Error Correlation Between CO2 and CO as Constraint for CO2 Flux Inversions Using Satellite Data

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Error correlation between CO\textsubscript{2} and CO as constraint for CO\textsubscript{2} flux inversions using satellite data

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Abstract. Inverse modeling of CO\textsubscript{2} satellite observations to better quantify carbon surface fluxes requires a chemical transport model (CTM) to relate the fluxes to the observed column concentrations. CTM transport error is a major source of uncertainty. We show that its effect can be reduced by using CO satellite observations as additional constraint in a joint CO\textsubscript{2}-CO inversion. CO is measured from space with high precision, is strongly correlated with CO\textsubscript{2}, and is more sensitive than CO\textsubscript{2} to CTM transport errors on synoptic and smaller scales. Exploiting this constraint requires statistics for the CTM transport error correlation between CO\textsubscript{2} and CO, which is significantly different from the correlation between the concentrations themselves. We estimate the error correlation globally and for different seasons by a paired-model method (comparing GEOS-Chem CTM simulations of CO\textsubscript{2} and CO columns using different assimilated meteorological data sets for the same meteorological year) and a paired-forecast method (comparing 48-h vs. 24-h GEOS-5 CTM forecasts of CO\textsubscript{2} and CO columns for the same forecast time). We find strong error correlations ($r^2>0.5$) between CO\textsubscript{2} and CO columns over much of the extra-tropical Northern Hemisphere throughout the year, and strong consistency between different methods to estimate the error correlation. Application of the averaging kernels used in the retrieval for thermal IR CO measurements weakens the correlation coefficients by 15% on average (mostly due to variability in the averaging kernels) but preserves the large-scale correlation structure. We present a simple inverse modeling application to demonstrate that CO\textsubscript{2}-CO error correlations can indeed significantly reduce uncertainty on surface carbon fluxes in a joint CO\textsubscript{2}-CO inversion vs. a CO\textsubscript{2}-only inversion.

1 Introduction

The joint Japan Aerospace Exploration Agency (JAXA), National Institute of Environmental Studies (NIES) and Ministry of the Environment (MOE) Greenhouse gases Observing SATellite (GOSAT or “Ibuki”) (http://www.jaxa.jp/projects/sat/gosat/index_e.html), launched in January 2009, is expected to greatly improve our knowledge of regional CO\textsubscript{2} sources and sinks by providing global measurements of CO\textsubscript{2} dry column mixing ratios ($X_{CO_2}$). It detects CO\textsubscript{2} by solar backscatter in the 1.61 and 2.06 $\mu$m bands, together with O\textsubscript{2} in the 0.76 $\mu$m band, resulting in $X_{CO_2}$ measurements with near-uniform sensitivity down to the surface. The National Aeronautics and Space Administration (NASA) Orbiting Carbon Observatory (OCO) was designed to provide global X\textsubscript{CO2} data with 0.3% (about 1 ppm) precision using the same channels (Crisp et al., 2004; Miller et al., 2007). Unfortunately, the February 2009 launch of OCO failed to reach orbit. Satellite observations of CO\textsubscript{2} from space are also available in the thermal IR from the AIRS (Crevoisier et al., 2003; Chahine et al., 2005, 2008; Tiwari et al., 2006; Maddy et al., 2008; Stow and Hannon, 2008), TES (Kulawik et al.,...
outflow from the TRACE-P aircraft campaign over the western Pacific in March–April 2001. Observed CO2 and CO concentrations showed correlation coefficients higher than 0.7 throughout the troposphere with distinct CO2/CO slopes depending on air mass origin (Suntharalingam et al., 2004). Palmer et al. (2006) found that exploiting this correlation in a joint CO2-CO flux inversion improved Asian CO2 flux estimates significantly relative to a CO2-only inversion. They assumed that the model transport error correlation between CO2 and CO would be identical to the observed correlation of concentrations, but as shown below this is not a good assumption in general.

Our aim in this paper is to develop an understanding of CO2-CO model transport error correlations as relevant to inversion of carbon fluxes from satellite observations. We present different methods for estimating the model error correlation and show that there is consistency and robustness across them. We examine the variability of the error correlation geographically, seasonally, and for satellite observations with different averaging kernels. We illustrate through a simple example how the error correlation can improve constraints on carbon fluxes.

2 Exploiting the CO2-CO error correlation in CO2 flux inversions

Consider the Bayesian inversion problem of constraining carbon fluxes from satellite measurements of the column mixing ratio \(X_{\text{CO2}}\). We follow the notation of Rodgers (2000). An ensemble of \(X_{\text{CO2}}\) observations \(y\), the observation vector) is used to optimize an ensemble of CO2 surface fluxes \(x\), the state vector) subject to prior knowledge of the fluxes (best estimate \(x_a\)). The state vector is related to the observation vector \(y\) through the CTM forward model:

\[
y = F(x) + \epsilon
\]

where \(\epsilon\) is the observational error, described in more detail below. The inverse model minimizes a cost function \(J(x)\) which is the least-squares sum of the observational error weighted by the observational error covariance matrix \(S = E(\epsilon \epsilon^T)\), where \(E\) denotes the expected value operator) and the a priori error \((x_a - x)\) weighted by the a priori error covariance matrix \(S_a = E(\epsilon_a \epsilon_a^T)\) (Rodgers, 2000):

\[
J(x) = (y - F(x))^T S^{-1} (y - F(x)) + (x - x_a)^T S_a^{-1} (x - x_a)
\]
vector as input. It includes contributions from instrument error ($\mathbf{e}_{I}$), representation error ($\mathbf{e}_{R}$), and forward model error ($\mathbf{e}_{M}$) (Heald et al., 2004; Engelen et al., 2002, 2006):

$$\mathbf{e} = \mathbf{e}_{I} + \mathbf{e}_{R} + \mathbf{e}_{M}$$

Components of the observational errors are not strictly independent. We will simplify here by ignoring their covariance. The error variances add quadratically (if the errors are independent). The instrument error includes measurement noise and retrieval error (Engelen et al., 2002, 2006). Smoothing error introduced by the averaging kernels of the satellite instrument is a source of retrieval error, but can be canceled by smoothing the CTM profiles with the same averaging kernels (Jones et al., 2003; Heald et al., 2004). Forward model error is the dominant source of observational error for CO observations from space (Heald et al., 2004) and may be dominant for CO$_2$ observations depending on data quality and averaging strategy (Baker et al., 2008).

The diagonal elements of the observational error covariance matrix $S$ are the variances of observational errors for the individual components of $y$. The off-diagonal elements are the corresponding observational error covariances, and can be obtained by scaling the error correlation coefficients with the corresponding square roots of error variances. One way to estimate the observational error variance is by the Relative Residual Error (RRE) method (Palmer et al., 2003; Heald et al., 2004). In this method, a forward model simulation using a priori fluxes ($\mathbf{K}_{x,a}$) is conducted and results compared to observation time series for individual domains (such as model grid squares). The mean differences for the time series (model bias) are assumed to be due to error in the a priori fluxes. The residual differences are taken to represent the observational error.

In a joint CO$_2$-CO inversion, the observational vector ($y$) consists of the CO$_2$ and CO observations, and the state vector ($x$) consists of CO$_2$ surface fluxes and CO sources. Coupling between the CO$_2$ and CO inversions occurs through the corresponding off-diagonal elements of the error covariance matrices. The observational error covariance matrix now takes the form (4), where $S_{CO_2}$ and $S_{CO}$ are the error covariance matrices for the single-species inversions:

$$S = \begin{pmatrix}
S_{CO_2} & E(\mathbf{e}_{CO_2}\mathbf{e}_{CO}^T) \\
E(\mathbf{e}_{CO}\mathbf{e}_{CO_2}^T) & S_{CO}
\end{pmatrix}$$

Since the instrument error for CO$_2$ and CO can be assumed independent, and the representation error can be assumed small (Heald et al., 2004), the observational error covariance between CO$_2$ and CO only comes from the model transport error. The CO$_2$-CO error covariance terms can be derived from the model error correlation coefficients by scaling by the square roots of model error variances of CO and CO$_2$. Although the model error variances obviously depend on the model, the correlation structure is more general as shown in Sect. 4.

In addition to observational error covariance, there could also be error correlation in the a priori emissions of CO$_2$ and CO due to the common combustion source. However, as shown by Palmer et al. (2006), this correlation is in fact very weak because the error in a priori CO emissions is mainly contributed by the emission factor (emission per unit fuel) rather than the activity rate (amount of fuel burned). A possible exception is biomass burning if uncertainty in activity rate exceeds a factor of two (Palmer et al., 2006). Palmer et al. (2006) found that this a priori error correlation was not useful in their inversion and we do not discuss it further here.

3 Estimating the CO$_2$-CO error correlation

We use two independent methods, which we call the paired-model and paired-forecast methods, to estimate the CO$_2$-CO model error correlation ($r_{M}$) and its geographical and seasonal distribution. In the paired-model method, we conduct otherwise identical CTM simulations of CO$_2$ and CO using different assimilated meteorological data sets for the same meteorological year. In the paired-forecast method, we compare 48-h vs. 24-h chemical forecasts of CO$_2$ and CO. The latter method has been used extensively for meteorological data assimilation and is often called the NMC method (Paris and Derber, 1992).

In both methods, each pair produces global 3-D concentration fields of CO$_2$ and CO for the same times that differ because of model transport error. A time series of model output for a given gridbox thus generates time series of concentration differences $\Delta$CO$_2$ and $\Delta$CO for the pair. We correlate the time series of $\Delta$CO$_2$ vs. $\Delta$CO for individual model grid boxes and individual months to estimate the corresponding CO$_2$-CO transport error correlation coefficients ($r_{M}$). The estimates may differ depending on the method and the data sets used, but by comparing the estimates obtained in different ways we can assess their robustness. The concentration fields are sampled as columns for the satellite overpass times and with and without instrument averaging kernels. Figure 1 shows typical averaging kernels for CO$_2$ from OCO (values for GOSAT are similar), CO from SCIAMACHY, and CO from AIRS. GOSAT, OCO and SCIAMACHY measure by solar backscatter in the near-IR and thus have near-unit sensitivity through the bulk of the atmosphere (i.e., nearly flat averaging kernels). AIRS, MOPITT, and TES measure in the thermal IR and have maximum sensitivity in the mid-troposphere. Infrared instruments can observe on both the night side and the day side of the orbit. On the dayside, all instruments observe at near 13:30 local time (“A-Train” constellation of satellites on the same orbit track) except for GOSAT (13:00), MOPITT (10:30) and SCIAMACHY (10:00).

For the paired-model method, we perform global simulations of CO$_2$ and CO using the GEOS-Chem CTM (v8-01-01. http://www-as.harvard.edu/chemistry/trop/geos) driven
by the same sources and sinks but different generations of Goddard Earth Observing System (GEOS) assimilated meteorological data produced by the NASA Global Modeling and Assimilation Office (GMAO). We compare simulations conducted with GEOS-5 vs. GEOS-4 for 2006, and GEOS-4 vs. GEOS-3 for 2001. GEOS-3, GEOS-4, and GEOS-5 differ in the underlying general circulation model, the methodology for data assimilation, and the data assimilated (Bloom et al., 2005; Rienecker et al., 2008; Ott et al., 2009). All GEOS data sets are 6-hourly (3-hourly for mixing depth and surface variables) and are regridded to 2° x 2.5° horizontal resolution for input to GEOS-Chem. The GEOS-Chem CO2 and CO simulations have been documented previously including extensive comparisons to observations (e.g., Sutharalingam et al., 2004; Duncan et al., 2007). Anthropogenic CO2 emissions are from Andres et al. (1996). Anthropogenic CO emissions are a combination of currently available inventories as used in Kopacz et al. (2009). Biomass burning emissions for both CO2 and CO are from the monthly Global Fire Emission Database version 2 (GFED2) inventory for the simulation year (van der Werf, 2006). Biofuel emissions of CO2 and CO are from Yevich and Logan (2003). All CO simulations use the same monthly 3-D OH concentration fields archived from a GEOS-Chem full-chemistry simulation (Fiore et al. 2003). Exchange of CO2 with the terrestrial biosphere follows the CASA balanced biosphere model with prescribed diurnal cycle (Randerson et al., 1997; Olsen and Randerson, 2004). Exchange of CO2 with the ocean follows Takahashi et al. (1997).

For the paired forecast method, we use GEOS-5 global chemical forecasts of CO and CO2 (1/2° x 2/3° horizontal resolution) for July 2008 generated by GMAO in support of the ARCTAS aircraft campaign (Jacob et al., 2009). These chemical forecasts were not custom designed for this paper. The CO simulation uses the same sources and OH fields as GEOS-Chem. The CO2 simulation differs in using daily averaged biospheric fluxes from CASA and no biomass burning. The 48-h and 24-h forecasts were sampled at 13:30 local time. No averaging kernels were applied.
4 CO₂-CO error correlation patterns

Figure 2 shows the global and seasonal patterns of the model error correlation between column CO₂ and column CO calculated with the paired-model method for GEOS-4 vs. GEOS-5 (2006) and GEOS-3 vs. GEOS-4 (2001). Both CO₂ and CO are sampled at 13:30 local time, corresponding to the A-Train overpass. Results are for actual columns (no averaging kernels) and would also apply to flat averaging kernels as obtained from the near-IR GOSAT and SCIAMACHY sensors (Fig. 1).

We find in Fig. 2 strong positive correlations ($r_M > 0.7$) prevailing during the non-growing season and in biomass burning regions. In January, 92%, 80% and 45% of the area north of 30° N has $r_M > 0.7$, 0.8, and 0.9, respectively. Similarly, strong negative correlations exist in the growing season in the absence of biomass burning. In July, 26%, 11%, and 3% of the area north of 30° N has $r_M < -0.6$, −0.7, and −0.8, respectively. Due to the magnitude and variability of the CASA balanced biospheric flux, the correlations are stronger and more coherent in winter than in summer. Error correlations extend far downwind of biomass burning and fossil fuel regions and over the scale of the Northern Hemisphere. Regions of weak model error correlations include but are not limited to regions of strong model error variances. Inverse model studies of CO₂ fluxes have pointed to model transport errors in northern extra-tropical land areas as a major limiting factor in flux optimization (Gurney et al., 2002, 2003, 2004; Baker et al., 2006). The strong CO₂-CO error correlations in that region offer promise for improvements through a joint CO₂-CO inversion.

We also find in Fig. 2 that error correlation patterns are very similar for the GEOS-4/GEOS-5 and GEOS-3/GEOS-4 pairs. The robustness of error correlation patterns indicates that the directions of the general gradients of column CO and CO₂ are similar between the two sets of models. Stronger positive correlation over Indonesia and the Indian Ocean in October for the GEOS-4/GEOS-5 pair can be explained by stronger biomass burning in Indonesia in 2006 (Logan et al., 2008). We find that correlation magnitudes and patterns are insensitive to time of day (not shown), even though the CO₂ surface flux changes sign between day and night during the growing season. This is consistent with observations by Washenfelder et al. (2006) that CO₂ columns (as opposed to surface concentrations) show little diurnal variability.

Figure 3 shows the model error correlations obtained from the paired-forecast method for July 2008. As in Fig. 2, no averaging kernels are applied. Despite the differences in meteorology, emissions, sampling time, and method (Sect. 3), the large scale model error correlations are very similar to those in Fig. 2. The error structure is finer because of the higher spatial resolution ($1/2° \times 2/3°$ vs. $2° \times 2.5°$).

Figure 4 shows the error correlation results including averaging kernels for OCO CO₂ and AIRS CO, as obtained by the paired-model method for January and July 2006. For OCO we use fixed land and ocean averaging kernels taken from Fig. 1; these do not significantly modify the CO₂ columns. Similar averaging kernels apply for GOSAT. For AIRS, we used the averaging kernels for each CO retrieval (AIRS data version 5, http://disc.sci.gsfc.nasa.gov/AIRS; McMillan et al., 2005), and averaged the resulting CO columns over the $2° \times 2.5°$ model grid. Application of AIRS averaging kernels degrades the error correlation because the CO₂ and CO columns are now observed with different and variable vertical weighting factors. Yet we find that the large-scale correlation structures are preserved (Fig. 4) with the correlation coefficients reduced on average by 15% (of which 9% is due to averaging kernel variation) relative to the results of Fig. 2.

In their previous joint CO₂-CO inverse analysis using TRACE-P aircraft data, Palmer et al. (2006) assumed that the CO₂-CO observational error correlation was the same as the correlation of concentrations. If this assumption was approximately correct it would greatly facilitate the generation of error correlation statistics. We examine its validity in Fig. 5 by showing the correlations between column CO₂ and column CO (without averaging kernels) simulated by GEOS-Chem for 2006. These can be compared to the error correlations shown in the left panels of Fig. 2. We find the same general patterns of strong positive correlations in combustion source regions, and strong negative correlations in regions of photosynthesis activity. But there are also large differences, particularly in the transition seasons (e.g., April). For the Palmer et al. (2006) conditions of Asian outflow over the NW Pacific in April, we find that the transport errors are much more strongly correlated than the columns themselves, which would increase the utility of the joint CO₂-CO inversion for constraining carbon fluxes. Overall, the differences between Fig. 2 and 5 are sufficiently large and complex that correlation of concentrations should not be used as error correlations in general.

![Fig. 3. Model error correlation coefficients between CO₂ and CO columns calculated with the paired-forecast method for July 2008 at 1/2° × 2/3° resolution. No averaging kernels were applied. Results can be compared to the July panels of Fig. 2.](http://www.atmos-chem-phys.net/9/7313/2009/)
5 Demonstration of error reduction in a CO2 flux inversion

We demonstrate the benefit of using CO2-CO model error correlations in CO2 flux inversions with a simple example. Pseudo data of column CO2 and column CO with OCO-like averaging kernels (Fig. 1, OCO-land) were generated along A-train orbits using 2° × 2.5° GEOS-Chem CO and CO2 simulations driven by GEOS-4 meteorology. Model error variances and correlation derived from the paired-model (GEOS-5 vs. GEOS-4) method were used to specify the observational error covariance matrix S. Since OCO averaging kernels essentially show uniform vertical sensitivity, we used the GEOS-5 vs. GEOS-4 correlation map without averaging kernels (Fig. 2). We assumed that the forward model error is the only source of observational error (\( \epsilon = \epsilon_M \)), and ignored spatial and temporal error correlations.

\[ \hat{S} = \left( K^T S^{-1} K + S_n^{-1} \right)^{-1} \]
The a posteriori errors are the square roots of the diagonal terms of $\hat{S}$. Figure 7 shows the ratios of a posteriori CO$_2$ flux errors between the CO$_2$-CO and CO$_2$-only inversion. In January, when strong positive model error correlations prevail in the Northern Hemisphere, a posteriori CO$_2$ combustion and biosphere flux uncertainties from the CO$_2$-CO inversion are 39–82% of those in the uncorrelated inversion, with a median of 56% for combustion sources and 69% for biosphere fluxes. In July, they typically decrease by 10–30% relative to the CO$_2$-only inversion. Larger improvements in January compared to July are due to generally larger absolute values of correlation coefficients and greater spatial coherence. Differences in improvement between source regions in Fig. 7 generally reflect differences in the strength of the correlation in Fig. 2. The present example indicates significant promise. Our example involves several simplifications such as neglecting instrument and representation errors, neglecting spatial and temporal correlations, and using the same model for both pseudo data and inversion. These simplifications may influence the benefits of the joint CO$_2$-CO inversion (Chevallier, 2007). A more extensive study will be needed to better understand their effects.

6 Conclusions

We explored the potential of using CO$_2$-CO transport error correlations to improve inversions of CO$_2$ surface fluxes from satellite observations of CO$_2$ columns. CO columns can be measured from space with high relative precision. Because of its relatively short lifetime, CO is more sensitive than CO$_2$ to model transport errors on synoptic and smaller scales. A joint CO$_2$-CO inversion including model transport error correlation could improve the inversion of CO$_2$ surface fluxes relative to a CO$_2$-only inversion. In this paper we showed how the CO$_2$-CO error correlation structure can be determined robustly on a global scale, and we presented an illustrative example to demonstrate its value for CO$_2$ flux inversions.

We used two independent methods to characterize the model transport error correlation for CO$_2$ and CO columns as measured from space. The first is a paired-model method in which we conducted CTM simulations of CO$_2$ and CO with the same sources and sinks for the same meteorological year but different assimilated meteorological data sets. We applied this method to GEOS-5 vs. GEOS-4 data sets for 2006 and to GEOS-4 vs. GEOS-3 data sets for 2001. The second is a paired-forecast method (often called the NMC method) in which we compared 48-h vs. 24-h CTM forecasts of CO$_2$. 
and CO for the same forecast times. We find that these different methods and data sets yield very similar large scale error correlation patterns. Strong positive error correlations are found over much of the Northern Hemisphere during the non-growing season, and over biomass burning regions of the tropics extending to the oceans far downwind. Strong negative error correlations are found over much of the Northern Hemisphere during the growing season. The correlations are largely insensitive to the time of day of the observations.

Satellite measurements by solar backscatter in the near-IR (OCO and GOSAT for CO₂, SCIAMACHY for CO) have vertically uniform sensitivities, but thermal-IR instruments (MOPITT, AIRS, TES, IASI) have greatest sensitivity in the mid-troposphere. We therefore examined the model error correlation of CO₂ with CO including variable AIRS averaging kernels for individual scenes as observed in 2006. We find that the CO₂-CO error correlation coefficients decrease by 15%, mostly due to variations in averaging kernels, but the large-scale correlation structure is preserved.

We examined whether simple correlation of concentrations could offer a suitable approximation to the error correlation since it is much easier to derive and can be constrained by observations. We find that the general patterns are often similar between the two but there are also sufficiently large differences to make the approximation inadequate.

We illustrated the potential of exploiting CO₂-CO error correlations in a joint CO₂-CO flux inversion with a simple example based on 14 days of pseudo satellite observations. We find that a posteriori CO₂ flux uncertainties are substantially reduced, implying significant improvement in the CO₂ flux inversion. Inversions using actual satellite observations are subject to measurement noise and model biases that complicate greatly the interpretation of results relative to our idealized example. Further work will be needed to demonstrate the value of CO₂-CO error correlations as constraints on CO₂ fluxes in real world applications.

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