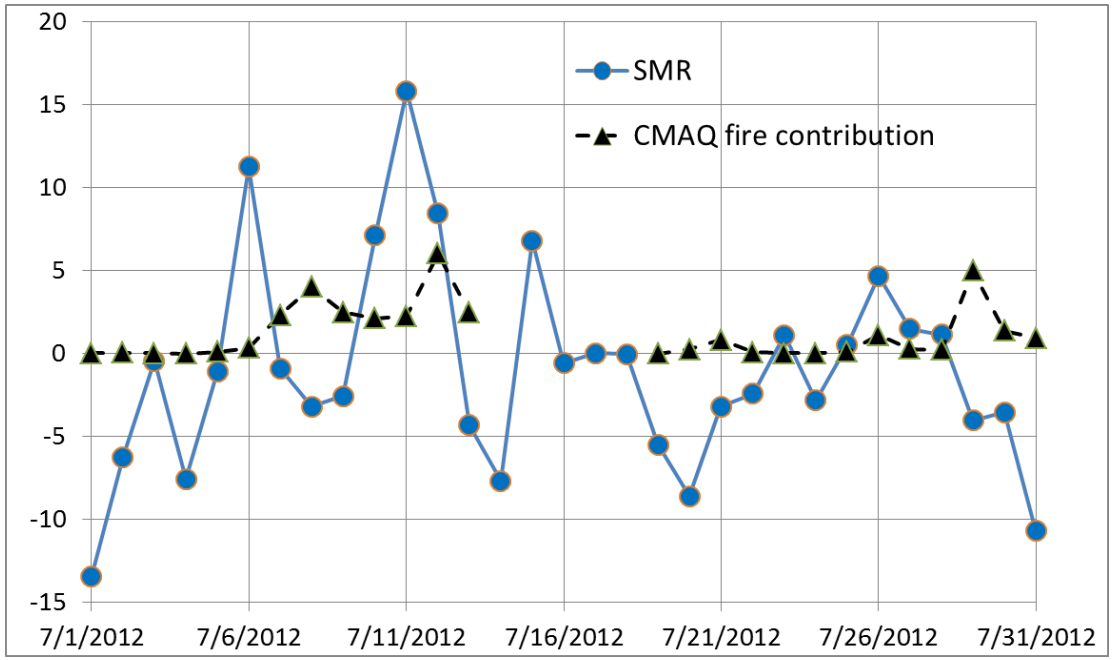
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Figure S3: Plot of SMR for SLC vs. daily maximum temperature. A horizontal line at 0.0 ppbv is shown for reference. The residuals show no relationship with daily max temperature.



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Figure S4. Observed MDA8 and daily average PM_{2.5} for the SLC monitoring sites during the fire influenced period. For this period the R² between the two observations is 0.27.



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58 **Figure S5. SMR and CMAQ fire contribution for Boise, ID in July 2012 in ppbv.**

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