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# Monitoring high-ozone events in the US Intermountain West using TEMPO geostationary satellite observations

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a value in the range of 60–70 ppbv (EPA, 2012). Ozone concentrations in this range are frequently observed at high-elevation sites in the western US with minimal local pollution influence (Lefohn et al., 2001). Although ozone levels have been decreasing over the eastern US for the past two decades due to emissions controls, there has been no such decrease in the West except for California (Cooper et al., 2012). Free tropospheric ozone at 3–8 km altitude over the western US has been increasing by 0.41 ppbvyr<sup>-1</sup> during the past two decades (Cooper et al., 2012) and this could affect background surface concentrations in the West (Zhang et al., 2008). There has been great interest in using satellite observations of ozone and related species to monitor and attribute background surface ozone (Lin et al., 2012a; Fu et al., 2013). This capability has been limited so far by the sparseness of satellite data and low sensitivity to the surface. All satellite measurements so far have been from low Earth orbit (LEO). Here we show that multispectral measurements from the NASA Tropospheric Emissions: Monitoring of Pollution (TEMPO) geostationary satellite mission over North America, scheduled for launch in 2018–2019, can provide a powerful ozone monitoring resource to complement surface sites, and can help to identify NAAQS exceedances caused by elevated background.

The North American background is defined by the EPA as the surface ozone concentration that would be present over the US in the absence of North American anthropogenic emissions. It represents the achievable benefits from domestic emissions control policies (including agreements with Canada and Mexico). It is particularly high in the Intermountain West, extending between the Sierra Nevada/Cascades to the west and the Rocky Mountains on the east, due to high elevation and arid terrain (Zhang et al., 2011). Subsidence of high-ozone air from the free troposphere can cause surface ozone concentrations in that region to approach or exceed the NAAQS (Reid et al., 2008). This is not an issue in the eastern US because of lower elevation, forest cover, and high moisture (Fiore et al., 2002).

Several chemical transport models (CTMs) and one chemistry-climate model (CCM) have been used to estimate the North American background including GEOS-Chem

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(Fiore et al., 2003; Zhang et al., 2011), GFDL AM3 CCM (Lin et al., 2012a, b), CMAQ (Mueller and Mallard 2011), and CAMx (Emery et al., 2012). Values average 30–50 ppbv in spring and summer over the Intermountain West with events exceeding 60 ppbv. However, there are large differences between models reflecting variable contributions from the stratosphere (Lin et al., 2012b), lightning (Kaynak et al., 2008; Zhang et al., 2011), and wildfires (Mueller and Mallard, 2011; Zhang et al., 2011; Jaffe and Wigder, 2012; Singh et al., 2012).

Background effects on surface ozone air quality are important to diagnose, as NAAQS exceedances can be dismissed as exceptional events if shown to be not reasonably controllable by local governances (EPA, 2013). Monitoring of ozone in the Intermountain West is mostly performed at urban stations designed to observe local pollution and not background influences. There is a limited network of Clean Air Status and Trends Network (CASTNet; [www.epa.gov/castnet/](http://www.epa.gov/castnet/)) sites located at national parks and other remote locations, and these have been used extensively to estimate background ozone and evaluate models (Fiore et al., 2002; Zhang et al., 2011; Lin et al., 2012b; Cooper et al., 2012). Langford et al. (2009) demonstrated that transport of stratospheric air contributed to surface one-minute average ozone concentrations in excess of 100 ppbv in Colorado in 1999. Analysis of ozonesonde and lidar measurements by Lin et al. (2012b) indicates thirteen stratospheric intrusions in spring 2010 leading to observed maximum daily 8 h average (MDA8) ozone of 70–86 ppbv at surface sites. Yates et al. (2013) similarly demonstrated a stratospheric origin for a NAAQS exceedance in Wyoming in June 2012 by using a combination of 3-D modeling, aircraft observations, LEO satellite data, and geostationary weather satellites. But the current air quality observing system is very limited in its ability to (1) monitor ozone at sites prone to high background, and (2) diagnose the origin of high-ozone events at these sites.

Geostationary satellites are a promising tool to address this limitation (Fishman et al., 2012). These satellites orbit around the Earth with a 24 h period in an equatorial plane, thus continuously staring at the same scenes. Depending on the observing strategy,

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they may provide hourly ozone data over a continental domain, while a LEO satellite may offer at best a 1 day return time. A global constellation of geostationary satellite missions targeted at air quality is planned to launch in 2018–2019 including TEMPO over North America (Chance et al., 2012), SENTINEL-4 over Europe (Ingmann et al., 2012), and GEMS over East Asia (Bak et al., 2013).

TEMPO will measure backscattered solar radiation in the 290–750 nm range, including the ultraviolet (UV) and visible Chappuis (Vis) ozone bands (Chance et al., 1997; Liu et al., 2005). Observation in the weak Chappuis bands takes advantage of the relative transparency of the atmosphere in the Vis to achieve sensitivity to near-surface ozone. This UV+Vis multispectral combination for ozone observation has not been used from space before. A theoretical study by Natraj et al. (2011) indicates that it should provide sensitivity to the lower troposphere. An observing system simulation experiment (OSSE) by Zoogman et al. (2011) shows that a UV+Vis instrument in geostationary orbit could provide useful constraints on surface ozone through data assimilation.

Here we conduct an OSSE to quantify the potential of geostationary ozone measurements from TEMPO to improve monitoring of ozone NAAQS exceedances in the Intermountain West and the role of background ozone in causing these exceedances. Our goal is to inform the TEMPO observing strategy and develop methods for exploitation of TEMPO data. OSSEs have previously informed mission planning for geostationary observations of atmospheric composition (Edwards et al., 2009; Timmermans et al., 2009; Zoogman et al., 2011, 2013; Claeys et al., 2011). An important feature of our work here is the inclusion of surface network and LEO satellite observations in the data assimilation system to properly quantify the added benefit of TEMPO observations. In this paper we describe our OSSE system, detail the method for assimilating surface measurements in conjunction with satellite observations, and analyze the potential of TEMPO for ozone monitoring and assessment.



et al., 2012b). This is important because the “true” model should reproduce the characteristics of the observations relevant to the OSSE. Lin et al. (2012a, b) used GFDL AM3 to investigate the effect of Asian transport and stratospheric intrusions on surface ozone in the Intermountain West during April–June 2010, and they quantified the ozone background through a sensitivity simulation with North American anthropogenic sources shut off. Here we use 3 hly concentrations archived from their standard simulation to provide the global 3-D ozone fields of the “true” atmosphere.

Our forward model for data assimilation is the GEOS-Chem CTM (Bey et al., 2001; <http://www.geos-chem.org>) The version used here (v8-02-03) was previously described by Zhang et al. (2011) in a study of background ozone influence on the Intermountain West during 2006–2008. It covers the North America domain with  $1/2^\circ \times 2/3^\circ$  horizontal resolution ( $10^\circ \text{ N}–60^\circ \text{ N}$ ,  $140^\circ \text{ W}–40^\circ \text{ W}$ ), nested within a global domain with  $2^\circ \times 2.5^\circ$  horizontal resolution. GEOS-Chem and GFDL AM3 have completely separate development heritages and use different driving meteorological fields, chemical mechanisms, and emission inventories. This independence between the two models used in the OSSE is important for a rigorous assessment (Arnold and Dey, 1986). The horizontal resolution of both models ( $\sim 50 \text{ km}$ ) is adequate for characterization of background ozone.

Figure 1 shows the maximum daily average 8 h (MDA8) ozone concentrations in surface air for each model, averaged over April–June 2010. GFDL AM3 has higher ozone concentrations than GEOS-Chem over the US as a whole and over the Intermountain West (bordered region) in particular. Zhang et al. (2011) previously showed that GEOS-Chem can reproduce ozone concentrations in the Intermountain West up to 70 ppbv with relatively little error, but cannot reproduce exceptional events of higher concentrations. GFDL AM3 is biased high in the mean but better simulates high-ozone events (Lin et al., 2012b).

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defines the sensitivity of  $\mathbf{x}'_p$  to  $\mathbf{x}_p$  ( $\mathbf{A} = \partial \mathbf{x}'_p / \partial \mathbf{x}_p$ ):

$$\mathbf{x}'_p = \mathbf{x}_s + \mathbf{A}(\mathbf{x}_p - \mathbf{x}_s) + \varepsilon \quad (1)$$

where  $\varepsilon$  is the instrument noise vector and  $\mathbf{x}_s$  is an independent a priori ozone profile used to regularize the retrieval.

Figure 2 shows typical clear-sky averaging kernel matrices for UV+Vis and TIR retrievals of tropospheric ozone taken from the Natraj et al. (2011) theoretical study. Also shown are the degrees of freedom for signal (DOFS) below given pressure levels. The DOFS are the number of independent pieces of information in the vertical provided by the retrieval, as determined from the corresponding trace of the averaging kernel matrix.

We generate synthetic TEMPO geostationary observations from the GFDL AM3 “true” atmosphere by sampling daytime vertical profiles over land in the North American domain (140–40° W, 10–70° N) with the averaging kernel matrix given in Fig. 2. TEMPO observations over the ocean are not included as the planned field of regard for the mission includes very little ocean and because the ocean surface is too dark for Vis retrievals. We similarly generate synthetic LEO IASI-3 observations over the North American domain twice a day (local noon and midnight) with the averaging kernel matrix given in Fig. 2. We omit scenes with cloud fraction > 0.3 (as given by the GEOS meteorology). We assume fixed averaging kernel matrices, acknowledging that in practice there is significant variability (Worden et al., 2013). Gaussian noise is added to the synthetic observations following Natraj et al. (2011) to simulate the random error associated with the spectral measurement. The noise from the TEMPO instrument (footprint of 4 km × 8 km) is reduced by the square root of the number of observations averaged over each GEOS-Chem grid square (~ 50 km × 50 km) in the data assimilation process. Since the TEMPO measurements are spatially dense we assume zero representation error during assimilation. IASI measurements have footprint diameters of 12–40 km with centers spaced 25–80 km apart (August et al., 2012); no reduction of the random error is applied to the LEO observations.

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## 2.3 Assimilation of surface and satellite measurements

The goal of our data assimilation system is to optimize an  $n$ -element state vector ( $\mathbf{x}$ ) of 3-D tropospheric ozone concentrations over the North American domain of GEOS-Chem, using surface and satellite observations to correct the GEOS-Chem simulation at successive time steps. CASTNet and TEMPO data are assimilated at discrete 3 h time steps, and LEO data are assimilated at 12 h time steps. We use a Kalman filter, as previously applied to ozone data assimilation by Khattatov et al. (2000), Parrington et al. (2008), and Zoogman et al. (2011). At each time step, we calculate an optimal estimate  $\hat{\mathbf{x}}$  of the true ozone concentrations  $\mathbf{x}$  as a weighted average of the model forecast  $\mathbf{x}_a$  (with corresponding error vector  $\boldsymbol{\varepsilon}_a$  relative to the true concentrations) and the observations  $\mathbf{x}'$  (with observational error  $\boldsymbol{\varepsilon}'$  and with  $\mathbf{x}'$  set to  $\mathbf{x}_a$  where there are no observations). The observational error includes both the instrument noise  $\boldsymbol{\varepsilon}$  and (for surface sites) the previously defined representation error. The errors are characterized by error covariance matrices  $\mathbf{S}_a = E[\boldsymbol{\varepsilon}_a \boldsymbol{\varepsilon}_a^T]$  and  $\mathbf{S}_\varepsilon = E[\boldsymbol{\varepsilon}' \boldsymbol{\varepsilon}'^T]$ , where  $E[\cdot]$  is the expected-value operator. Assuming Gaussian error distributions for  $\boldsymbol{\varepsilon}_a$  and  $\boldsymbol{\varepsilon}$  we obtain (Rodgers, 2000):

$$\hat{\mathbf{x}} = \mathbf{x}_a + \mathbf{G}(\mathbf{x}' - \mathbf{K}\mathbf{x}_a) \quad (2)$$

where  $\mathbf{K}$  is the observation operator that maps the model forecast to the observations. For satellite measurements  $\mathbf{K}\mathbf{x}_a = \mathbf{x}_s + \mathbf{A}(\mathbf{x}_a - \mathbf{x}_s)$  (Eq. (1) with no noise term), while for surface measurements  $\mathbf{K}\mathbf{x}_a = \mathbf{x}_a$ . The gain matrix  $\mathbf{G}$  is given by

$$\mathbf{G} = \mathbf{S}_a \mathbf{K}^T (\mathbf{K} \mathbf{S}_a \mathbf{K}^T + \mathbf{S}_\varepsilon)^{-1} \quad (3)$$

and determines the relative weight given to the observations and the model. The instrument error covariance matrix  $\mathbf{S}_\varepsilon$  is assumed diagonal and set to an arbitrarily large number in locations where there are no observations. For surface measurements we include the 5 % representation error in quadrature with the 2 % instrument error so that

the corresponding error variances are additive. The optimal estimate  $\hat{\mathbf{x}}$  has error  $\hat{\boldsymbol{\varepsilon}}$  with error covariance  $\hat{\mathbf{S}} = E[\hat{\boldsymbol{\varepsilon}}\hat{\boldsymbol{\varepsilon}}^T]$ :

$$\hat{\mathbf{S}} = (\mathbf{I}_n - \mathbf{GK})\mathbf{S}_a \quad (4)$$

5 Where  $\mathbf{I}_n$  is the identity matrix of dimension  $n$ .

The model error covariance matrix  $\mathbf{S}_a$  expresses the error in the forward model at each assimilation time step and is given by:

$$\mathbf{S}_a = \begin{pmatrix} \text{var}(\boldsymbol{\varepsilon}_{a,1}) & \cdots & \text{cov}(\boldsymbol{\varepsilon}_{a,1}, \boldsymbol{\varepsilon}_{a,n}) \\ \vdots & \ddots & \vdots \\ \text{cov}(\boldsymbol{\varepsilon}_{a,n}, \boldsymbol{\varepsilon}_{a,1}) & \cdots & \text{var}(\boldsymbol{\varepsilon}_{a,n}) \end{pmatrix} \quad (5)$$

10 where  $\boldsymbol{\varepsilon}_a = (\boldsymbol{\varepsilon}_{a,1}, \dots, \boldsymbol{\varepsilon}_{a,n})^T$ , with  $\boldsymbol{\varepsilon}_{a,i}$  representing the error for GEOS-Chem gridbox  $i$ . At each assimilation time step the model error is decreased as described by the a posteriori error covariance matrix  $\hat{\mathbf{S}}$  (Eq. 4). The diagonal terms of  $\hat{\mathbf{S}}$  are initialized using an a priori model error of 29% and transported as tracers in GEOS-Chem to the next assimilation time step and are augmented by a model error variance reflecting the time-dependent divergence of the model from the true state following Zoogman et al. (2011). This yields the diagonal terms  $\text{var}(\boldsymbol{\varepsilon}_{a,i})$  of  $\mathbf{S}_a$ . The off-diagonal terms (error covariances) describe the propagation of information from each observation over a spatial domain of influence. We compute  $\text{cov}(\boldsymbol{\varepsilon}_{a,i}, \boldsymbol{\varepsilon}_{a,j})$  for each pair of gridboxes  $(i, j)$  as a function of the horizontal and vertical distance between the two gridboxes using the error correlation length scales from Sect. 2.4.

20 In practice the dimension of the matrices used in the assimilation must be limited to make the computation tractable. This is done by solving Eq. (2) column by column and including only measurements at a horizontal distance less than 510 km (the horizontal error correlation length scale, see below) in the model error covariance matrix.

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## 2.4 Error correlation length scales

The spatial extent of information provided by an observation to correct the GEOS-Chem model simulation through data assimilation can be quantified by correlating the GEOS-Chem errors relative to in situ observations at different sites in the Intermountain West (for the horizontal scale) and ozonesonde profiles (for the vertical scale). To define a horizontal error correlation length scale we used actual CAST-Net surface measurements from our period of study (April–June 2010), downloaded from [www.epa.gov/castnet/](http://www.epa.gov/castnet/). We compute the time series of model error during day-time (09:00–17:00 LT) at each surface site, and from there derive the model error correlation between each pair of surface sites. Figure 3 (left) shows the correlation coefficients plotted against the distance  $d$  between sites (binned every 100 km). We find  $R = \exp(-d/510 \text{ km})$ . We also show the error correlation length scale calculated when comparing GEOS-Chem and GFDL AM3 (in red) sampled over the Intermountain West region. The model-model error correlation length scale is similar to the model-observation length scale and this provides some support for the realism of error patterns in our OSSE. We assume that the horizontal error correlation length scale is invariant with altitude.

To estimate the vertical correlation length scale we compare GEOS-Chem ozone concentrations to in situ vertical profiles from May–June 2010 ozonesondes at six locations in California (Cooper et al., 2011). Figure 3 (right) shows the correlation coefficients plotted against the vertical distance  $z$  (binned every 500 m) for the time series of model errors at each ozonesonde station from the surface to 8 km altitude. We find  $R = \exp(-z/1.7 \text{ km})$ . Again, the model-model length scale (red) is not significantly different from the model-observation length scale.

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### 3 TEMPO observation of high-ozone events in the Intermountain West

We now apply our OSSE system to evaluate the benefit of TEMPO observations to monitor and attribute ozone exceedances in the Intermountain West. We compare the “true” concentrations in surface air over the Intermountain West to GEOS-Chem CTM ozone concentrations without data assimilation (a priori) and with assimilation of synthetic CASTNet, TEMPO, and IASI-3 LEO observations. We also performed an assimilation of CASTNet and TEMPO observations without a LEO instrument and found no significant difference in results. Thus the LEO instrument does not add significant information beyond TEMPO for constraining surface ozone concentrations in the Intermountain West. Its value for tracking exceptional events will be discussed in Sect. 4.

Figure 4 examines the ability of the data assimilation system to monitor daily MDA8 ozone over the Intermountain West at the  $1/2^\circ \times 2/3^\circ$  ( $\sim 50 \text{ km} \times 50 \text{ km}$ ) GEOS-Chem grid resolution. The top panel shows a scatterplot of a priori GEOS-Chem MDA8 ozone concentrations in April–June 2010, for individual grid squares over the Intermountain West domain of Fig. 1 and individual days, vs. the “true” concentrations from the GFDL AM3 model. The GEOS-Chem a priori is biased low and performs poorly in reproducing the “true” variability ( $R^2 = 0.12$ , bias =  $-9.0$  ppbv). Assimilation of synthetic CASTNet surface measurements reduces the low bias from 9.0 to 2.8 ppbv, but still does not capture much of the variability ( $R^2 = 0.34$ ). Adding the synthetic TEMPO geostationary observations eliminates the low bias and captures over half of the variability ( $R^2 = 0.58$ ).

The ability of TEMPO observations to capture high-ozone events is of particular interest. Figure 5 shows a map of the number of days in April–June 2010 with MDA8 ozone in excess of 70 ppbv for individual GEOS-Chem gridsquares in the Intermountain West. Values are shown for the “true” atmosphere, the GEOS-Chem a priori without data assimilation, and the data assimilation results including only the CASTNet observations and with the addition of TEMPO observations. The “truth” shows an average of 5.7 high-ozone events per gridsquare in the Intermountain West over the April–June 2010

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period. The a priori model has only 0.8 event-days per gridsquare and the spatial pattern is very different (spatial correlation  $R^2 = 0.09$  for the ensemble of Intermountain West gridsquares). Assimilation of surface measurements improves both the average number of high-ozone events (3.6 event-days) and the spatial pattern ( $R^2 = 0.62$ ). The inability to fully correct the bias is due in part to the large impact of free tropospheric air in driving high-ozone events, and in part to the limited coverage from the sparse surface network. Adding TEMPO satellite observations almost fully corrects the bias (mean of 5.4 event-days) and captures most of the spatial distribution of high-ozone events ( $R^2 = 0.82$ ).

#### 4 Attribution of exceptional events using TEMPO observations

TEMPO will provide continuous observation in the free troposphere as well as in the boundary layer, with separation between the two (Fig. 2). Thus it could be particularly powerful in quantifying free tropospheric background contributions to NAAQS exceedances. This would assist in the designation of exceptional events where an exceedance of the NAAQS is considered to be outside local control.

We examine a case study of a stratospheric intrusion on 13 June in the GFDL AM3 model taken as the “truth”. Figure 6 shows a time series for June 2010 of MDA8 ozone concentrations at a location in northern New Mexico ( $107^\circ$  W,  $36^\circ$  N). We choose this event as it was diagnosed by ozonesonde observations and meteorological tracers as a deep stratospheric intrusion event (Lin et al., 2012b). Actual observations at nearby CASTNet locations indicate ozone in excess of 75 ppbv during the 12–15 June period.

Evidence of free tropospheric origin for the 13 June event is critical to achieving an “exceptional event” designation. Figure 7 (top left) shows a longitude-altitude cross section of ozone concentrations in the GFDL AM3 model taken as the “truth”. The stratospheric intrusion is manifest at  $103$ – $109^\circ$  W. The a priori GEOS-Chem model (top right) also shows a stratospheric ozone enhancement extending to the surface but of much smaller magnitude. Assimilation of surface measurements (not shown) makes

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These are in close agreement with error correlation length scales between the two models used in our OSSE.

We find that the CASTNet surface observations are too sparse to adequately monitor high-ozone events in the Intermountain West even after data assimilation. We show that the TEMPO geostationary observations will provide a greatly improved observing system for monitoring such events. In addition, because of the information they provide on the vertical distribution of ozone, they can effectively diagnose NAAQS exceedances caused by background ozone. A LEO satellite instrument flying concurrently with TEMPO provide no significant added value for monitoring the ozone background over the US but could be useful for tracking transpacific plumes.

Use of the complete observing system described here (surface, geostationary, and LEO) will provide a powerful tool for future air quality policy. Planning is underway to combine this system with regional air quality models to supply the public with near real time pollution reports and forecasts. These reports and forecasts would be much the same as currently available weather information, also provided in large part from geostationary satellite observations.

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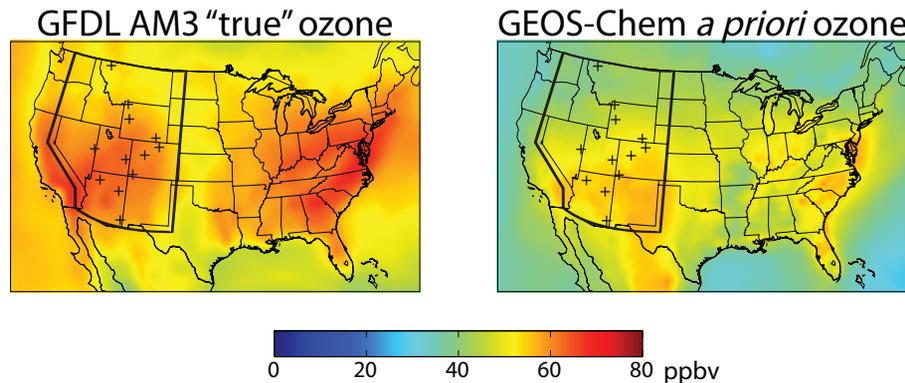
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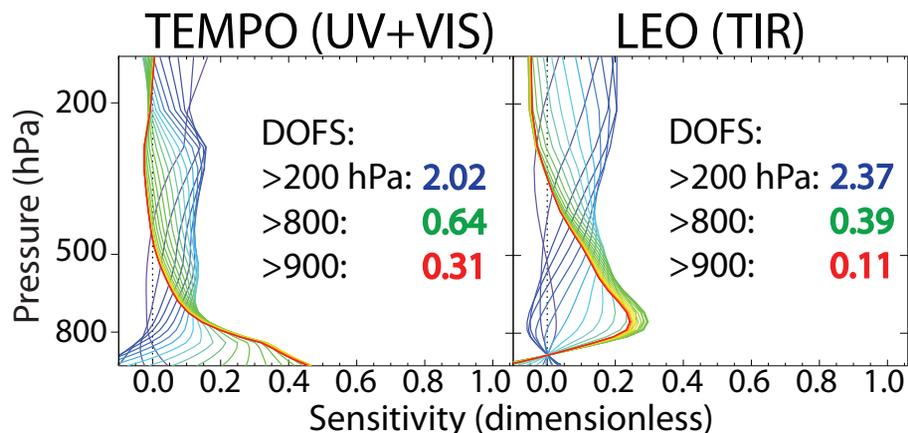


**Fig. 1.** Mean values of the daily maximum 8 h average (MDA8) ozone concentrations for April–June 2010 in surface air. Left panel shows values from the GFDL AM3 CCM used as the “true” atmosphere in our OSSE. Right panel shows the a priori values from the GEOS-Chem CTM used for data assimilation. The black lines delineate the Intermountain West and black crosses show CASTNet surface measurement sites in the region.

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**Fig. 2.** Averaging kernel matrices assumed in this study (from Natraj et al., 2011) for clear-sky retrievals of tropospheric ozone from space in the UV+Vis (left) and the TIR (right). UV+Vis in our study corresponds to TEMPO, while TIR corresponds to a future LEO instrument flying concurrently with TEMPO. Lines are matrix rows for individual vertical levels, with the color gradient from red to blue corresponding to vertical levels ranging from surface air (red) to 200 hPa (blue). Inset are the degrees of freedom for signal (DOFS) for the atmospheric columns below 200, 800, and 900 hPa.

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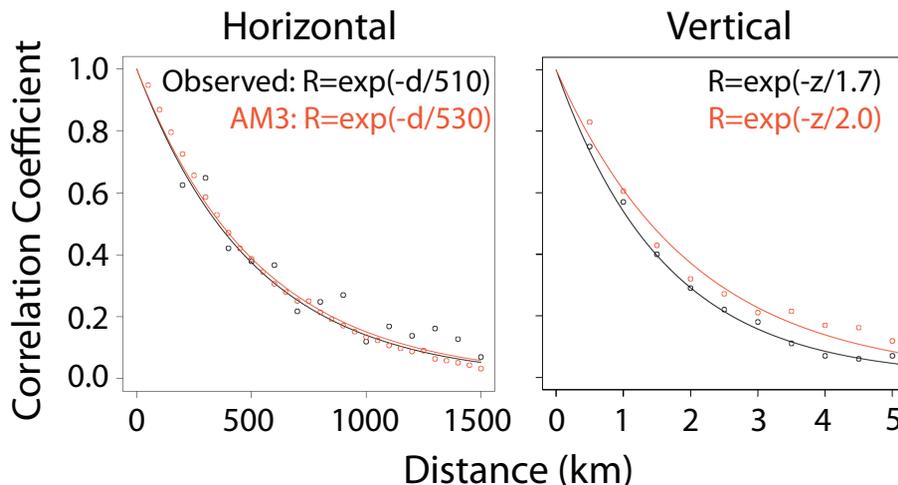
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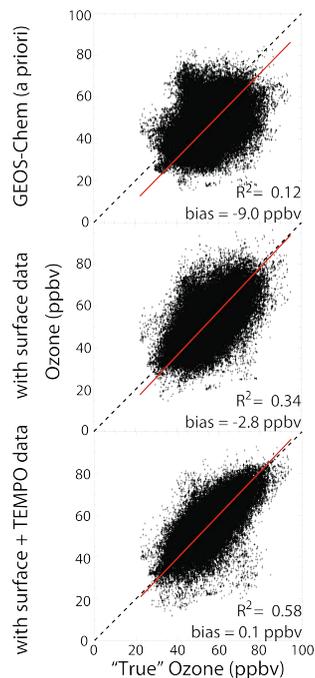


**Fig. 3.** Error correlation length scales for the GEOS-Chem model simulation of tropospheric ozone in the US Intermountain West. The error correlations are relative to actual CASTNet and ozonesonde observations (in black) and relative to the GFDL AM3 model sampled in the Intermountain West region (in red). Statistics are computed for April–June 2010. The left panel shows the correlation coefficient ( $R$ ) of the model error between pairs of CASTNet sites, plotted against the distance between sites. Values are for the 12 CASTNet sites in the Intermountain West (Fig. 1). The right panel shows the correlation coefficient of the model error between pairs of vertical levels (up to 8 km altitude) for ozonesonde measurements from the IONS-2010 campaign in California (Cooper et al., 2011), plotted against distance between levels. Exponential fits to the data are shown inset, where  $d$  and  $z$  are horizontal and vertical distances in km.

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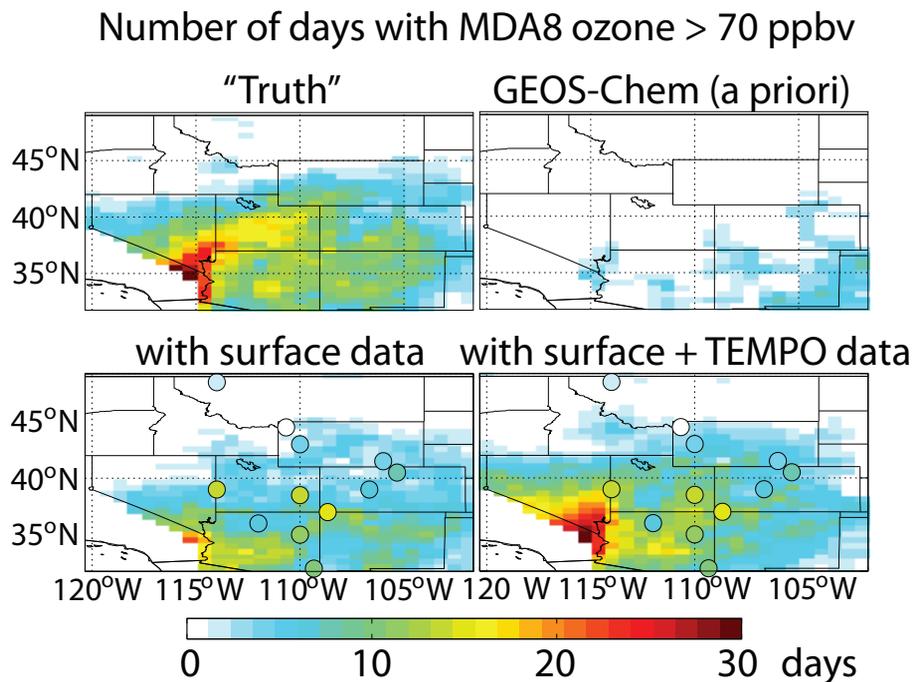


**Fig. 4.** Improved monitoring of surface ozone across the Intermountain West from data assimilation of synthetic CASTNet (surface) and TEMPO (geostationary satellite) observations. The figure shows scatterplots of simulated (GEOS-Chem) vs. “truth” (GFDL AM3) daily maximum 8 h (MDA8) surface ozone for April–June 2010 for all  $1/2^\circ \times 2/3^\circ$  grid squares in the region (Fig. 1) and for individual days. Results are for GEOS-Chem without data assimilation (top), with assimilation of CASTnet synthetic surface data (middle), and with additional assimilation of TEMPO, and LEO synthetic satellite data (bottom). Comparison statistics are inset. Also shown are the reduced-major-axis (RMA) regression line and the 1 : 1 line.

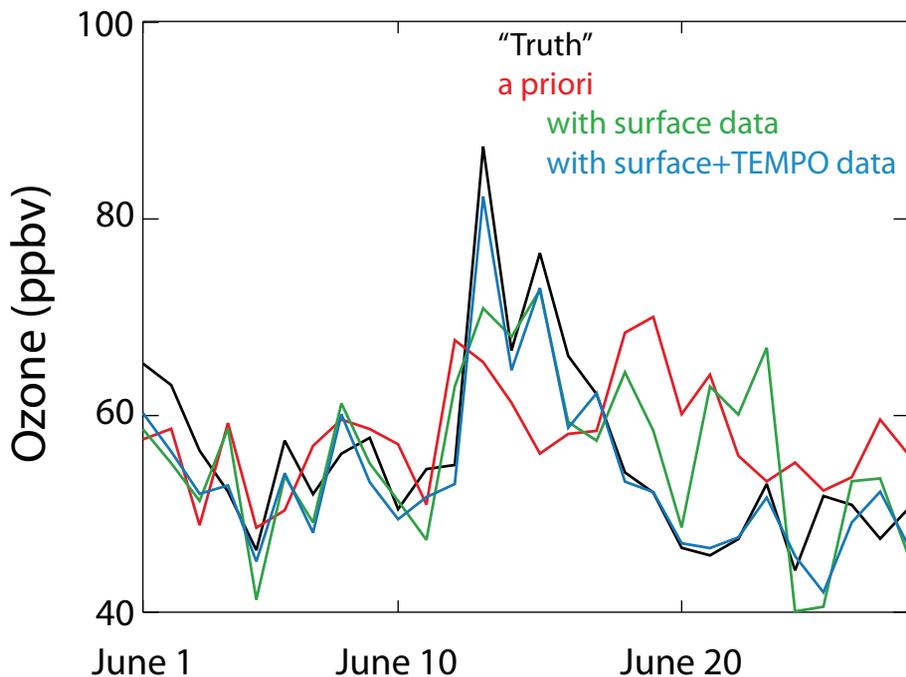
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**Fig. 5.** Improved detection of high-ozone events in the Intermountain West by data assimilation. The figure shows the number of events (daily maximum 8 h ozone > 70 ppbv) in April–June 2010 on the GEOS-Chem grid. The “truth” defined by the GFDL AM3 model (top left panel) is compared to GEOS-Chem simulations without data assimilation (top right), with assimilation of synthetic CASTNet surface data (bottom left), and with additional assimilation of synthetic TEMPO and LEO satellite data (bottom right). Locations of CASTNet surface sites used for assimilation with their “true” values are overlain in the bottom panels.



**Fig. 6.** Detection of an exceptional ozone event by TEMPO. The figure shows the June 2010 time series of daily maximum 8 h (MDA8) ozone concentrations at a location in northern New Mexico ( $107^{\circ}$  W,  $36^{\circ}$  N) featuring a major stratospheric intrusion on 13 June in the GFDL AM3 model taken as the “truth” (black line). The ability to capture this event is examined for the GEOS-Chem model without data assimilation (a priori, red line) and with assimilation of surface measurements only (green line) and satellite measurements added (blue line).

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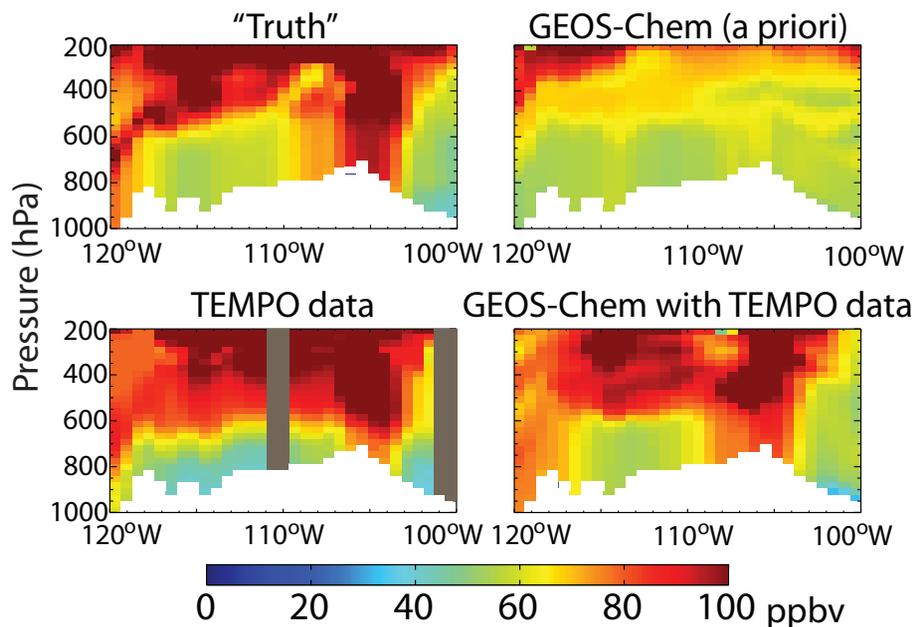
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**Fig. 7.** Longitude–altitude cross-section of ozone concentrations ( $36^{\circ}$  N, 3 GMT on 14 June 2010) associated with the stratospheric intrusion of Fig. 6. The “true” state from the GFDL AM3 model (top left) is compared to the GEOS-Chem model without data assimilation (top right) and with assimilation of surface and satellite data (bottom right). The bottom left panel shows synthetic TEMPO observations of the “true” state (gray regions indicate cloudy scenes) without data assimilation. Local topography is shown in white.