

# Ozone Data Assimilation with GEOS-Chem: a Comparison Between 3D-Var, 4D-Var, and Suboptimal Kalman Filter Approaches

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**Abstract.** Chemistry transport models determine the evolving chemical state of the atmosphere by solving the fundamental equations that govern physical and chemical transformations. Models are imperfect and cannot capture completely the complex dynamics of the atmosphere. As a result there is always a mismatch between model generated estimates and reality. Data assimilation is the procedure to combine  
5 data from observations with model predictions to obtain a more accurate representation of the state of the atmosphere.

Two families of data assimilation methods are currently widely used: variational and Kalman filter (KF). The variational approach is based on control theory, and formulates data assimilation as a minimization problem of a cost functional that measures the model-observations mismatch. The Kalman filter approach  
10 is rooted in statistical estimation theory and provides the analysis covariance together with the best state estimate. Suboptimal Kalman filters employ different approximations of the covariances in order to make the computations feasible with large models. Each family of methods has both merits and drawbacks.

This paper compares several data assimilation methods used for global chemical data assimilation. Specifically, we evaluate data assimilation approaches for improving estimates of the summertime global tropo-  
15 spheric ozone distribution; the data is provided by retrievals from Tropospheric Emission Spectrometer (TES) and the model is GEOS-Chem. The resulting analyses are compared against an independent data set provided by ozonesonde measurements to assess the effectiveness of each assimilation method. All assimilation methods provide notable improvements over the free model run results, which differ from the ozonesonde measurements by about 20% (below 200 hPa). Four dimensional variational data assim-

20 ilation with window lengths between five days and two weeks is the most accurate method, with mean differences between analysis profiles and ozonesonde measurements of 1-5%. Two sequential assimilation approaches (three dimensional variational and suboptimal KF), although derived from different theoretical considerations, provide similar ozone estimates, with relative differences of 5-10% between the analyses and ozonesonde measurements.

## 25 **1 Introduction**

Understanding the distribution of tropospheric ozone is of considerable interest (Li et al., 2005; Cooper et al., 2006, 2007; Hudman et al., 2007). Ozone is an integral constituent of the troposphere that plays a significant role in determining the chemical and radiative state of the lower atmosphere. Ozone in the stratosphere absorbs high energy radiation, thus preventing the disintegration of DNA molecules and  
30 supporting the existence of life on the planet. In the upper troposphere ozone is a greenhouse gas through absorption of upwelling long wave radiation. In the mid-troposphere ozone is a precursor to OH radicals which moderate pollution levels. At the surface ozone is a pollutant causing respiratory problems and affecting crop yields.

Numerous attempts to quantify the amount of tropospheric ozone and characterizing its distribution  
35 through chemical transport models and general circulation models have been made (Horowitz et al., 2003; Horowitz, 2006). The findings from these studies vary significantly due to the strong variability in ozone lifetimes and uncertainties in determining the amount of ozone lost through dry deposition, entered through upper troposphere-stratosphere exchanges, or evolved due to chemical reactions of trace gas and emission precursors. The ozone lifetime varies from a few minutes at the surface, to a few days  
40 in the lower troposphere, to months in the upper troposphere. In such situations, it is important to validate the accuracy of model predictions against observed state of the atmosphere. Studies of variations in tropospheric ozone have been conducted through ozonesonde measurements and surface observations (Logan, 1994, 1999; Tarasick et al., 2005; Oltmans et al., 2006), while through satellite measurements (Munro et al., 1998; Tellmann et al., 2004).

45 Chemical data assimilation is a process of optimally combining imperfect observations of reality with imperfect model results to produce a better estimate of the chemical state of the atmosphere. Considerable experience with data assimilation has been accumulated in the field of numerical weather prediction (Daley, 1991; Courtier et al., 1998; Rabier et al., 2000; Kalnay, 2002; Navon, 2009). In this work we focus on chemical data assimilation, i.e., on assimilation of observations of pollutant levels in the atmosphere.  
50 Chemical data assimilation poses specific challenges related to the multiphysics nature of the system, the stiffness of chemical kinetic equations, the sparseness of chemical observations, and the uncertainty in the levels of anthropogenic and natural pollutants emitted into the atmosphere. Throughout this paper we

will refer to model results as model predictions or model forecasts even when a past period is simulated.

Previous studies have employed various approaches to assimilating observations of trace gases for improved tropospheric chemistry representations. Data assimilation has been used to improve initial conditions, emissions, and boundary values, and has resulted in improved air quality forecasts. The base concepts of the variational approach to chemical data assimilation, and the construction of adjoint chemical transport models are discussed in detail in (Sandu et al., 2005; Carmichael et al., 2008). Early work in chemical data assimilation using variational techniques has been reported in (Khattatov et al., 2000; Elbern and Schmidt, 2001). Since then there is a growing body of literature with applications of 3D-Var and 4D-Var chemical data assimilations. 3D-Var was first used by (Derber et al., 1991; Parrish and Derber, 1992) and later applied by most of the meteorological centers (Courtier et al., 1998; Cohn et al., 1998; Gauthier et al., 1999a). A study on ozone improvement using 3D-Var assimilation is presented in (Bei et al., 2008). Adjustment of gas phase chemical tracer initial conditions has been studied in (Chai et al., 2007; Zhang et al., 2008). Adjustment of pollutant emissions through 4D-Var chemical data assimilation has been discussed in (Chai et al., 2009). Data assimilation studies involving particle measurements to improve aerosol fields have been performed in (Hakami et al., 2005; Henze et al., 2009). Suboptimal Kalman filters have been employed successfully for chemical data assimilation (Khattatov et al., 2000; Menard et al., 2000; Lamarque et al., 2002; Segers et al., 2005; Clark et al., 2006; Pierce et al., 2007; Parrington et al., 2009). The use of the ensemble Kalman filter (EnKF) (Evensen, 1994) in chemical data assimilation has been studied in (Constantinescu et al., 2007b,c).

Different approaches to data assimilation are rooted in different theories (control, statistical estimation), have different implementation and computational costs, and yield different performance on large scale problems of practical interest. A discussion on relationship between optimality of variational data assimilation and Kalman filters is presented in (Li and Navon, 2001). (Houtekamer, 2005) compared the quality of background statistics in 3D-Var and EnKF using radiance observations from satellite, while, (Laroche et al., 2005) compared the characteristics 3D-Var and 4D-Var introduced in the operational suite of the Canadian Meteorological Center (CMC). (Constantinescu et al., 2007c) and (Wu et al., 2008) compare the performances of EnKF with 4D-Var for chemical transport models on a regional scale using ground-level ozone measurements, while, (Geer et al., 2006) provides an intercomparison of tropospheric ozone estimates obtained through 3D-Var, 4D-Var and Kalman filter assimilation systems for both chemical transport and global circulation models as part of the Assimilation of ENVISAT Data (ASSET) project.

The Tropospheric Emission Spectrometer (TES) (Beer et al., 2001) is the first dedicated infrared instrument from which information of the global and vertical distribution of tropospheric ozone can be retrieved. (Parrington et al., 2009) provided the first set of results from the assimilation of vertical profiles of ozone from TES into the GEOS-Chem using suboptimal Kalman filter. We have developed 3D-Var and 4D-Var data assimilation capabilities for GEOS-Chem v7. The goal of this paper is to provide the first direct

comparison of global tropospheric ozone distribution estimated through 3D-Var, 4D-Var and suboptimal KF assimilation systems showcasing the potential of TES profile retrievals. The assessment of analyses generated through different assimilation systems are on the similar lines of (Geer et al., 2006; Parrington et al., 2009).

This work differs from (Wu et al., 2008) in several aspects. First, we study the global ozone distribution assimilating satellite observations into GEOS-Chem as compared to the ozone forecasts over western Europe through assimilation of observations from ground stations into Polair3D (Boutahar et al., 2004). We evaluate the quality of tropospheric ozone analyses estimated by our assimilation systems through ozonesonde measurement data, an independent observation dataset not used in assimilation, while (Wu et al., 2008) has a forecast scoring scheme, where the scores are calculated as root mean square of the differences between analyses and assimilated observations. Not surprisingly, their optimal interpolation (OI) scheme fetched the best overall score while 4D-Var performed average. Such a scoring scheme does not provide a fair assessment of the performance of an assimilation system since there is no provision to adjudge whether observations are erroneous. In addition, the average performance of their 4D-Var assimilation could be attributed to the limitations of their study as they adjusted only the initial conditions while boundary conditions remained unchanged, making the assimilation ineffective especially for long range simulations. We believe this plays a significant role in the decrease of their 4D-Var performance with increase in assimilation window lengths, although they have attributed it solely to the model errors. Such a situation does not arise in our case as global assimilations are not restricted by any horizontal boundaries.

This paper is structured as follows. Section 2 provides the mathematical overview of how observations are integrated into the model in different data assimilation systems. Section 5.2 discusses the characteristics of background error covariance matrices used in this study. Section 3 provides a brief overview of the global chemical transport model (GEOS-Chem) and its adjoint development. A description of the TES instrument, its observation operator and profile retrieval formulation is provided in Section 4. Section 6 details the experimental settings, computational costs and assessment of tropospheric ozone estimates through different assimilation systems. Summary and points of future work are discussed in Section 7.

## 115 2 Chemical data assimilation

Variational methods solve the data assimilation problem in an optimal control framework (Sasaki, 1958; Le Dimet and Talagrand, 1986; Courtier and Talagrand, 1987; Lions, 1971). Specifically, they attempt to find the control variable values (e.g., initial conditions) by minimizing the discrepancy between the model forecast and observations; the minimization is constrained by the governing dynamic equations. In contrast, statistical estimation methods (generically known as Kalman filters/smoothers) solve the data

assimilation problem in a Bayesian framework by combining probability densities of errors from different sources (Khattatov et al., 2000; Menard et al., 2000; Lamarque et al., 2002; Segers et al., 2005; Clark et al., 2006; Pierce et al., 2007; Parrington et al., 2009; Constantinescu et al., 2007b,c). In the following discussion, for simplicity of presentation, we focus on discrete models (in time and space) where the initial conditions are the control variables.

Data assimilation provides best estimates of the state of the atmosphere by combining the following three sources of information.

1. The apriori, or background state  $\mathbf{x}^b$  represents the best estimate of the true state  $\mathbf{x}^t$  available before any measurements are taken. This estimate is assumed unbiased, and the random background (estimation) errors  $\varepsilon^b$  are typically assumed to have a normal probability density with a background error covariance matrix  $\mathbb{B}$

$$\varepsilon^b = \mathbf{x}^b - \mathbf{x}^t \in \mathcal{N}(0, \mathbb{B}). \quad (1)$$

2. The model encapsulates our knowledge about physical and chemical laws that govern the evolution of the system. The model evolves an initial state  $\mathbf{x}_0 \in \mathbb{R}^n$  at the initial time  $t_0$  to future state values  $\mathbf{x}_i \in \mathbb{R}^n$  at future times  $t_i$ ,

$$\mathbf{x}_i = \mathcal{M}_{t_0 \rightarrow t_i}(\mathbf{x}_0). \quad (2)$$

The size of the state space in realistic chemical transport models is very large. For example, a GEOS-Chem simulation at the  $2^\circ \times 2.5^\circ$  horizontal resolution has  $n \in \mathcal{O}(10^8)$  variables.

3. Observations  $\mathbf{x}_i^{\text{obs}} \in \mathbb{R}^m$  of the state are taken at times  $t_i$ ,  $1 = 1, \dots, N$

$$\mathbf{x}_i^{\text{obs}} = \mathcal{H}(\mathbf{x}_i) + \varepsilon_i^{\text{obs}}. \quad (3)$$

The observation operator  $\mathcal{H}$  maps the state space onto the observation space. In many practical situations  $\mathcal{H}$  is a highly nonlinear mapping (as is the case, e.g., with satellite observation operators). The observations are characterized by measurement and representativeness errors  $\varepsilon_i^{\text{obs}}$ . The observation errors at each time are assumed to be independent of background errors, and independent of the observation errors at other times. They are typically assumed to have a normal distribution with mean zero and covariance  $\mathbb{R}_i$ ,

$$\varepsilon_i^{\text{obs}} \in \mathcal{N}(0, \mathbb{R}_i). \quad (4)$$

Based on these three sources of information data assimilation computes the posterior estimate  $\mathbf{x}^a$  of the true state;  $\mathbf{x}^a$  is called the “analysis” state.

## 2.1 Three dimensional variational (3D-Var) data assimilation

In the 3D-Var data assimilation the observations (3) are considered successively at times  $t_1, \dots, t_N$ . The background state (i.e., the best state estimate at time  $t_i$ ) is given by the model forecast, starting from the previous analysis (i.e., best estimate at time  $t_{i-1}$ ):

$$\mathbf{x}_i^b = \mathcal{M}_{t_{i-1} \rightarrow t_i}(\mathbf{x}_{i-1}^a).$$

The discrepancy between the model state  $\mathbf{x}_i$  and observations at time  $t_i$ , together with the departure of the state from the model forecast  $\mathbf{x}_i^b$ , are measured by the 3D-Var cost function:

$$\mathbb{J}(\mathbf{x}_i) = \frac{1}{2}(\mathbf{x}_i - \mathbf{x}_i^b)^T \mathbb{B}^{-1}(\mathbf{x}_i - \mathbf{x}_i^b) + \frac{1}{2}(\mathcal{H}(\mathbf{x}_i) - \mathbf{x}_i^{obs})^T \mathbb{R}_i^{-1}(\mathcal{H}(\mathbf{x}_i) - \mathbf{x}_i^{obs}) \quad (5)$$

While in principle a different background covariance matrix should be used at each time, in practice the same matrix is re-used throughout the assimilation window. The 3D-Var analysis is computed as the state which minimizes (5)

$$\mathbf{x}_i^a = \operatorname{argmin} \mathbb{J}(\mathbf{x}_i). \quad (6)$$

Typically a gradient-based numerical optimization procedure is employed to solve (6). The gradient  $\nabla \mathbb{J}$  of the cost function (5) is

$$\nabla \mathbb{J}(\mathbf{x}_i) = \mathbb{B}^{-1}(\mathbf{x}_i - \mathbf{x}_i^b) + (\mathcal{H}'(\mathbf{x}_i))^T \mathbb{R}_i^{-1}(\mathcal{H}(\mathbf{x}_i) - \mathbf{x}_i^{obs}) \quad (7)$$

135 Note that the gradient requires to computation of the linearized observation operator  $\mathcal{H}'$  about the current state.

Preconditioning is often used to improve convergence of the numerical optimization problem (6). A change of variables is performed, for example, by shifting the state and scaling it with the square root of covariance:

$$\hat{\mathbf{x}}_i = \mathbb{B}^{1/2}(\mathbf{x}_i - \mathbf{x}_i^b), \quad (8)$$

The optimization is then carried out on the new variables  $\hat{\mathbf{x}}_i$ .

## 2.2 Four dimensional variational (4D-Var) data assimilation

140 In strongly-constrained 4D-Var data assimilation all observations (3) at all times  $t_1, \dots, t_N$  are simultaneously considered. The control parameters are the initial conditions  $\mathbf{x}_0$ ; they uniquely determine the state of the system at all future times via the model equation (2).

The discrepancy between model predictions and observations at all future times  $t_1, \dots, t_N$ , together with the departure of the initial state from the background state, are measured by the 4D-var cost function:

$$\mathbb{J}(\mathbf{x}_0) = \frac{1}{2}(\mathbf{x}_0 - \mathbf{x}_0^b)^T \mathbb{B}^{-1}(\mathbf{x}_0 - \mathbf{x}_0^b) + \frac{1}{2} \sum_{i=1}^N (\mathcal{H}(\mathbf{x}_i) - \mathbf{x}_i^{obs})^T \mathbb{R}_i^{-1}(\mathcal{H}(\mathbf{x}_i) - \mathbf{x}_i^{obs}) \quad (9)$$

Note that the departure of the initial conditions from the background is weighted by the inverse background covariance matrix,  $\mathbb{B}^{-1}$ , while the differences between the model predictions  $\mathcal{H}(\mathbf{x}_i)$  and observations  $\mathbf{x}_i^{obs}$  are weighted by the inverse observation error covariances,  $\mathbb{R}_i^{-1}$ .

The 4D-Var analysis is computed as the initial condition which minimizes (9) subject to the model equation constraints (2)

$$\mathbf{x}_0^a = \arg \min \mathbb{J}(\mathbf{x}_0) \quad \text{subject to (2)}. \quad (10)$$

145 The model (2) propagates the optimal initial condition (9) forward in time to provide the analysis at future times,  $\mathbf{x}_i^a = \mathcal{M}_{t_0 \rightarrow t_i}(\mathbf{x}_0^a)$ .

The optimization problem (10) is solved numerically using a gradient-based technique. The gradient of (9) reads

$$\nabla \mathbb{J}(\mathbf{x}_0) = \mathbb{B}^{-1}(\mathbf{x}_0 - \mathbf{x}_0^b) + \sum_{i=1}^N \left( \frac{\partial \mathbf{x}_i}{\partial \mathbf{x}_0} \right)^T (\mathcal{H}'(\mathbf{x}_i))^T \mathbb{R}_i^{-1} (\mathcal{H}(\mathbf{x}_i) - \mathbf{x}_i^{obs}) \quad (11)$$

The 4D-Var gradient requires not only the linearized observation operator  $\mathcal{H}'$ , but also the transposed derivative of future states with respect to the initial conditions. The 4D-Var gradient can be obtained effectively by forcing the adjoint model with observation increments, and running it backwards in time.

150 The construction of an adjoint model requires considerable effort, time, and know-how.

### 2.3 Suboptimal Kalman filter

The suboptimal Kalman filter is a sequential data assimilation approach (Khattatov et al., 2000) in which corrections in the concentration state vector are performed as soon as observations become available. Similar to 3D-Var, for every observation time  $t_i$ , this technique starts with the model forecast state ( $\mathbf{x}_i^f$ ) and provides an expected analysis state ( $\mathbf{x}_i^a$ ) that reduces the discrepancy between the model forecast and the observations  $\mathbf{x}_i^{obs}$ . The analysis state vector is obtained as

$$\mathbf{x}_i^a = \mathbf{x}_i^f + K_i (\mathbf{x}_i^{obs} - \mathcal{H}(\mathbf{x}_i^f)) \quad (12)$$

where  $K$  is the Kalman gain matrix,  $\mathcal{H}$  is the observation operator defined in equation (3), and  $\mathbf{x}^{obs}$  the vector of observations at a given time. The Kalman gain matrix is defined as

$$K_i = \mathbb{P}_i^f \mathcal{H}^T (H_i \mathbb{P}_i^f H_i^T + \mathbb{R}_i)^{-1} \quad (13)$$

where  $\mathbb{P}_i^f$  is the forecast error covariance matrix,  $\mathbb{R}_i$  is the observation error covariance matrix (4), and  $H_i = \mathcal{H}'(\mathbf{x}_i^f)$  is the linearized observation operator about the forecast state. If a diagonal or block-diagonal approximation of the error covariance matrix  $\mathbb{P}^f$  is used in equation (13), the analysis state generated  
155 through equation (12) is suboptimal. A description on the structure of  $\mathbb{P}^f$  is provided in Section 5.2.

At each observation time, along with the analysis state, the analysis error covariance matrix  $\mathbb{P}_i^a$  is also calculated as

$$\mathbb{P}_i^a = (\mathbb{I} - K_i H_i) \mathbb{P}_i^f \quad (14)$$

where  $\mathbb{I}$  is the identity matrix. There are multiple ways in which this analysis covariance matrix is made available to the next observation window; here we will transport variances as passive tracers following (Menard et al., 2000).

### 3 GEOS-Chem

160 In this paper we specifically consider GEOS-Chem (<http://geos-chem.org>), a global three-dimensional chemical transport model (CTM) driven by assimilated meteorological fields from Goddard Earth Observing System(GEOS-4) at the NASA Global Modeling and Assimilation Office (GMAO). It is being widely used by research groups world-wide for performing global atmospheric chemistry studies. The model along with comparison of model predictions with observations was first described in (Bey et al., 2001).  
165 GEOS-Chem accounts in detail for emissions from both natural and anthropogenic sources, for gas phase chemistry, aerosol processes, long range transport of pollutants, troposphere-stratosphere exchanges, etc. Anthropogenic emissions are obtained from the Global Emissions Inventory Activity (GEIA) (Benkovitz et al., 1996) while lightning NO<sub>x</sub> source emissions are estimated using (Price and Rind, 1992), based on deep convective cloud top heights provided with the GMAO meteorological fields. Biomass burning emissions  
170 are based on (Duncan et al., 2003) while biofuel emissions are from (Yevich and Logan, 2003). The meteorological fields have a horizontal resolution of 1° along latitude and 1.25° along longitude with 55 vertical levels, and a temporal resolution of 6 hrs (3 hrs for surface fields).

The GEOS-Chem Adjoint system ([http://wiki.seas.harvard.edu/geos-chem/index.php/GEOS-Chem\\_Adjoint](http://wiki.seas.harvard.edu/geos-chem/index.php/GEOS-Chem_Adjoint)) has been developed through a joint effort of groups at Virginia Tech, University of Colorado, Caltech, Jet  
175 Propulsion Laboratory, and Harvard (Henze et al., 2007; Singh et al., 2009a,b; Eller et al., 2009). The system can perform adjoint sensitivity analyses and 4D-Var chemical data assimilation. Inverse modeling studies with GEOS-Chem-Adjoint are exemplified in (Henze et al., 2009; Kopacz et al., 2007; Zhang et al., 2009).

### 4 Tropospheric Emission Spectrometer (TES) observations

TES (Beer et al., 2001), one of four science instruments aboard NASA's Aura satellite, measures the  
180 infrared-light energy (radiance) emitted by Earth's surface, and by the chemical tracers in the atmosphere (<http://tes.jpl.nasa.gov>). Vertical profiles of chemical concentrations are retrieved from the radiance measurements using an off-line inversion process. In this work we assimilate the retrieved ozone vertical profiles. Figure 1 shows the location of TES profiles for two days.

A-priori information about the vertical concentration profile of the species of interest is needed to solve  
 185 the retrieval inverse problem (the prior information does not come from the measurement). Let  $\mathbf{x}^{\text{prior}}$  be  
 the prior vertical ozone concentration profile (in volume mixing ratio units), and let  $\mathbf{z}^{\text{prior}} = \log \mathbf{x}^{\text{prior}}$ . Let  
 $\mathbf{z}^{\text{radiance}} (= \log \mathbf{x}^{\text{true}})$  be the atmospheric profile as resulting directly from the radiances.

The vertical ozone profile retrieval can be expressed according to the formula

$$\hat{\mathbf{z}} = \mathbf{z}^{\text{prior}} + \mathbb{A} \left( \mathbf{z}^{\text{radiance}} - \mathbf{z}^{\text{prior}} \right) + \mathbb{G} \eta, \quad \hat{\mathbf{x}} = \exp(\hat{\mathbf{z}}). \quad (15)$$

Here  $\mathbb{A}$  is the averaging kernel matrix,  $\mathbb{G}$  is the gain matrix, and  $\eta$  is the spectral measurement error  
 (assumed to have mean zero and covariance  $S_\eta$ ). More details can be found in (Bowman et al., 2002; Jones  
 190 et al., 2003; Worden et al., 2004).

The corresponding TES observation operator  $\mathcal{H}$  is linear with respect to the logarithm of the concentrations,  
 but nonlinear with respect to the concentration profile:

$$\mathcal{H}(\mathbf{x}) = \mathbf{z}^{\text{prior}} + \mathbb{A} \left( \log(L(\mathbf{x})) - \mathbf{z}^{\text{prior}} \right) \quad (16)$$

where  $L$  is an interpolation operator that transforms  $\mathbf{x}$  from the GEOS-Chem  $N$ -level vertical grid to the  
 TES profile retrieval  $P$ -level grid.

For this reason several chemical data assimilation studies based on TES retrieved profiles (Jones et al.,  
 2003; Bowman et al., 2006; Parrington et al., 2009) have opted to perform the suboptimal Kalman filtering  
 step (12) in the logarithm of the concentrations:

$$\log \mathbf{x}^a = \log \mathbf{x}^f + K \left( \hat{\mathbf{z}} - \mathcal{H}(\mathbf{x}^f) \right)$$

Here  $K$  is the Kalman gain matrix,  $\mathcal{H}$  is the observation operator defined in equation (16), and  $\hat{\mathbf{z}}$  is the  
 ozone profile retrievals from TES as described in equation (15). The analysis state is calculated in natural  
 195 logarithm of volume mixing ratio (log VMR) at each observation grid point since the TES profile retrievals  
 are in log VMR. An exponential operator and a linear interpolation operator based on pressure is then  
 applied to this logarithm of analysis state in succession to regain the actual analysis state in GEOS-Chem  
 grid domain. The points which do not lie on the observation grid remain unaffected by the assimilation.

The observation operator  $\mathcal{H}$  that transforms higher resolution model state to the TES profile vertical grid  
 200 (observation grid) domain is expressed by equation (16). The Kalman gain matrix  $K$  is defined by equation  
 (13), particularized to the case where the state is the logarithm of volume mixing ratio.

For variational data assimilation the forcing calculation is carried out in concentrations. For this reason, an  
 adjoint of the observation operator needs to be derived to update the gradients as described in equations

(7) and (11)

$$(\mathcal{H}'(\mathbf{x}))^T \cdot v = \left( \frac{\partial}{\partial \mathbf{x}} (\mathbb{A} \log(L(\mathbf{x}))) \right)^T \cdot v = \left( \frac{\partial L}{\partial \mathbf{x}} \right)^T \cdot \begin{pmatrix} (L\mathbf{x})_0^{-1} & 0 & \dots & 0 \\ 0 & (L\mathbf{x})_1^{-1} & \dots & 0 \\ \vdots & \vdots & \ddots & \vdots \\ 0 & 0 & \dots & (L\mathbf{x})_p^{-1} \end{pmatrix} \cdot \mathbb{A}^T \cdot v$$

Here,  $(\mathcal{H}'(\mathbf{x}))^T$  is a matrix and  $v = \mathbb{R}^{-1}(\mathcal{H}(\mathbf{x}) - \mathbf{x}^{obs})$ . The TES averaging kernel  $A$  is usually a non-symmetric matrix, and the result of  $\mathbb{A}^T \cdot v$  is fed to the interpolation operator to construct the diagonal matrix with the  $i$ -th element being  $1/(L\mathbf{x})_i$ . The term  $(\partial L/\partial \mathbf{x})^T$  is the adjoint of the interpolation operator and brings entities from the TES profile retrieval domain back to the GEOS-Chem model domain.

Note that the TES data can be biased by as much as 10% (Nassar et al., 2008). We have estimated the bias using the technique proposed in (Nassar et al., 2008) and have removed it before assimilating the data.

## 5 Experimental setting for data assimilation

For numerical experiments, we employ GEOS-Chem v7-04-10 adjoint code package (Singh et al., 2009b), capable of performing both 3D-Var and 4D-Var data assimilations with real data. It also incorporates sub-optimal Kalman filter approach of data assimilation developed in Parrington et al. (2009). We assimilate Tropospheric Emission Spectrometer (TES) satellite ozone profile retrievals into the GEOS-Chem model and validate the generated analyses against an independent observation dataset provided by direct ozone profile measurements from ozonesondes. The numerical optimization method used in all variational experiments is the limited memory bound-constrained BFGS (Zhu et al., 1997). This quasi-Newton approach has become the “gold standard” in solving large scale chemical data assimilation problems (Sandu et al., 2005).

Simulations with GEOS-Chem v7 adjoint can be carried out at  $4^\circ \times 5^\circ$  and  $2^\circ \times 2.5^\circ$  resolutions. We have used  $4^\circ \times 5^\circ$  resolution in all our experiments. There are  $46 \times 72$  latitude-longitude grid boxes at this resolution, and 55 vertical levels; near the equator and at ground level each grid box covers an area of about  $400 \text{ km} \times 500 \text{ km}$ . The current GEOS-Chem model does not capture well the dynamics of ozone distribution in the upper troposphere and in the stratosphere. Therefore, we performed data assimilation for only the first 23 model levels (for up to about 50 hPa). The model has been modified to use the linearized ozone (linoz) scheme (McLinden et al., 2000) for a better estimation of ozone exchanges at troposphere-stratosphere boundary. This scheme is going to be used in the next release of the standard GEOS-Chem model.

## 5.1 Data assimilation schemes

The 3D-Var data assimilation experiments were performed for a period of two weeks in the month of August 2006, starting at 00:00(GMT) on August 1st. The TES data was read once every four simulation  
230 hours; the observation operator called at model time  $t$  (hours) reads in all the measurements collected within the interval  $t - 2$  (hours) to  $t + 2$  (hours). 3D-Var data assimilation treats all observations in this interval as instantaneous, and assimilates them in the same optimization run. In all our 3D-Var experiments, we performed 8 iterations per analysis since the cost function decreased significantly within the first few iterations. It is important to note that 3D-Var does not involve any model adjoint calculations; gradients  
235 require only the adjoint of the observation operator. The optimization adjusts ozone concentrations. The generated analysis profile at the end of each observation window is evolved through the forward model that becomes the initial condition for the next observation window. It is also important to mention here that a new background error covariance matrix (17) is constructed for every observation window.

The setup for data assimilation using the suboptimal Kalman filter is quite similar to 3D-Var where we  
240 assimilated TES profile retrievals into GEOS-Chem over a two week period from 00:00 GMT on August 1, 2006 to 00:00 GMT on August 15, 2006. Observations were read every 4 hours and analysis states were generated for each observation window through the sequential update formula (12).

The 4D-Var data assimilation experiments were performed for two different assimilation window lengths to adjudge if model errors hamper the quality of assimilations in GEOS-Chem involving longer assimila-  
245 tion windows; 4D-Var is strongly constrained by the forward model equation (10). Starting at 00:00 GMT on August 1, 2006, the first assimilation window is considered to be of five days while the second window is of two weeks. All the three assimilation systems had the same initial conditions to start with and were generated through a free GEOS-Chem model run. There were 12 optimization iterations performed in order to improve the ozone initial condition. Each iteration during 4D-Var assimilation includes a forward  
250 model and a backward model adjoint run. TES profile retrievals were read every 4 hours during the model adjoint run, and the cost function and adjoint gradients accumulated the impact of all 4 hour data sets throughout the assimilation window. Contrary to 3D-Var and suboptimal KF, where analysis states are generated sequentially every observation window, in 4D-Var the analysis is generated only at the initial time and accounts for the mismatch between observations and model predictions over all the observations  
255 in the assimilation window.

## 5.2 Specification of error variances

We consider a diagonal background error covariance matrix ( $\mathbb{B}$ ) in all our variational data assimilation experiments for simplicity. The initial variances (the diagonal entries of the  $\mathbb{B}$  matrix) are constructed

from the average background concentrations  $\mathbf{x}_0^B$  on each of the  $Nlev$  model vertical layers

$$\mathbb{B} = \begin{bmatrix} \mathbb{B}^{(0)} & 0 \dots & 0 \\ 0 & \mathbb{B}^{(1)} \dots & 0 \\ \vdots & \ddots & \vdots \\ 0 & 0 \dots & \mathbb{B}^{(Nlev)} \end{bmatrix} \quad (17)$$

where

$$\mathbb{B}^{(\ell)} = \begin{bmatrix} \sigma_\ell^2 & 0 \dots & 0 \\ 0 & \sigma_\ell^2 \dots & 0 \\ \vdots & \ddots & \vdots \\ 0 & 0 \dots & \sigma_\ell^2 \end{bmatrix}_{dim \times dim}, \quad dim = Nlon \cdot Nlat, \quad (18)$$

with

$$\sigma_\ell = \frac{\alpha_{rel}}{dim} \sum_{i=1}^{Nlon} \sum_{j=1}^{Nlat} \mathbf{x}_0^B(i, j, \ell, s_{O3}), \quad \ell = 1, \dots, Nlev, \quad s_{O3} = \text{index of ozone} \quad (19)$$

The relative uncertainty level in the background initial conditions is taken to be 50%, i.e.,  $\alpha_{rel} = 0.5$ .

The forecast error covariance matrix  $\mathbb{P}^f$  used in our suboptimal Kalman filter approach is diagonal. The initial forecast error is assumed to be 50% of the initial forecast field that is supposed to capture the representativeness error as well. In matrix form,  $\mathbb{P}_0^f$  is represented as

$$\mathbb{P}_0^f = \begin{bmatrix} \mathbb{P}_0^{f(0)} & 0 \dots & 0 \\ 0 & \mathbb{P}_0^{f(1)} \dots & 0 \\ \vdots & \ddots & \vdots \\ 0 & 0 \dots & \mathbb{P}_0^{f(Nobs)} \end{bmatrix} \quad (20)$$

where  $Nobs$  is the number of observation points (in our case, the number of grid points in the TES retrieval domain). The initial forecast error covariance matrix block corresponding to each observation grid point is given as

$$\mathbb{P}_0^{f(i)} = \alpha_{rel} \cdot \begin{bmatrix} \mathbf{x}_0^f(i, 1, s_{O3}) & 0 & \dots & 0 \\ 0 & \mathbf{x}_0^f(i, 2, s_{O3}) & \dots & 0 \\ \vdots & \ddots & & \vdots \\ 0 & 0 & \dots & \mathbf{x}_0^f(i, Nret, s_{O3}) \end{bmatrix}_{Nret \times Nret}, \quad i = 1, 2, \dots, Nobs \quad (21)$$

where  $Nret$  is the number of vertical TES profile retrieval levels. Although the initial forecast error covariance matrix  $\mathbb{P}^f$  and all analysis  $\mathbb{P}^a$ 's henceforth are diagonal and there are no horizontal correlations being accounted for, the averaging kernels in the observation operator of TES as defined in equation (16) provide vertical correlations when operated on  $\mathbb{P}^f$  through equation (13). A detailed discussion on how to efficiently extend the background error covariance matrices to non-diagonal forms that capture spatial error correlations is provided in Singh et al. (2010a).

## 6 Data assimilation results

### 265 6.1 Computational costs

The 3D-Var and suboptimal KF frameworks are built on top of GEOS-Chem v7 package which uses Sparse Matrix Vectorized GEAR (SMVGEAR) solver for chemistry. However, to construct the adjoint of chemistry required by the 4D-Var, we implemented Kinetic PreProcessor (KPP) solver (Damian et al., 2002) into GEOS-Chem which not only provides a suite of high performance chemical solvers to choose from but  
270 also generates automatically the continuous and discrete adjoint codes (Daescu, 2000, 2003; Sandu et al., 2003). A detailed discussion on interfacing KPP with GEOS-Chem and comparison with native SMVGEAR solver for accuracy and computational performance is presented in Eller et al. (2009). As pointed out in (Henze et al., 2007), the computational cost of Rosenbrock solver increases significantly with the tolerance levels; higher tolerances use smaller internal time steps requiring more computation. In our experiments,  
275 we have set  $RTOL=10^{-3}$  and  $ATOL=10^{-2}$  to achieve moderate to high accuracy.

The suboptimal Kalman filter is less expensive than 3D-Var since it generates the analysis through the single update formula (12), while 3D-Var requires a few iterations before the optimization routine could generate a stable optimal analysis field. This is true however as long as the forecast error covariance matrix is diagonal. Once we move to non-diagonal matrices, the cost of calculating Kalman gain matrix (13) can  
280 be high, although this can be parameterized following, for example, Khattatov et al. (2000). In the case of 3D-Var and 4D-Var, using even full  $\mathbb{B}$  matrix adds a minimal cost to the overall simulation since the complete matrix is never constructed; at each step only a matrix vector product is required and efficient techniques are employed to derive the inverse and other powers of  $\mathbb{B}$  matrix (Singh et al., 2010a). The 4D-Var assimilation is the most expensive of all the assimilation systems under consideration. The reason  
285 is attributed to the fact that a single 4D-Var iteration performs both the forward and adjoint model runs, where, several variables on which the adjoint equation depends on, are written in checkpoint files in the forward model run.

Table 1 provides a comparison of the computational costs of the different data assimilation systems and the cost of free running model for a 24 hour simulation. All the simulations are performed on a Dell  
290 Precision T5400 workstation with 2 quadcore Intel(R) Xeon(R) processors with clock speed 2.33GHz and a RAM of 16GB shared between the two processors.

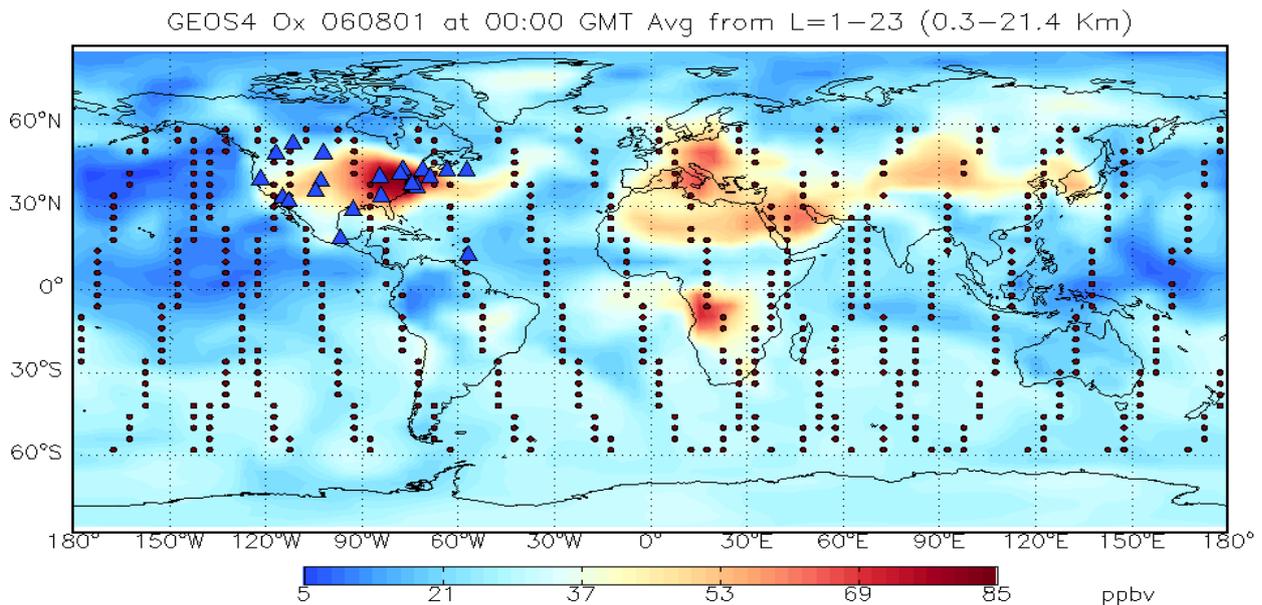
### 6.2 Comparison with ozonesonde measurements

In order to assess the quality of analysis fields generated through different assimilation systems, we use ozonesonde profiles measured by the INTEX Ozonesonde Network Study 2006 (IONS-6) (<http://croc.gsfc.nasa.gov/intexb/ions06.html>) (Thompson et al., 2007a, 2007b)) for the month of August, assuming  
295

**Table 1.** Timing results for GEOS-Chem free model runs using SMVGEAR and KPP chemistry, suboptimal Kalman filter, 3D-Var and 4D-Var data assimilations with diagonal background error covariance matrix for a 24 hour simulation starting 00:00 GMT August 1, 2006.

Experiment Description	CPU Time
Free model run, SMVGEAR chemistry solver	2 min 50 sec
Free model run, KPP chemistry solver	3 min 18 sec
Suboptimal Kalman filter with diagonal $\mathbb{P}^f$	3 min 08 sec
3D-Var with diagonal $\mathbb{B}$	3 min 57 sec
4D-Var with diagonal $\mathbb{B}$ (per model run)	16 min 51 sec

that these measurements provide values close to the true state of the atmosphere. There are 418 ozonesondes launched from 22 stations across North America as shown in the Figure 1. A detailed description of the number of ozonesondes launched per station with longitude and latitude information can be found in (Parrington et al., 2008). The ozonesonde observations are not used in data assimilation, and therefore provide an independent data set against which the analysis results are validated. Forecast scoring techniques using assimilated data as described in (Wu et al., 2008; Constantinescu et al., 2007c) do not provide a fair assessment of the quality of assimilation if the observation measurements involved high observation errors.



**Fig. 1.** Ozonesonde sounding stations (triangles) used during IONS06 campaign and AURA/TES satellite trajectory snapshots (dots) plotted over the global ozone distribution on August 1st, 2006.

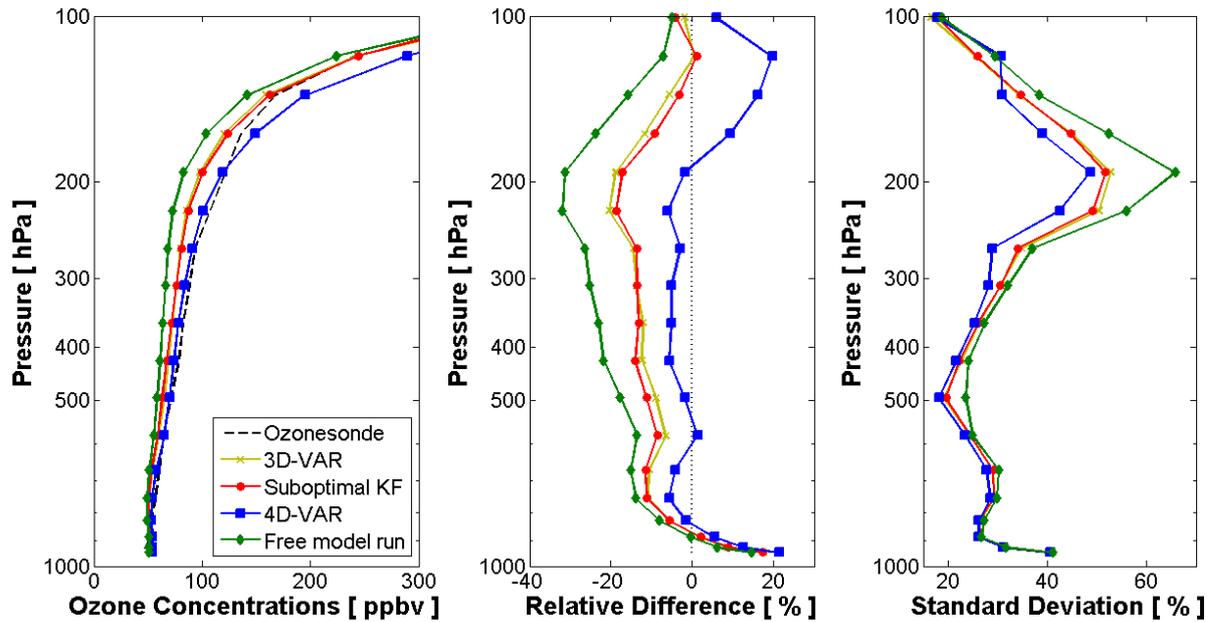
We first consider the case where the assimilation window length is five days. As per the property of  
305 sequential data assimilation algorithms, the model forecast is corrected as soon as an observation is avail-  
able. Ingesting observations every four simulation hours, we obtain an analysis field every four hours that  
accounts for the mismatch between the model prediction and the observations within that observation  
window. However, it is important to note that the model prediction at any observation window incor-  
porates implicitly the corrections from all previous observations. Thus, as we move forward in time, the  
310 analysis field agrees better with the true state of the atmosphere. 4D-Var on the other hand accumulates  
the forcing due to mismatch between model forecast and observations throughout the assimilation win-  
dow to produce an initial condition that, when evolved forward in time through the model, will best fit  
the observations. Therefore, in the case of sequential assimilation approaches, to obtain a stable analysis  
state that resembles the true chemical state of the atmosphere at a particular instant, we need to start the  
315 simulation days or months prior to that instant to benefit from earlier observations. 4D-Var is advanta-  
geous in situations where past observations are not available, as it provides the best estimate using only  
the observations available in the assimilation window under consideration.

We present in Figure 2, a comparison of analysis profiles obtained from different assimilation systems, and  
free GEOS-Chem model run against ozonesonde measurement data. The left panel shows vertical ozone  
320 profiles (concentrations against pressure levels); the model predictions are sampled at the locations and  
times of ozonesonde measurements available in the 5-day assimilation window. The differences between  
model results and ozonesonde data reflect model prediction errors; one error vertical profile is obtained  
for each ozonesonde launch. The center and right panels show the mean and the standard deviation of  
these errors. The plots provide an assessment of the quality of tropospheric ozone estimates given by the  
325 free model run, and by data assimilation systems based on suboptimal Kalman filter, 3D-Var and 4D-Var  
approaches. The errors also reflect the impact of TES profile retrievals on these assimilation systems.

It is evident from the plots in Figure 2 that 4D-Var provided the best estimate for lower and mid tro-  
posphere ozone concentrations. The relative difference between the mean ozone analysis field and the  
ozonesonde measurements were decreased to less than 4% up to 180 hPa as compared to 5-20% in cases  
330 of suboptimal KF and 3D-Var. The overestimation of ozone in the upper troposphere by 4D-Var is intrigu-  
ing and could be attributed to the fact that the TES averaging kernels are bringing stratospheric ozone  
values into the analysis at these altitudes. A detailed analysis on the information brought in by TES pro-  
file retrievals into the 4D-Var assimilation system at different pressure levels is provided in (Singh et al.,  
2010b).

335 Figure 3 provides the global tropospheric ozone distribution as estimated by GEOS-Chem free model run  
and different assimilation systems. The ozone concentration values are averaged over 10 GEOS-Chem  
levels (from the surface to about 370 hPa) for each longitude-latitude grid point on the horizontal domain.

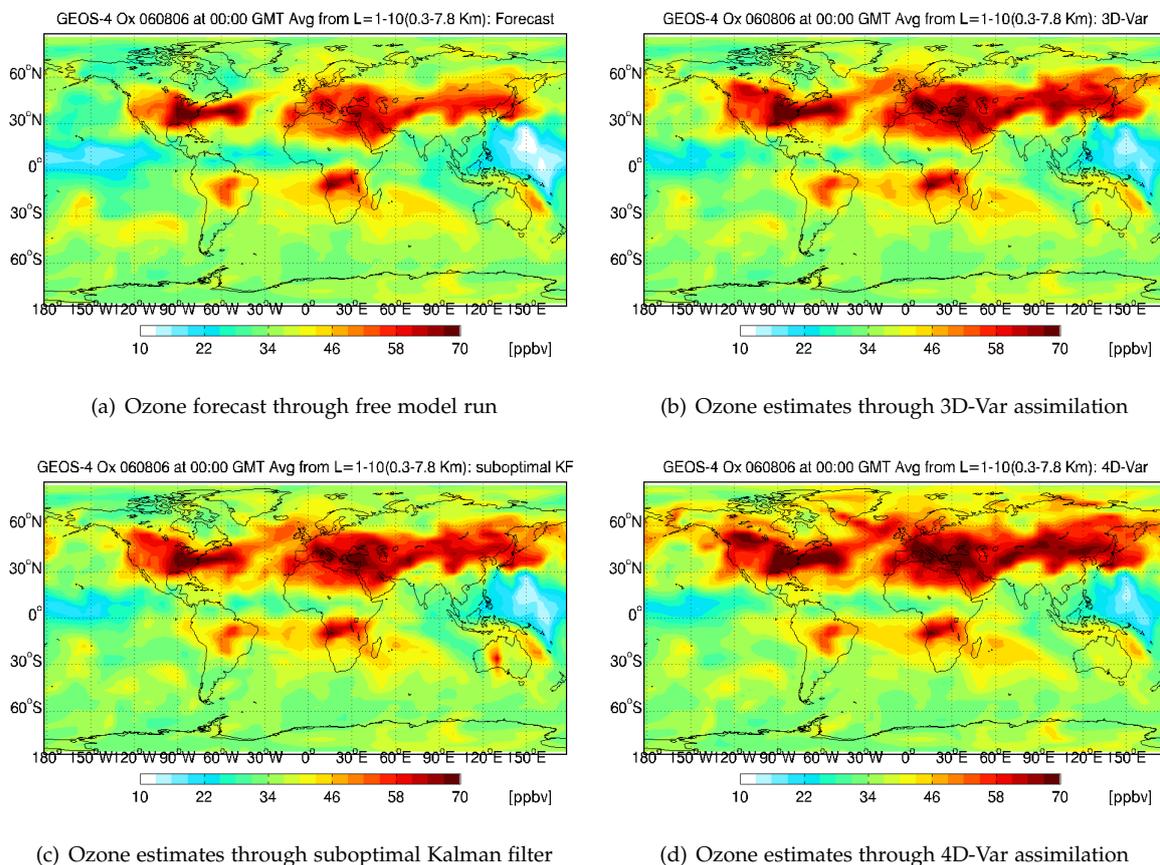
As seen in Figure 3, all the assimilation systems seem to have caused an increase in the tropospheric



**Fig. 2.** The impact of ozone profile retrievals from TES on data assimilation systems for GEOS-Chem. Left panel: mean ozone concentrations sampled at ozonesonde locations and times for 3D-Var, 4D-Var, suboptimal KF analyses and free model trajectories. Center panel: relative mean errors of predicted ozone concentrations with respect to ozonesonde measurements. Right panel: standard deviation of absolute values of errors with respect to ozonesonde measurements. The data is averaged over all ozonesonde launches. These plots were generated from 5 days simulation from 00:00 GMT August 1, 2006 to 00:00 GMT August 6, 2006 and compared against ozonesonde data available for the month of August.

ozone as compared to the model forecast with 4D-Var bringing the highest amount. The gain seems to be prominent in the 30° N to 60° N latitude region in case of suboptimal KF and 3D-Var, while it is extended up to 90° N in case of 4D-Var. For a clear demonstration of these changes, we provide in Figure 4, the plots of differences in the tropospheric ozone estimates through free model run and different assimilation systems.

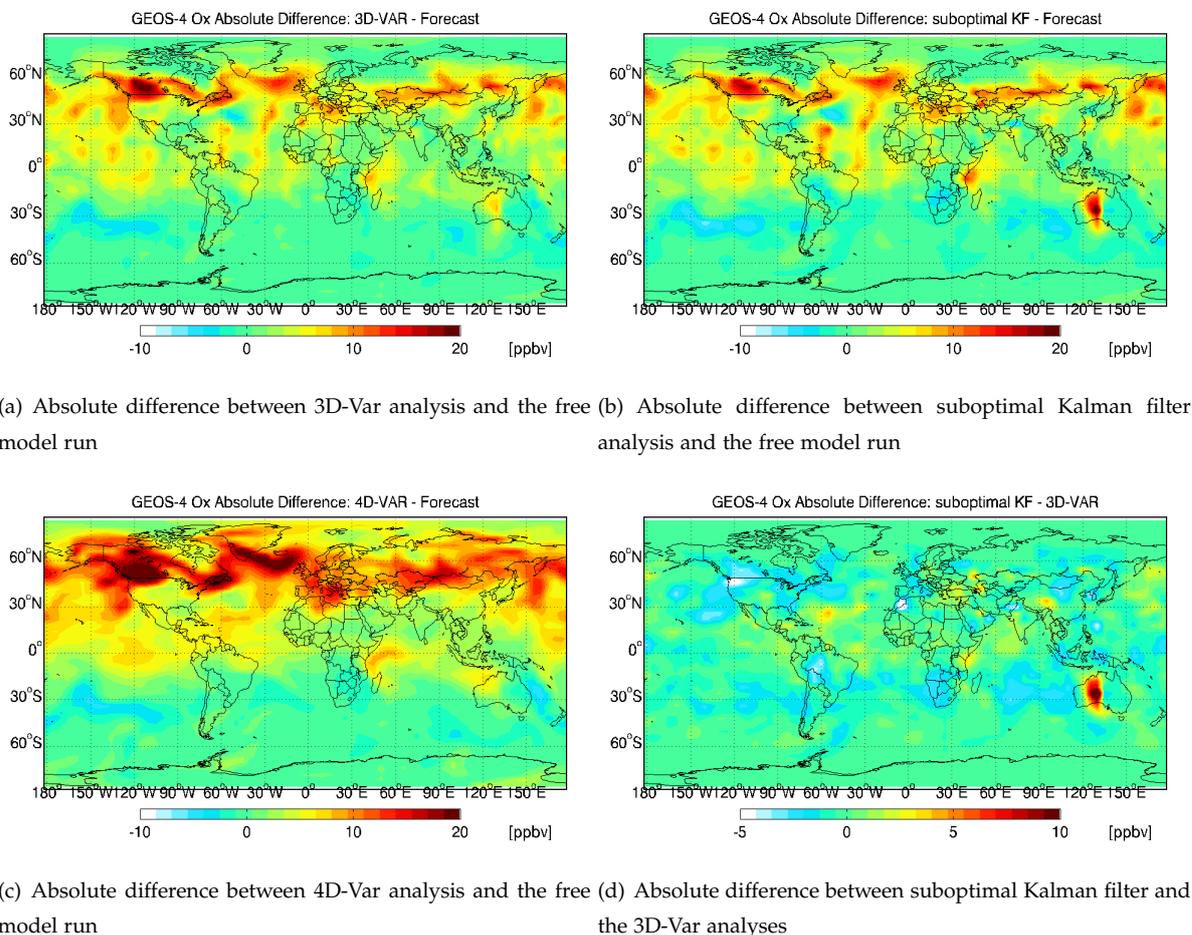
In Figure 4, panels (a) and (b) show that the structure of corrections in the ozone concentrations through 3D-Var and suboptimal KF data assimilation are quite similar. The reason behind such a structure is that these sequential algorithms bring in instantaneous corrections based solely on the mismatch between the model predictions and the observations in an observation window (analysis cycle). The localized corrections here are mostly along the Aura satellite orbit. Panel (c) on the other hand showcases the smoother correction profile of 4D-Var. In each 4D-Var optimization iteration, the cost function and gradients are accumulated for all the observation windows where the adjoint variable (gradient) is flown backwards in time as governed by the model adjoint equation. The corrections brought in by the optimization routine



**Fig. 3.** Global ozone distribution at 00:00 GMT on August 6, 2006 averaged over the first 10 GEOS-Chem vertical levels. Panels (a)-(d): Global tropospheric ozone estimates provided by free model run and suboptimal KF, 3D-Var, and 4D-Var data assimilation systems from a 5-day simulation.

therefore are no more localized. We also plot the difference in the analysis fields obtained by 3D-Var and suboptimal KF showcasing their close resemblance (panel (d)). Interestingly, there seems to be a localized overcorrection in the mid west Australian region by the suboptimal Kalman filter.

355 We next consider simulations with assimilation window length of 2 weeks. A longer assimilation window provides an insight into how ozone estimates due to assimilation evolve with time and if the corrections maintain structures similar to 5-day case. It also helps adjudge if model errors in GEOS-Chem cause any degradation in the assimilation systems, especially the strongly constrained 4D-Var. Similar to Figure 2, we present in Figure 5, a comparison of analysis profiles obtained from different assimilation systems against  
 360 ozonesonde measurement data. The plots reflect that the accuracy of suboptimal Kalman filter and 3D-Var assimilations start to differ with longer assimilation window. While suboptimal KF underestimates ozone concentrations in the lower and mid troposphere, it performs better than 3D-Var in the mid and upper tropospheric region. 4D-Var still provided the best ozone estimate of all the assimilation systems, and,



**Fig. 4.** Differences in global ozone concentrations at 00:00 GMT on August 6, 2006, the end of 5-day simulation, averaged over first 10 GEOS-Chem vertical levels. Panels (a)-(c): Differences between suboptimal KF, 3D-Var, and 4D-Var analysis fields and the model forecast (solution without data assimilation). Panel (d): Difference between suboptimal KF and 3D-Var analysis fields.

unlike the 5 days assimilation window length case, it performed well in the upper tropospheric region as well except near the tropopause. Panel (c) suggests that the standard deviation of 4D-Var analysis from the ozonesonde measurements stayed the least among all the assimilation systems. The relative difference between the mean ozone analysis field and the ozonesonde measurements were decreased to less than 4% up to 150 hPa as compared to 4-16% in cases of suboptimal KF and 3D-Var. With longer assimilation window, all the assimilation systems seem to have benefited from more observations being assimilated.

Figure 6 provides the global tropospheric ozone distribution as estimated by GEOS-Chem free model run and different assimilation systems. Similar to the 5 days assimilation window case, 4D-Var leads to the maximum increase in the tropospheric ozone.

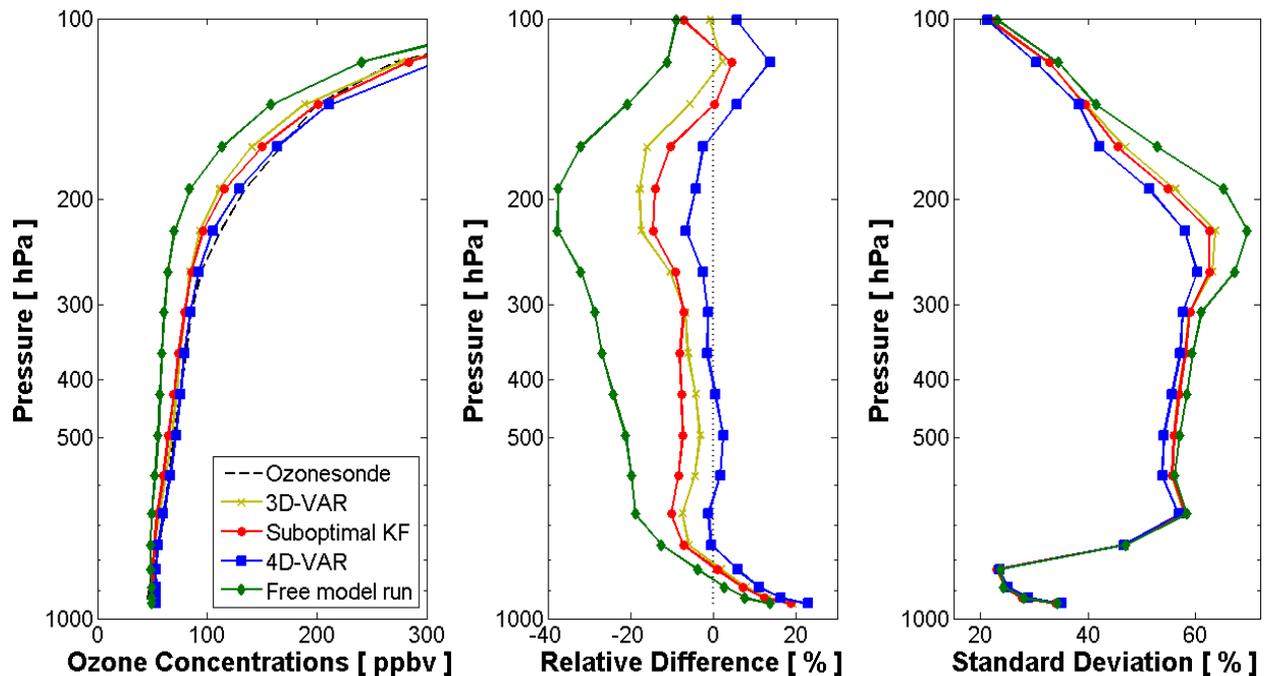
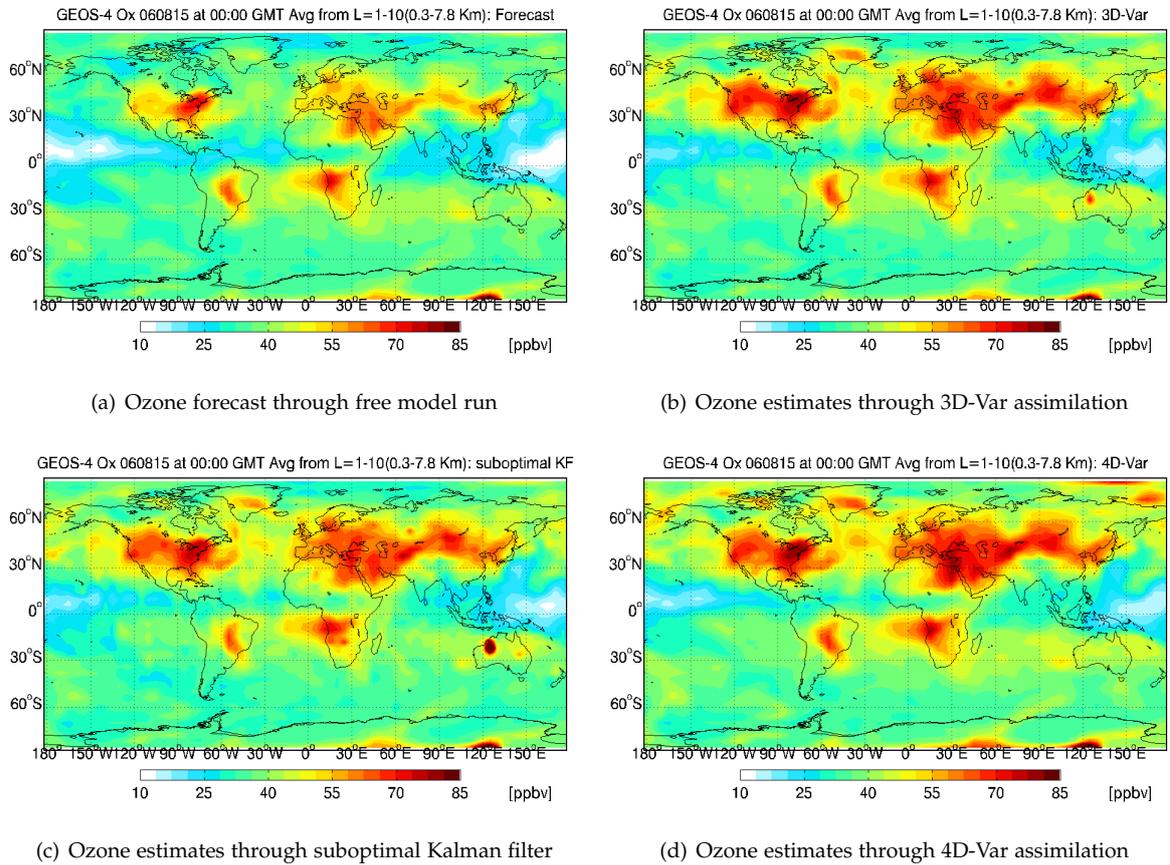


Fig. 5. The impact of ozone profile retrievals from TES on data assimilation systems for GEOS-Chem. Left panel: mean ozone concentrations at ozonesonde locations for 3D-Var, 4D-Var, suboptimal KF analyses and free model trajectories. Center panel: relative mean errors of predicted ozone concentrations with respect to ozonesonde measurements. Right panel: standard deviation of absolute values of errors with respect to ozonesonde measurements. The data is averaged over all ozonesonde launches. These plots were generated from 2 weeks simulation from 00:00 GMT August 1, 2006 to 00:00 GMT August 15, 2006 and compared against ozonesonde data available for the month of August.

Figure 7 showcases the structure of corrections in model predicted ozone through different assimilation systems. The ozone corrections are up to 20 ppbv, and are consistent among the three assimilation schemes. The localized correction structure in 3D-Var and suboptimal KF cases still persists with longer assimilation window. 4D-Var provides larger corrections with a significant increase in ozone concentrations in the 30° N to 90° N latitude region. Interestingly, the localized correction in the mid west Australian region which was not visible in the 3D-Var case for 5 days assimilation window case, seems to be prominent in longer assimilation, while, in the case of suboptimal KF, it has been accentuated.

Contrary to what was observed in (Wu et al., 2008) for the 4D-Var assimilation in Polair3D case where the accuracy of the ozone estimates decreased with increase in the assimilation window length, our findings show that the performance of the 4D-Var system improves with increase of the assimilation window. It seems that assimilating more meaningful observations keeps the effect of model errors from compromising the quality of assimilation. There is however one case where the accuracy of ozone estimates decrease with increase in assimilation window length for 4D-Var and that is when the model adjoints are inaccurate. We have studied this case in detail in (Singh et al., 2010c) and have utilized inaccurate gradients



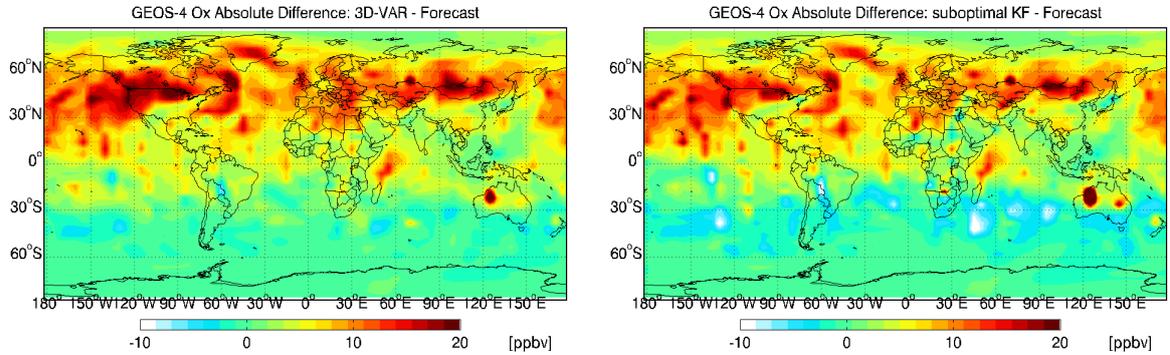
**Fig. 6.** Global ozone distribution at 00:00 GMT on August 15, 2006 averaged over the first 10 GEOS-Chem vertical levels. Panels (a)-(d): Global tropospheric ozone estimates provided by free model run and suboptimal KF, 3D-Var, and 4D-Var data assimilation systems from a 2-week simulation.

to work towards our benefit in terms of reducing significantly the memory and computational costs, still maintaining the quality of the analysis.

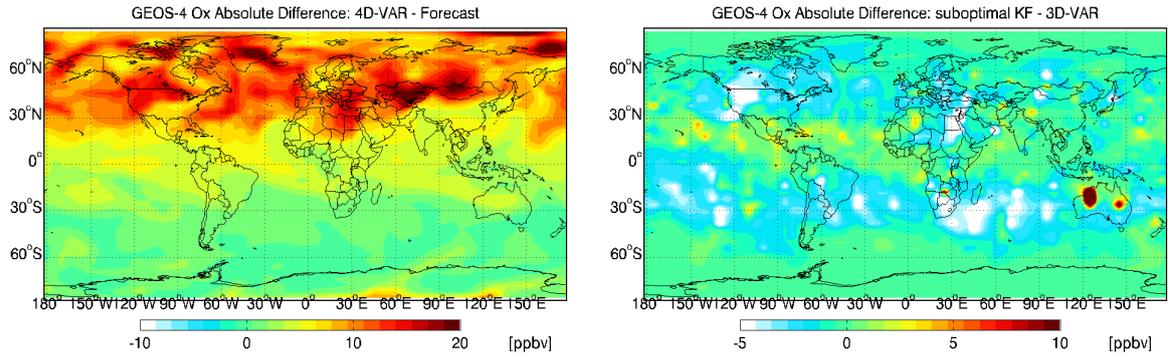
### 6.3 Comparison of 3D-Var and 4D-Var

390 As described in section 2, 3D-Var considers observations at successive intervals and a new analysis vector is generated every such interval. These 3D-Var corrections are forward in direction similar to the model flow and become closer to the reality with more sets of observations being considered. This behavior is well depicted in Figures 2 and 5. Up to authors knowledge, no such attempt has been made to measure the net effect of 3D-Var corrections at the initial time. Such a measurement would provide an opportunity

395 to assess the impacts of 3D and 4D-Var assimilations on the initial conditions. Provided below are the first set of formulations to calculate directly the difference of 3D and 4D-Var analyses, and 3D-Var analysis and



(a) Absolute difference between 3D-Var analysis and the free model run (b) Absolute difference between suboptimal Kalman filter analysis and the free model run



(c) Absolute difference between 4D-Var analysis and the free model run (d) Absolute difference between suboptimal Kalman filter and the 3D-Var analyses

**Fig. 7.** Differences in global ozone concentrations at 00:00 GMT on August 15, 2006, the end of 2-week simulation, averaged over first 10 GEOS-Chem vertical levels. Panels (a)-(c): Differences between suboptimal KF, 3D-Var, and 4D-Var analysis fields and the model forecast (solution without data assimilation). Panel (d): Difference between suboptimal KF and 3D-Var analysis fields.

initial condition at the start of an assimilation.

Let the background, the 3D-Var, and the 4D-Var solutions at  $t_k$  be

$$\mathbf{x}_k^B, \quad \mathbf{x}_k^{a(3D-Var)}, \quad \mathbf{x}_k^{a(4D-Var)},$$

respectively.

Define the cost function that measures the discrepancy between the 3D-Var and the 4D-Var solutions at all times

$$\mathcal{J}(\mathbf{x}_0^{a(4D-Var)}) = \frac{1}{2} \sum_{k=0}^N \left\| \mathbf{x}_k^{a(4D-Var)} - \mathbf{x}_k^{a(3D-Var)} \right\|^2$$

The gradient is

$$\nabla_{\mathbf{x}_0^{a(4D-Var)}} \mathcal{J}(\mathbf{x}_0^{a(4D-Var)}) = \sum_{k=0}^N \left( \frac{\partial \mathbf{x}_k^{a(4D-Var)}}{\partial \mathbf{x}_0^{a(4D-Var)}} \right)^T \cdot \left( \mathbf{x}_k^{a(4D-Var)} - \mathbf{x}_k^{a(3D-Var)} \right)$$

Therefore the gradient w.r.t. the initial conditions is given by the adjoint model forced by the differences  
400 between the 4D-Var and the 3D-Var solutions. The forward solution along which this adjoint is computed starts with the 4D-Var optimal initial condition.

We compute the adjoint sensitivities with respect to initial conditions and emissions

$$\nabla_{\mathbf{x}_0^{a(4D-Var)}} \mathcal{J}(\mathbf{x}_0^{a(4D-Var)}) \quad \text{and} \quad \nabla_{Emissions} \mathcal{J}(\mathbf{x}_0^{a(4D-Var)})$$

both of which are implemented in Geos Chem v7.

In a similar fashion, the cost function that measures the discrepancy between the 3D-Var and the model forecast at all times could be defined as

$$\mathcal{J}(\mathbf{x}_0^B) = \frac{1}{2} \sum_{k=0}^N \left\| \mathbf{x}_k^B - \mathbf{x}_k^{a(3D-Var)} \right\|^2$$

The respective gradient is

$$\nabla_{\mathbf{x}_0^B} \mathcal{J}(\mathbf{x}_0^B) = \sum_{k=0}^N \left( \frac{\partial \mathbf{x}_k^B}{\partial \mathbf{x}_0^B} \right)^T \cdot \left( \mathbf{x}_k^B - \mathbf{x}_k^{a(3D-Var)} \right)$$

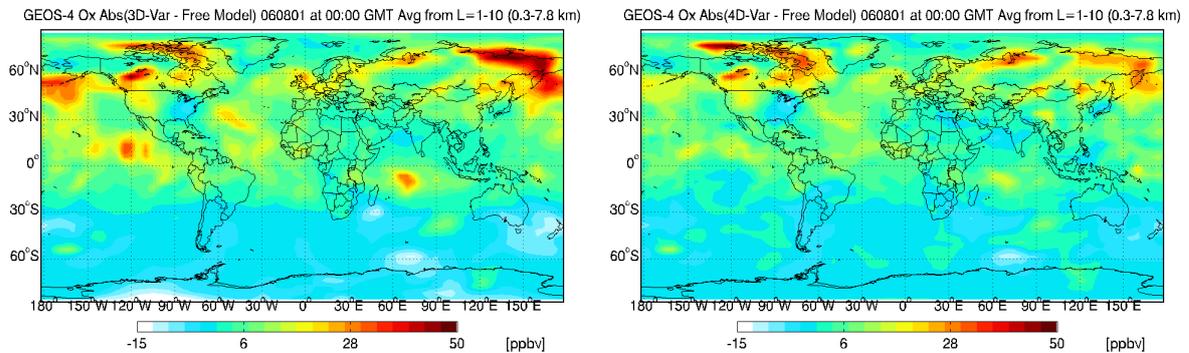
Therefore the gradient w.r.t. the initial conditions is given by the adjoint model forced by the differences  
405 between the model forecast and the 3D-Var. The forward solution along which this adjoint is computed starts with the model initial condition.

Figure 8 represents \*\*\*\*

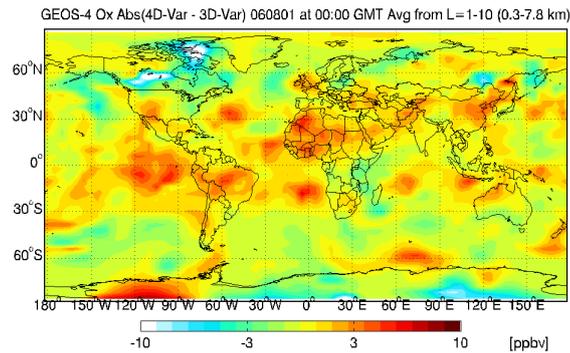
**Discuss why 4D-Var is better than 3D-Var and Multispecies\*\*\*\***

## 7 Conclusions

We have successfully implemented 3D-Var and 4D-Var data assimilation frameworks into GEOS-Chem v7  
410 adjoint package to carry out assimilations of TES ozone profiles. The current study involved estimation of global tropospheric ozone distribution. We provide the first set of results of direct comparison between 3D-Var, 4D-Var and suboptimal Kalman filter data assimilation systems. If using diagonal background error covariance matrix, suboptimal KF is computationally least expensive followed by 3D-Var. Memory and computational costs for 4D-Var is highest as it requires checkpointing dependent variables and an  
415 additional adjoint model run every iteration. The quality of estimated ozone from different assimilation systems were verified using ozonesonde measurements, an independent data set.



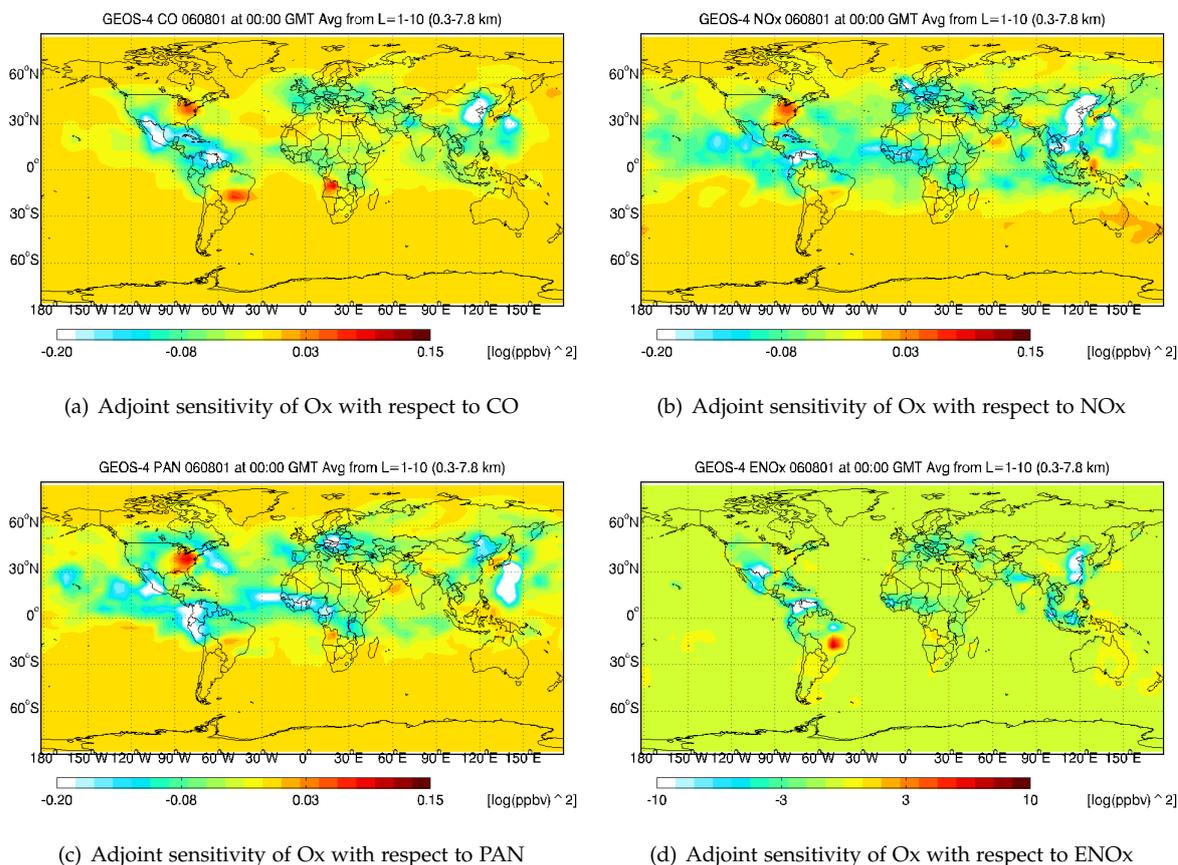
(a) Absolute Difference 3D-Var and Free model initial conditions  
 (b) Absolute Difference 4D-Var and Free model initial conditions



(c) Absolute Difference 4D-Var and 3D-Var optimal initial conditions

**Fig. 8.** Global ozone distribution at 00:00 GMT on August 15, 2006 averaged over the first 10 GEOS-Chem vertical levels. Panels (a)-(d): Global tropospheric ozone estimates provided by free model run and suboptimal KF, 3D-Var, and 4D-Var data assimilation systems from a 2-week simulation.

Two different assimilation window lengths were considered. Ozone estimates with all three assimilation systems have benefited from TES profile retrievals. Sequential assimilation methods, 3D-Var and suboptimal KF, perform similarly for smaller assimilation window decreasing the relative difference between mean analysis and ozonesonde measurements to about 5-20%. 4D-Var on the other hand brought down this difference to less than 4% for up to 180 hPa. For larger assimilation window length, the sequential approaches seem to differ a bit with suboptimal KF underestimating the ozone concentrations in the lower and mid troposphere, however performing better than 3D-Var in the mid and upper troposphere. The relative difference was brought down to 4-16% by sequential approaches while to less than 4% up to 150 hPa by 4D-Var. The structure of corrections in ozone concentration due to sequential assimilation methods seem to be localized along the satellite orbit, while smoother and well distributed in case of 4D-Var. The latitude region 30° N to 60° N seem to have gained maximum from all the assimilation systems extending



**Fig. 9.** Global ozone distribution at 00:00 GMT on August 1, 2006 averaged over the first 10 GEOS-Chem vertical levels. Panels (a)-(d): Global tropospheric ozone estimates provided by free model run and suboptimal KF, 3D-Var, and 4D-Var data assimilation systems from a 2-week simulation.

up to 90° N in case of 4D-Var.

The developed data assimilation frameworks and comparison results would enable users of GEOS-Chem  
 430 to obtain better estimates for all available trace gases at surface, tropospheric and stratospheric levels, depending on the memory and computational requirements. The framework is currently built to assimilate TES profile retrievals, however, it could easily be extended to use data from any instrument. Another interesting idea is to carry out assimilation with respect to surface emissions; a good estimate of emission rates is significantly important for policy making. Interesting and completely new applications such as  
 435 efficient information content estimation of observations and construction of full rank covariance matrices for variational data assimilation methods are studied in companion papers (Singh et al., 2010a,b,c).

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