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Assessing 5 years of GOSAT Proxy XCH₄ data and associated uncertainties

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Abstract

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Assessing 5 years of GOSAT Proxy XCH₄ data and associated uncertainties

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The uncertainty relating to the model XCO₂ component of the Proxy XCH₄ is assessed through the use of an ensemble of XCO₂ models. While each individual XCO₂ model is found to agree well with the TCCON validation data ($r = 0.94\text{--}0.97$), it is not possible to select one model as the best from our comparisons. The median XCO₂ value of the ensemble has a smaller scatter against TCCON (a standard deviation of 0.92 ppm) than any of the individual models whilst maintaining a small bias (0.15 ppm). This model median XCO₂ is used to calculate the Proxy XCH₄ with the

maximum deviation of the ensemble from the median used as an estimate of the uncertainty.

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1 Introduction

Atmospheric methane (CH₄) contributes significantly to the Earth's radiative forcing budget (Myhre et al., 2013), making it the second most important anthropogenic greenhouse gas after carbon dioxide (CO₂). The major sources of atmospheric methane include wetland emission, rice production, enteric fermentation (cattle), termites, biomass burning, fossil fuel production, and waste (Bousquet et al., 2006). There remains, however, a large degree of uncertainty on the magnitude of these individual sources (Kirschke et al., 2013).

The lifetime of CH₄ in the atmosphere is mainly controlled by its reaction with the hydroxyl free radical (OH), resulting in an atmospheric lifetime of approximately 9 years (Prather et al., 2012). Given its long atmospheric lifetime, there is a need for long-term global measurements to fully understand how the atmospheric distribution of CH₄ is evolving with time. Indeed, recent unexpected variability in the atmospheric growth rate of methane has emphasised gaps in our current understanding (Rigby et al., 2008; Dlugokencky et al., 2009; Nisbet et al., 2014).

In order to begin to understand the spatio-temporal distribution of atmospheric methane, regular global satellite observations of CH₄ can be coupled with highly precise but geographically sparse surface concentration data. Through the combination of both data sources, the large uncertainties related to the upscaling of surface concentration data can be minimised whilst also obtaining information in remote regions where surface measurements are not available.

Various studies have demonstrated the utility of such space-borne measurements in determining the regional surface fluxes of methane using data from the SCIAMACHY (Bergamaschi et al., 2007, 2009, 2013; Houweling et al., 2014) and Greenhouse gases Observing SATellite (GOSAT) (Fraser et al., 2013; Cressot et al., 2014; Monteil et al., 2013; Alexe et al., 2015) instruments.

The SCIAMACHY instrument operated onboard ENVISAT and provided a 9-year record (2003–2012) of global methane total column observations (Schneising et al., 2011; Frankenberg et al., 2011). The continuation of this time series of space-based observations was ensured by the launch of the first dedicated greenhouse gas measuring satellite, the Japanese GOSAT, in 2009 (Yokota et al., 2009). GOSAT provides global coverage with a 3-day repeat cycle and was designed with the intention of characterising continental-scale sources and sinks.

In a previous work (Parker et al., 2011) we presented the first year of our global short-wave infrared (SWIR) measurements of the dry-air column-averaged mole fraction of CH₄ (XCH₄) from the GOSAT mission using the “proxy” retrieval approach. This data product has subsequently been developed (Buchwitz et al., 2013) and validated (Dils et al., 2014) as part of the ESA Climate Change Initiative Greenhouse Gas project and we now report an assessment of the

full 5-year data set for version 5.0 of the University of Leicester GOSAT Proxy XCH₄ data product.

This work is motivated by the desire to better understand the uncertainty characteristics of the Proxy XCH₄ data for use within flux inversion systems, especially relating to uncertainties introduced by the model XCO₂.

In Sect. 2 we describe the retrieval approach, including details of the updates since the original version of the University of Leicester GOSAT Proxy XCH₄ data (Parker et al., 2011). In Sect. 3 we compare both the Proxy XCH₄ and the XCH₄/XCO₂ ratio against the ground-based validation data. In Sect. 4 we assess the CO₂ model component of the Proxy XCH₄ for the first time, with Sect. 5 then discussing the associated uncertainty of the final Proxy XCH₄ product and its utility in constraining surface fluxes within an inversion framework. Finally, we conclude the paper in Sect. 6 and provide recommendations for data users.

2 University of Leicester GOSAT Proxy XCH₄ retrieval updates

The University of Leicester GOSAT Proxy XCH₄ retrieval utilises the Orbiting Carbon Observatory (OCO) “full physics” retrieval algorithm, developed for the original NASA OCO mission to retrieve XCO₂ (dry-air column-averaged mole fraction of CO₂) from a simultaneous fit of SWIR O₂ and CO₂ bands and has subsequently been modified to operate on GOSAT spectral data.

Full details of the OCO retrieval algorithm can be found in O'Dell et al. (2012). In short, the retrieval algorithm utilises an iterative retrieval scheme based on Bayesian optimal estimation to estimate a set of atmospheric, surface, and instrument parameters from the measured spectral radiances, referred to as the state vector. The state vector of our retrieval consists of 20-level profiles for CH₄ and CO₂ volume mixing ratios (vmr), profile scaling factors for H₂O vmr, and temperature, surface albedo, and spectral dispersion.

Rather than perform the “full physics” retrieval as typically used for CO₂ (Connor et al., 2008; Boesch et al., 2011), an alternative approach is possible for CH₄, the so-called “proxy” method. First used for the retrieval of XCH₄ from SCIAMACHY (Frankenberg et al., 2006), this approach uses the fact that there exists CO₂ and CH₄ spectral signatures located close together at around 1.6 μm and hence the majority of atmospheric scattering and instrument effects will be similar between the two bands. The ratio of the retrieved XCH₄ / XCO₂ should cancel modifications to the length of the light path that are experienced due to scattering (Butz et al., 2010), with the CO₂ effectively acting as a “proxy” for the unknown light-path enhancements. As CO₂ is known to vary much less than CH₄, the final XCH₄ product can be obtained by multiplying this XCH₄ / XCO₂ ratio by a model CO₂ value, typically taken from a global chemistry transport model (Eq. 1).

$$\text{Proxy}_{\text{XCH}_4} = \frac{[\text{XCH}_4]}{[\text{XCO}_2]} \times \text{Model}_{\text{XCO}_2} \quad (1)$$

The “proxy” retrieval approach has various advantages over the full physics approach (Schepers et al., 2012). Because there is no reliance on an explicit a priori knowledge of the aerosol distribution, the proxy approach is more robust in the presence of aerosols and also far less sensitive to instrumental issues or inconsistent radiometric calibration between the spectral bands than is the case for the full physics approach. Additionally, as moderate scattering from aerosols will be cancelled out and still result in an accurate retrieval of XCH₄, the number of successful soundings for the proxy approach is typically much higher than for the full physics approach which requires far stricter post-filtering. This leads not only to more soundings in general but also to more soundings over regions where very little full physics data may be available, such as in the tropics.

The main disadvantage with the proxy approach is that it is reliant on an accurate, unbiased model XCO₂ data set to convert the XCH₄ / XCO₂ ratio back into an XCH₄ quantity; otherwise errors relating to the model XCO₂ may be folded into the final XCH₄ result. Here we present assessments of the different uncertainties to determine the importance of this aspect of the Proxy XCH₄ data.

We process the latest versions of the GOSAT Level 1B files (version 161.160) acquired directly from the NIES Large Volume Data Server and apply the recommended radiometric calibration and radiometric degradation correction as per Kuze et al. (2014).

For the spectroscopic inputs we use v4.2.0 of the OCO line lists with CH₄ taken from the Total Carbon Column Observing Network (TCCON) line lists (version “20120409”). The a priori pressure, temperature, and water vapour is taken from the ECMWF ERA-Interim data (Dee et al., 2011). For the CO₂ a priori we use the MACC-II CO₂ inversion (v13r1) and for the CH₄ we use the MACC-II CH₄ inversion (v10-SINOAA, using 2012 data for 2013) but here we adjust the stratospheric methane using a specialised full chemistry run (run ID 563) of the TOMCAT stratospheric chemistry model from the University of Leeds (Chipperfield, 1999). This TOMCAT model run has been validated against ACE-FTS observations and was found to provide a more accurate representation of the stratosphere.

The spectral noise is estimated from the standard deviation of the out-of-band signal. Spectra over ocean or with a signal-to-noise ratio (SNR) below 50 are removed. Cloud-contaminated scenes are removed by the comparison of a clear-sky surface pressure retrieval from the O₂ A-band to the ECMWF surface pressure for the relevant measurement time and location. A scene is determined to be cloudy when the retrieved surface pressure differs by more than 30 hPa from the estimated ECMWF surface pressure. This relatively loose threshold is allowed as the proxy retrieval

approach remains relatively robust in the presence of near-surface clouds. The average difference between our retrieved surface pressure and ECMWF after filtering for cloud is approximately 3 hPa with a standard deviation of below 10 hPa, with the offset from 0 hPa being attributed to spectroscopic uncertainties in the O₂ cross-sections. The Proxy XCH₄ retrieval is performed for all scenes that are deemed to be sufficiently cloud free.

After filtering for signal-to-noise, cloud, and data quality we are left with 1 032 760 XCH₄ retrievals over land between April 2009 and December 2013. Figure 1 shows global maps of the Proxy XCH₄ for each season and compares it to the MACC-II model XCH₄ data. Both model and observation show the XCH₄ variability in time and space, in particular with the large emissions of methane from wetland and rice cultivation over India and S.E. Asia.

3 Validation of the Proxy XCH₄ and XCH₄/XCO₂ ratio

This section presents the validation of the University of Leicester GOSAT Proxy XCH₄ v5.0 data through comparison to observations from the ground-based TCCON. In addition, for the first time the XCH₄ / XCO₂ ratio itself, the core component of the Proxy XCH₄ data, is validated against the corresponding TCCON data.

TCCON is a global network of ground-based high-resolution Fourier transform spectrometers recording direct solar spectra in the near-infrared spectral region (Wunch et al., 2011a). The TCCON data are calibrated to World Meteorological Organization (WMO) standards by calibration against aircraft measurements (Wunch et al., 2010). Although it should be noted that this aircraft calibration does not measure the whole column, the TCCON data are the standard against which current satellite observations of greenhouse gases are validated (Cogan et al., 2012; Wunch et al., 2011b; Dils et al., 2014).

To date, all previous validation of satellite greenhouse gas observations against TCCON has used TCCON data that were affected by instrumental biases relating to a laser sampling error which resulted in an XCO₂ error of approximately 0.26 % (1 ppm) (Messerschmidt et al., 2010). Although the corresponding XCH₄ error was not quantified, it is expected that it would be of similar magnitude (i.e. 1 part in 400). The latest, recently released, version of the TCCON data (GGG2014) incorporates a correction for the laser sampling errors and any remaining bias is expected to be small.

Figure 2 shows the GGG2014 TCCON XCH₄ data and the Proxy XCH₄ plotted as time series for each TCCON site. The mean GOSAT–TCCON difference, the standard deviation of the GOSAT–TCCON difference, the correlation coefficient, and the number of soundings are all provided for each site.

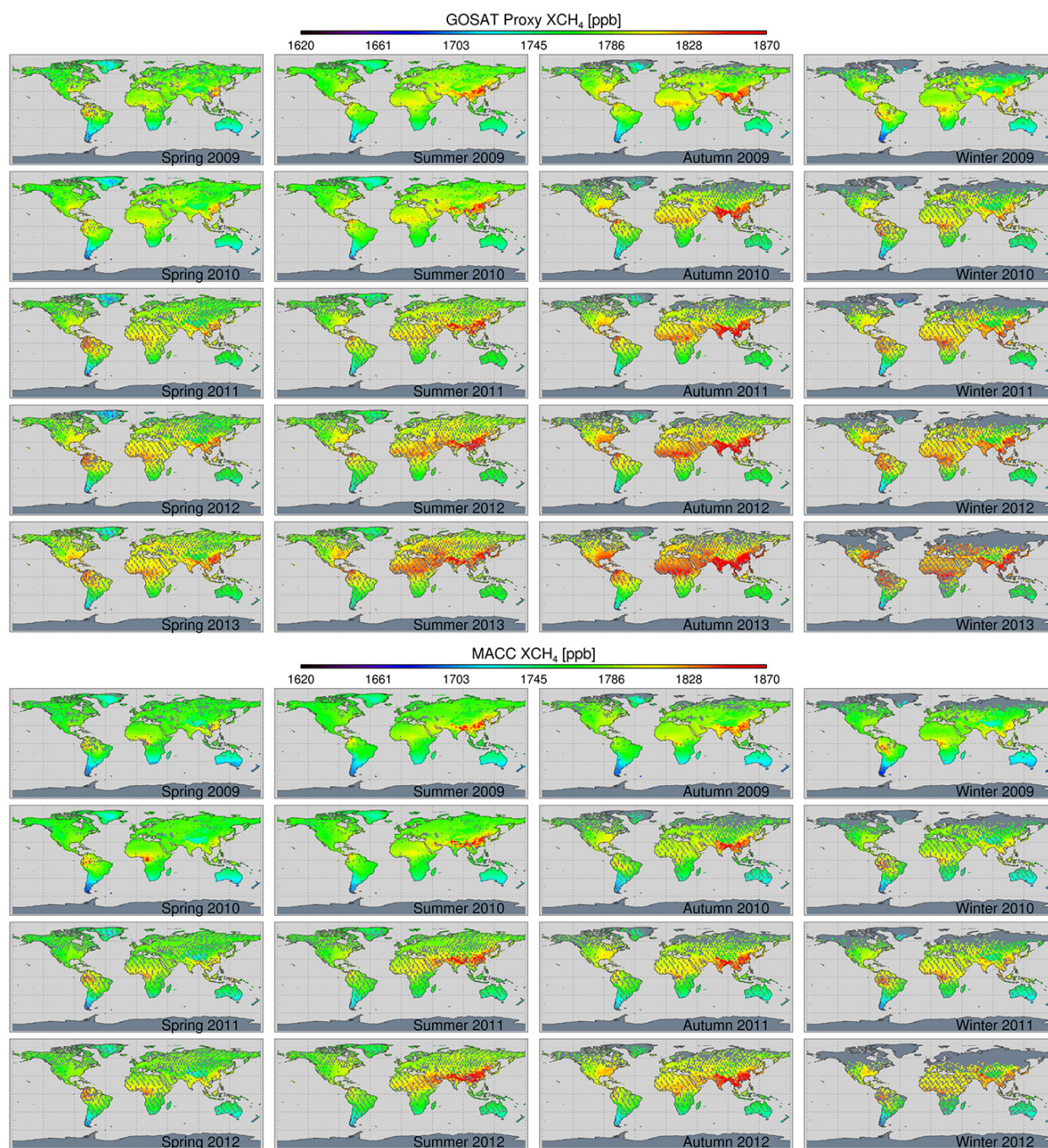


Figure 1. Seasonal global maps of the University of Leicester GOSAT Proxy XCH₄ (top) and the MACC-II (bottom) model XCH₄ data (v10-S1NOAA). Both model and observation show the XCH₄ variability in time and space, in particular with the large emissions of methane from wetland and rice cultivation over India and S.E. Asia. Note that GOSAT changed their pointing pattern in August 2010 from five across-track points to three across-track points, resulting in a change in spatial coverage.

Figure 3 (top) shows the correlation between the GGG2014 TCCON XCH₄ data and the Proxy XCH₄ values within $\pm 5^\circ$ of each TCCON site and a temporal coincidence of ± 2 h. It should also be noted that for all TCCON comparisons, the difference inherent in the data due to using different a priori has been compensated for (as discussed in

Rodgers (2000), by replacing the a priori used in the GOSAT retrievals with the TCCON a priori after the retrieval has been performed) which typically increases the GOSAT XCH₄ data by an average of between 0 and 5 ppb with the larger effect seen at more northerly TCCON stations. We use all TCCON sites where version GGG2014 has been processed at the time

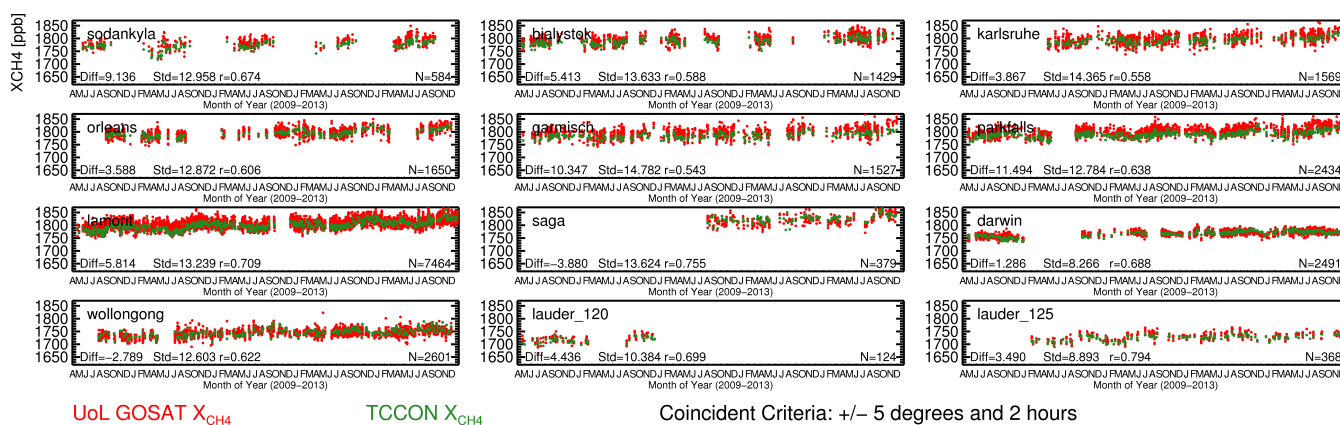


Figure 2. GGG2014 TCCON XCH₄ data and the Proxy XCH₄ plotted as time series for each TCCON site. The mean GOSAT–TCCON difference, the standard deviation of the GOSAT–TCCON difference, the correlation coefficient, and the number of soundings are all provided for each site.

of writing that contain data during the GOSAT time period (2009–2014). This results in 11 TCCON stations ranging from Sodankylä, Finland, at 67.4° N to Lauder, New Zealand, at 45.0° S. The correlation between the GOSAT and TCCON data is reasonable/good across all sites, ranging from 0.54 at Karlsruhe to 0.79 at Lauder with an overall correlation coefficient of 0.87 between 22 619 points. The overall bias is found to be 4.8 ppb with an overall single measurement precision of 13.4 ppb (ranging from 8.3 ppb at Darwin to 14.9 ppb at Garmisch). The station-to-station bias, which is an indication of the relative accuracy, is calculated as the standard deviation of the individual site biases and is found to be just 4.2 ppb.

In addition to the validation of the Proxy XCH₄ data, we also present for the first time the validation of the XCH₄ / XCO₂ ratio. This ratio is the quantity directly retrieved from the satellite measurement, is independent of any model XCO₂, and has recently itself been used directly within a flux inversion study (Fraser et al., 2014). The correlation coefficient across all stations is found to be 0.88 (ranging from 0.6 at Wollongong to 0.88 at Sodankylä) with a mean bias of 0.014 ppb ppm⁻¹ and a single-sounding precision of 0.033 ppb ppm⁻¹ (ranging from 0.20 ppb ppm⁻¹ at Darwin to 0.037 ppb ppm⁻¹ at Garmisch). The statistics for the XCH₄ / XCO₂ ratio are therefore comparable to those of the Proxy XCH₄ itself, suggesting that the majority of the variation is from the satellite retrieval itself and not introduced by the model XCO₂. The next section investigates this aspect in more detail.

4 Assessing the CO₂ model ensemble component

In Sect. 3 the final Proxy XCH₄ and the XCH₄ / XCO₂ component were both validated against the TCCON data. In this section we validate the remaining component of the proxy product from Eq. (1), namely the model XCO₂.

As discussed in Sect. 2, this update to the University of Leicester GOSAT Proxy XCH₄ data uses an ensemble of model XCO₂ data to act as the model XCO₂ component. We utilise the XCO₂ from three state-of-the-art global transport models which all assimilate surface in situ measurements; GEOS-Chem (University of Edinburgh – Feng et al., 2011, v1.50), MACC-II (Chevallier et al., 2010, v14r1) and CarbonTracker (NOAA – Peters et al., 2007, vCT2013B). These model runs have assimilated similar surface measurements but not necessarily from all of the same data sets or the same locations. The models also have different spatial resolutions and different temporal coverage (GEOS-Chem: 2009–2011, 5° × 4°; CarbonTracker: 2009–2012, 3° × 2°; MACC-II: 2009–2012, 3.75° × 1.89°). Where the model does not cover the full GOSAT time period studied here, the data from the previous year are used and adjusted by the NOAA annual growth rate.

The main concern with using modelled XCO₂ data for the proxy method is that the additional uncertainty added to the final proxy data product is difficult to determine. Where the model XCO₂ data are constrained by surface data there can be a high degree of confidence that the model data are close to representing the true value of CO₂; however, it is away from such regions where there is a possibility of adding additional biases into the Proxy XCH₄ data. The TCCON stations are mostly in regions that are also well constrained by surface in situ measurements and hence the model CO₂ data should be well constrained, at least at the surface level, and it is therefore expected to reasonably reproduce the TCCON column data. Figure 4 confirms that this is the case. As the model XCO₂ is used as a component in the proxy retrievals, the models are treated as “pseudo-measurements” and validated in the same way as the satellite data in order to maintain consistency with the satellite validation. The model XCO₂ data sampled at each GOSAT measurement point within ± 2° of each TCCON station are found to agree well with the

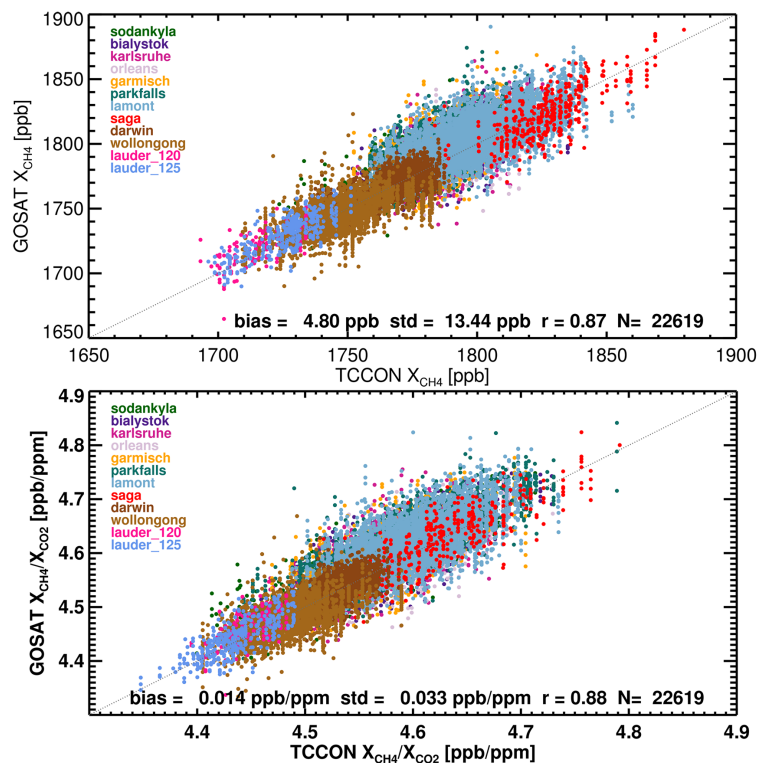


Figure 3. Correlation plot of the Proxy XCH₄ (top) and the XCH₄/XCO₂ ratio (bottom) data against TCCON ground-based FTS data at 11 TCCON sites. The overall bias, standard deviation (single-sounding precision), correlation coefficient, and total number of soundings are provided. Note that the Lauder TCCON station upgraded the instrument from a Bruker 120 to a Bruker 125 in February 2010 and these two data sets are displayed separately.

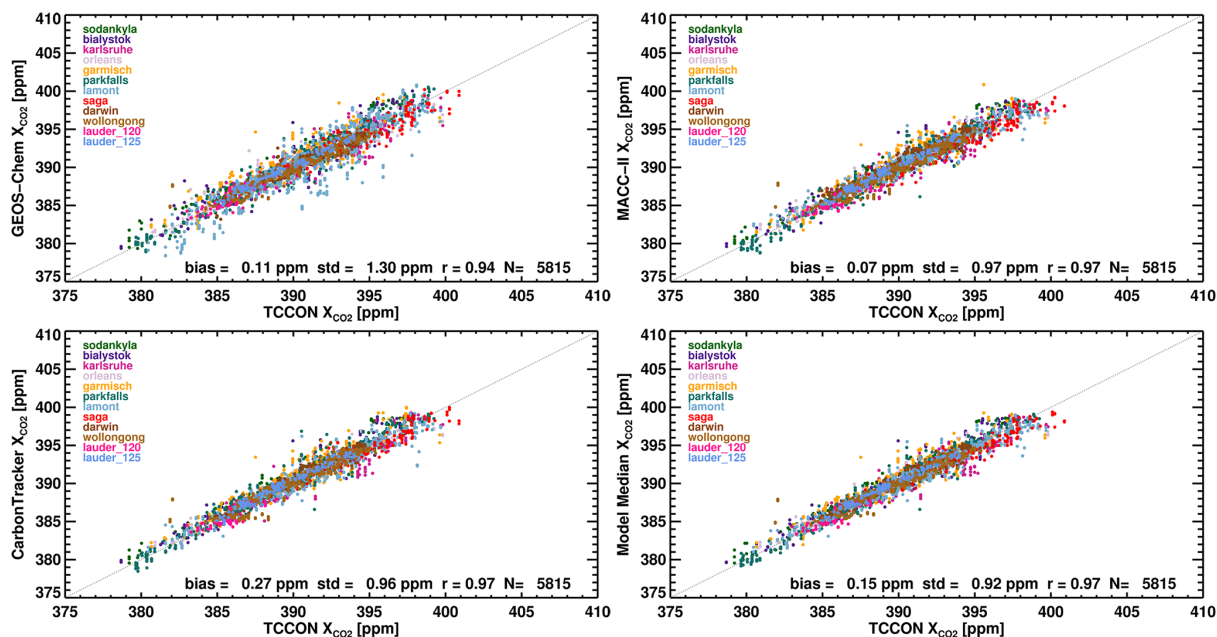


Figure 4. Correlation plot of the model XCO₂ data for GEOS-Chem, MACC-II, CarbonTracker, and the ensemble median against TCCON ground-based FTS data at 11 TCCON sites. The overall bias, standard deviation (single-measurement precision), correlation coefficient, and total number of soundings are provided separately.

TCCON data, with the correlation coefficients ranging from 0.94 (GEOS-Chem) to 0.97 (MACC-II and CarbonTracker). Similarly the precision and bias to TCCON are both found to be small (ranging from 0.97 to 1.3 and 0.07 to 0.27 ppm respectively). The relative accuracies (the standard deviation of the individual site biases) are similar at around 0.5 ppm, with CarbonTracker and GEOS-Chem performing slightly better than MACC-II. Another metric to assess the models is how often they provide the median value of the ensemble. CarbonTracker (41 %) and MACC-II (36 %) tend to provide the median value more often than GEOS-Chem (22 %) but this can vary per site with the contribution from MACC-II as low as 27 % at Darwin (and CarbonTracker at 60 %) and conversely as high as 44 % at Wollongong (with CarbonTracker only 21 %). This provides further indication that no one model can be determined to be the “best”.

For a more detailed analysis of the performance of the different XCO₂ models please see Table A1 in Appendix A. In short, none of the models are found to consistently be superior over the other models. GEOS-Chem typically has the highest scatter against TCCON but also has the smallest bias at 5 out of 12 of the sites. MACC-II has the smallest bias at seven sites but the highest bias at four of the sites. CarbonTracker has the highest bias at seven of the sites but also has the smallest scatter at eight of the sites. Whilst the absolute bias in the calculated median XCO₂ is typically not quite as small as the best of the individual models, the scatter in the median is better than (or the same as) the best scatter from any of the individual models at every site except Lauder_120 (where the time series is the shortest) and even there it is only worse than the best model by less than 0.1 ppm.

The above has demonstrated that it is not a simple decision to determine which model most accurately represents the true atmosphere, even in locations where all of the models have been constrained by (often the same) surface measurements and high-quality validation data are available. In more remote regions where we neither have validation data nor surface measurements to constrain the models, this inconsistency between the models becomes more pronounced. It is this uncertainty in model XCO₂ in regions away from the available validation data that we attempt to address through the use of the XCO₂ model ensemble. Each of the three XCO₂ models are sampled at every GOSAT time and location and convolved with the scene-specific GOSAT averaging kernels. The median value of the three model values is used as the model XCO₂ in calculating the final Proxy XCH₄. However, we also define the uncertainty on this median XCO₂ as the maximum of the absolute differences of each individual model to the median value.

We have already demonstrated that the models all well reproduce the validation data at TCCON sites without any one model identified as being better than the others from our comparisons. Where the models all agree well with each other away from the validation sites, the assumption is that the models are accurately representing the true atmosphere.

Where the models disagree with each other, we do not know which model is correct in the absence of further validation data and in some cases the discrepancy between models can be very large (i.e. > 4 ppm). In such cases where no validation is possible, the best estimate of the uncertainty in the model XCO₂ data is obtained by examining the difference of the model data around the median value. Figure 5 shows global maps of this estimated model uncertainty for each season. There are clear spatial/temporal patterns in the distribution of this model uncertainty. During March–May (boreal “spring”), there is a large uncertainty (> 2 ppm) over India and the African regions typically associated with biomass burning. There is also a moderate level of uncertainty (> 1 ppm) over Europe, South America, and for the latter years over North America and Australia. For the summer months (June–August) it is the Eurasian region, extending from the Ural mountains eastwards through Siberia and northern China, where the model uncertainty is largest at over 2 ppm. This is to be expected as in the Northern Hemisphere it is the period of greatest photosynthetic activity and the model sensitivity to the underlying mechanisms is likely to be largest. During boreal autumn (September–November), the uncertainty in the Northern Hemisphere is vastly reduced again, with India being the major region of uncertainty along with South America and regions of biomass burning in Africa. Winter is similar to autumn, with all three models in very good agreement with each other in the Northern Hemisphere, with only S.E. Asia showing a moderate level of uncertainty. In the Southern Hemisphere, again South America and southern Africa show moderate uncertainty which appears to be linked to emissions from biomass burning.

This section has shown that the estimated uncertainty of the model XCO₂ can vary greatly in time and space. When considering the implication of this uncertainty on flux inversions of the Proxy XCH₄ data, the relative importance of the different uncertainties must be considered. The following section investigates the distribution of the model XCO₂ uncertainty and judges its relative importance against the a posteriori error from the retrieval itself. Finally, both of these uncertainties are assessed against the difference to modelled XCH₄ already constrained by surface observations to determine the utility of the satellite data despite the presence of these uncertainties.

5 Assessing the relative uncertainties

In order to assess the importance of the uncertainty of the model XCO₂, we bin the three model fields into 4° × 5° grid boxes over 8-day time steps and calculate the maximum difference of the three-model ensemble from the median value to use as an estimate of the uncertainty in the model values. We convert this uncertainty in model XCO₂ into an uncertainty in XCH₄ by multiplying each point by its respective retrieved XCH₄ / XCO₂ amount. We also calculate the av-

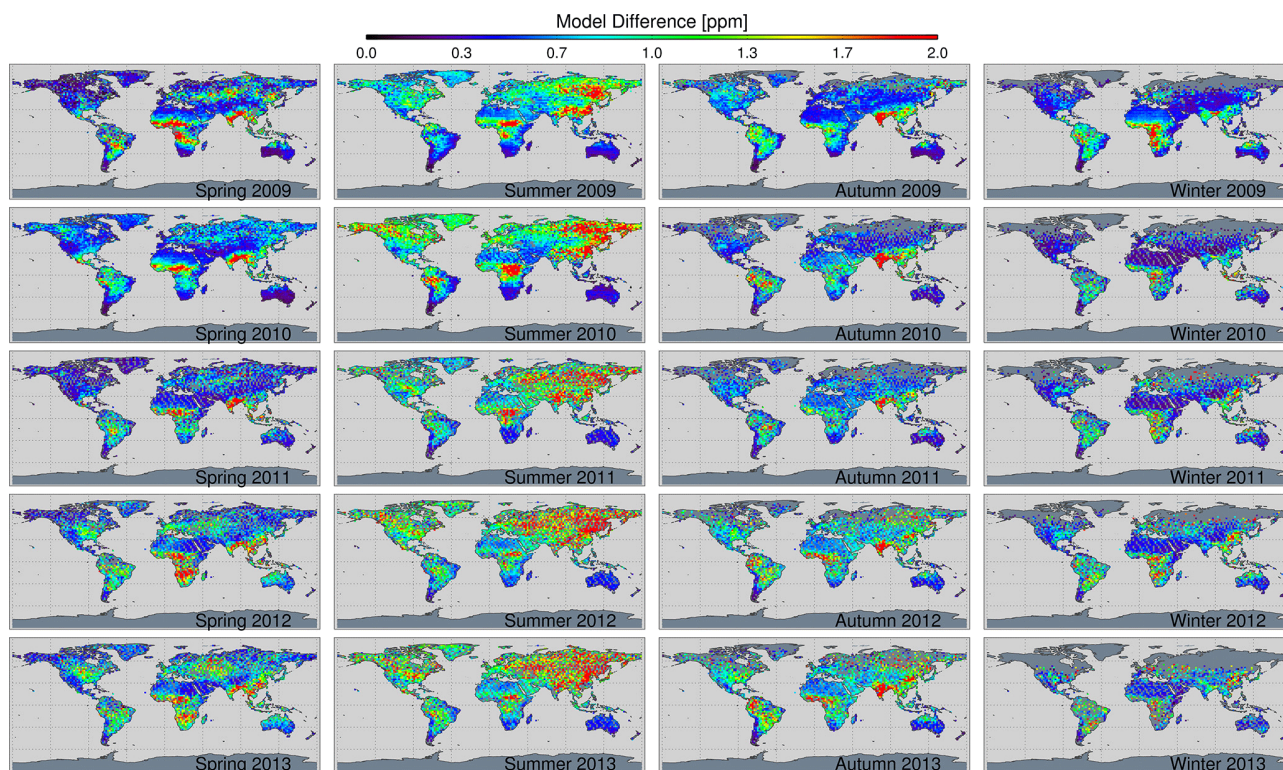


Figure 5. Seasonal maps of the model difference, defined as the maximum absolute difference of the three-model ensemble from the median. All individual soundings have been averaged into $2^\circ \times 2^\circ$ grid boxes over each season. The largest uncertainties occur in regions where the CO₂ variability is expected to be highest and the models are unconstrained by surface measurements.

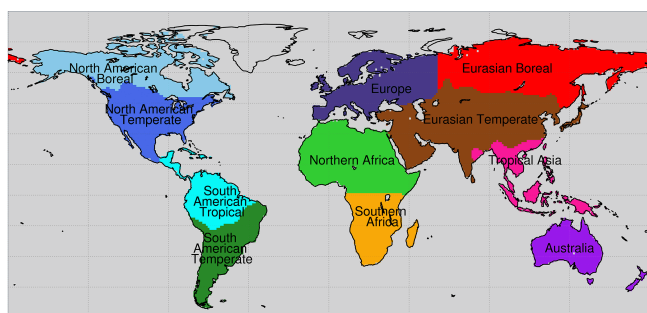


Figure 6. The Transcom regions over which the $4^\circ \times 5^\circ$ gridded data are then averaged in Fig. 7.

erage a posteriori error for the same data. Unlike the more systematic XCO₂ model uncertainty, the a posteriori error should be close to random and hence reduce approximately with the square root of the number of soundings being averaged. If the error does not reduce as much, the model XCO₂ component would then contribute even less to the total, leading to this assumption being a “worst case” scenario for the effect of the model XCO₂ uncertainty. These $4^\circ \times 5^\circ$ grid boxes are then themselves averaged over the Transcom regions (Gurney et al., 2002) as defined in Fig. 6.

In Fig. 7, the red line shows the mean of the Proxy XCH₄ random (a posteriori) error from each $4^\circ \times 5^\circ$ box averaged over each Transcom region with the green line representing the estimated uncertainty related to the model XCO₂. The majority of regions exhibit a similar trend over time. The a posteriori error peaks in the winter months when the SNR of the measurement is at its lowest and is at a minimum during the summer months when the SNR is at a maximum. This seasonal effect is more pronounced at higher latitudes which experience a greater degree of variability of sunlight throughout the year. Conversely, the XCO₂ model uncertainty follows biospheric activity with the uncertainty largest during the summer months when the XCO₂ variability is at a maximum and reduces to a minimum in the winter months when biospheric activity is lower. This leads to the situation where the a posteriori error dominates the model uncertainty in winter months but during summer months the model uncertainty can be comparable to, or even exceed, the a posteriori error. Taking the North America Temperate region as an example, during winter the a posteriori error can reach up to 8 ppb with the error from the model XCO₂ significantly lower with values less than 2 ppb. In contrast, during the summer months, the a posteriori error reduces to around 5 ppb but the error for the model XCO₂ increases to 5 ppb, meaning that both become significant components of the overall uncertainty.

We have shown that the uncertainty related to the XCO₂ model can, particularly in the Northern Hemisphere during summer months, be of comparable magnitude to the a posteriori retrieval error. However, that in itself does not preclude the data from adding useful information to a CH₄ flux inversion.

The MACC-II model XCH₄ (v10-S1NOAA) data have assimilated NOAA surface measurements at background sites and hence are well constrained in the remote atmosphere (Bergamaschi et al., 2013). Here we calculate the difference between the MACC XCH₄ model field and the GOSAT Proxy XCH₄ data for each GOSAT measurement (referred to from here as ΔXCH_4). We then aggregate these differences in the same way as the model XCO₂ uncertainties. Note that the MACC XCH₄ model data are currently only available until the end of 2012. As some inversion systems will perform a simple (e.g. latitudinal) bias correction, the calculated retrieval a posteriori and model XCO₂ uncertainties can potentially be much lower than the ΔXCH_4 value but still not provide information to the inversion. For this reason, it is also important to consider both the mean ($\mu_{\Delta\text{XCH}_4}$) and the standard deviation ($\sigma_{\Delta\text{XCH}_4}$) of the ΔXCH_4 . To determine whether the GOSAT data are capable of providing information to the inversion, we compare the a posteriori and model XCO₂ uncertainties to the $\mu_{\Delta\text{XCH}_4}$ and $\sigma_{\Delta\text{XCH}_4}$ values as shown in Fig. 7, with the seasonal averages for all of these values presented in Table A2.

It should be noted here that the absolute values are not necessarily quantitatively comparable when taking into account how an inversion system will use the two different quantities. The a posteriori error of the retrieved XCH₄ is an indication of the weighting that the inversion will give to an observation over the a priori, with a smaller value indicating that the inversion will “trust” the observation more. The ΔXCH_4 is an indication of how much the inversion needs to adjust the fluxes in order to match the observation. However, if the estimated uncertainties are significantly less than the $\mu_{\Delta\text{XCH}_4}$ and $\sigma_{\Delta\text{XCH}_4}$ values it is expected that the observations should provide value to the inversion. It should also be noted that this bias term ($\mu_{\Delta\text{XCH}_4}$) may also reflect systematic biases in the XCH₄ model due to, for example, errors in the vertical model profile whilst the sigma term ($\sigma_{\Delta\text{XCH}_4}$) may also relate to subgrid-scale variations which are unresolved at the model resolution.

For the North American Boreal region, both the $\mu_{\Delta\text{XCH}_4}$ and $\sigma_{\Delta\text{XCH}_4}$ values are very similar in terms of phase and magnitude to the a posteriori uncertainty with the $\sigma_{\Delta\text{XCH}_4}$ ranging from an average of 9.6 ppb in summer to 14.5 ppb in winter compared to the a posteriori uncertainty that ranges from 6.6 ppb in summer to 10.8 ppb in winter. This suggests that regardless of the contribution to the uncertainty from the XCO₂ model, it would be difficult for the satellite data to inform the inversion any further than the in situ data already do. However, this is not the case for the North American Temperate region where the $\mu_{\Delta\text{XCH}_4}$ (7.4–11.6 ppb) and

$\sigma_{\Delta\text{XCH}_4}$ (8.4–10.0 ppb) are far larger than the total uncertainty (6.0–6.9 ppb) for much of the year. Both South American regions exhibit more complicated behaviour with far less of an apparent seasonality in the $\mu_{\Delta\text{XCH}_4}$. Instead, for most years $\mu_{\Delta\text{XCH}_4}$ is much higher than the uncertainties (which themselves do not exhibit much seasonality in these regions). However, the year 2010 seems to be an anomalous year where the $\mu_{\Delta\text{XCH}_4}$ data are much more in agreement and in this year the difference is of comparable magnitude to the uncertainties with values between 4 and 8 ppb. The $\sigma_{\Delta\text{XCH}_4}$ does exhibit more seasonality than the $\mu_{\Delta\text{XCH}_4}$ and is again considerably higher than the estimated uncertainties (7–20 ppb vs. 5–6 ppb). In combination, this suggests that the GOSAT observations over South America should add considerable information to the inversion.

For Northern Africa, both the a posteriori error and the uncertainty related to the XCO₂ model are small due to the high SNR over the Sahara and the low CO₂ variability respectively (with seasonal average values ranging from 3.3 to 3.6 ppb for the a posteriori error and from 2.8 to 4.6 ppb for the model XCO₂ error). In contrast, the $\mu_{\Delta\text{XCH}_4}$ (7.2–12.3 ppb) and $\sigma_{\Delta\text{XCH}_4}$ (4.9–8.4 ppb) values over this region are relatively large with a high degree of temporal variability, suggesting that the satellite data should add considerable value in constraining the inversion over this region. One complication is that GOSAT operates in a “medium gain” mode over the desert and hence may exhibit different instrumental biases over such regions but, due to the proxy method, any such differences in instrumental biases that relate to light-path modification should be minimised. Southern Africa shows similar behaviour with the total uncertainty being low (seasonal averages of 5.1–7.3 ppb) compared to the much larger $\mu_{\Delta\text{XCH}_4}$ (12.6–19.8 ppb) and $\sigma_{\Delta\text{XCH}_4}$ (5.7–10.7 ppb) values, again indicating that considerable value is present in the satellite data.

The Eurasian Boreal region behaves similarly to the North American Boreal region. The $\mu_{\Delta\text{XCH}_4}$ and $\sigma_{\Delta\text{XCH}_4}$ is of similar phase and magnitude to the retrieval a posteriori error, suggesting little information will be added to any inversion over this region beyond what is available from the in situ measurements. In contrast, the $\mu_{\Delta\text{XCH}_4}$ values over the Eurasian Temperate region show a large variability with the differences in winter months much larger than the total uncertainty (9.9 ppb vs. 5.8 ppb), while in summer months the magnitudes become much more similar (5.5 ppb vs. 7.3 ppb). Interestingly, the $\sigma_{\Delta\text{XCH}_4}$ values appear to be of similar magnitude (5–20 ppb) but directly out of phase with the $\mu_{\Delta\text{XCH}_4}$ values. Even during summer months when the a posteriori (4.3 ppb) and model XCO₂ (5.8 ppb) uncertainties are comparable to the $\mu_{\Delta\text{XCH}_4}$ (5.5 ppb), the high variability in the ΔXCH_4 values, as indicated by $\sigma_{\Delta\text{XCH}_4}$ values of up to 20 ppb (and a summertime mean value of 15.0 ppb), suggests that the observations are capable of providing useful information to the inversion.

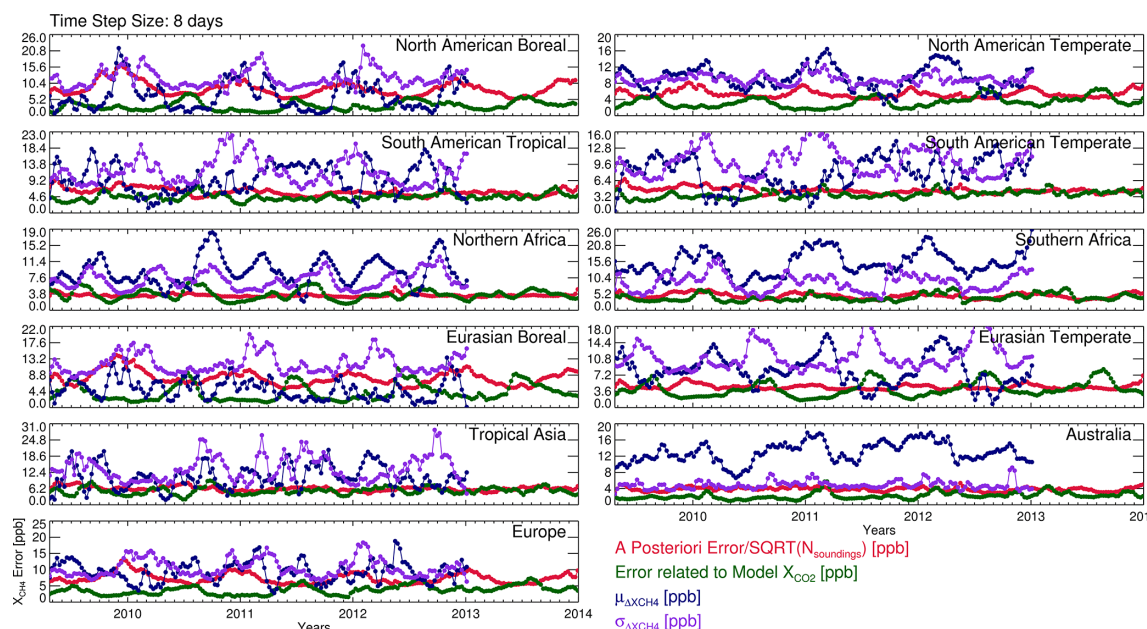


Figure 7. Time series for each Transcom region showing the a posteriori retrieval error (red), the estimated uncertainty from the model XCO₂ (green), and the mean (navy) and standard deviation (purple) of the difference between the GOSAT and MACC-II XCH₄. The a posteriori error is assumed to be a random error and hence reduces with the square root of the number of measurements whilst the XCO₂ model uncertainty is expected to be a systematic error and hence does not reduce.

The Tropical Asian region, which encompasses parts of India, China, and Indonesia, typically has low values for both the a posteriori (5.6–6.9 ppb) and XCO₂ model (4.4–6.0 ppb) uncertainties, with neither exhibiting much seasonal variability. The $\mu_{\Delta XCH_4}$ and $\sigma_{\Delta XCH_4}$ values however are much more variable (8.9–11.7 and 9.0–16.8 ppb) and generally much higher than the uncertainties, suggesting that useful information from the satellite data is present.

The European Transcom region has uncertainties in the satellite data (seasonal averages of 7.6–10.0 ppb) that are of comparable magnitude to the $\mu_{\Delta XCH_4}$ values (8.1–10.7 ppb), especially when considering the combination of the a posteriori and model XCO₂ uncertainties. However, the standard deviation of the $\mu_{\Delta XCH_4}$ values is highly variable (8.7–13.2 ppb) which suggests that there is scope for the observational data to aid in constraining the European XCH₄ fluxes.

Finally, the Australian Transcom region shows very small uncertainties in the satellite data. The uncertainty associated with the model XCO₂ is comparable to the a posteriori error during the Australian spring months but even in those circumstances, the $\mu_{\Delta XCH_4}$ values are far larger (11.4 ppb vs. 4.7 ppb), demonstrating that the satellite data are capable of providing some information to the inversion over Australia, although this may be limited in its ability to provide specific information on Australian sources as the $\sigma_{\Delta XCH_4}$ values over this region are similar to the estimated uncertainties with seasonal averages of 4.5–5.2 ppb compared to the total uncertainty values of 4.1–5.0 ppb.

6 Summary and conclusions

We present details of the update to the University of Leicester GOSAT Proxy XCH₄ v5.0 data set with 5 years of GOSAT data now processed. The data are validated against the latest ground-based TCCON data and found to agree well with on average a small bias of 4.8 ppb ($\sim 0.27\%$), a single-sounding precision of 13.4 ppb ($\sim 0.74\%$), and a relative accuracy of 4.2 ppb. For the first time the XCH₄ / XCO₂ ratio component of the proxy retrieval is validated and also found to agree well with TCCON with a bias of 0.014 ppb ppm⁻¹ ($\sim 0.3\%$) and a single-sounding precision of 0.033 ppb ppm⁻¹ ($\sim 0.72\%$).

A major unknown uncertainty in previous Proxy XCH₄ products was the uncertainty associated with the model XCO₂. In this work we validate three separate state-of-the-art chemistry transport models against the TCCON data and find that although the models can differ greatly (> 4 ppm) away from the TCCON stations, at the validation locations it is difficult to distinguish which model performs better from our comparisons. We therefore decide to use the median of the three models to act as the model XCO₂ in the calculation of the Proxy XCH₄ and use the maximum difference to the median as a measure of the uncertainty in the model XCO₂. This model uncertainty is found to vary greatly in time and space but is typically largest over regions associated with biomass burning such as central Africa and in particular over the Eurasian regions during summer months where large uptake in CO₂ leads to large differences between the models.

In order to assess the relative importance of these uncertainties, we compare this model XCO₂ uncertainty to the a posteriori retrieval error over the different Transcom regions and find typically that where there is seasonality in the uncertainties, it is typically directly out of phase between the two, resulting in the model XCO₂ uncertainty becoming significant during summer months where the a posteriori error is at its lowest. This relates to the fact that more sunlight leads to a reduction in the a posteriori uncertainty (by virtue of providing a greater signal in the SWIR) and at the same time is associated with an increase in photosynthesis and, hence, more potential for differences in the model XCO₂.

We assess the significance of these uncertainties on any flux inversion by comparing the mean and standard deviation of the GOSAT–MACC differences ($\mu_{\Delta\text{XCH}_4}$ and $\sigma_{\Delta\text{XCH}_4}$) to the estimated uncertainties. We find that for the majority of regions the mean and standard deviation of the ΔXCH_4 values are much larger than the estimated uncertainties, even taking into account the uncertainty related to the model

XCO₂. Our findings show that useful information will be provided to the inversions for the majority of regions, with the exceptions being the boreal regions (North American Boreal and Eurasian Boreal) where the uncertainty is of a similar magnitude to the $\mu_{\Delta\text{XCH}_4}$ and $\sigma_{\Delta\text{XCH}_4}$ values. It is important to note that the MACC data are already constrained by NOAA background sites.

One final consideration for users of the Proxy XCH₄ data who are performing atmospheric inversions is that, should they have their own XCO₂ model which they believe is consistent with their XCH₄ model, it may be beneficial to only take the GOSAT XCH₄ / XCO₂ ratio and apply their own model XCO₂ (with appropriate averaging kernels) in order to minimise transport model errors between the different models. Alternatively the XCH₄ / XCO₂ ratio can also be inverted directly as shown in Fraser et al. (2014) and Pandey et al. (2015).

Appendix A: Data sets

The GOSAT Proxy XCH₄ data used in this publication are freely available from http://www.leos.le.ac.uk/GHG/ghg_cci/CRDP/data_2/ESACCI/GHG/GOSAT/CH4_GOS_OCP/5.1/ upon request of a password. An updated version of this data set is now available (version 6.0) covering 2009–2014. Additionally, these data now contain both the raw XCH₄ and XCO₂ values as well as the uncertainty associated with the model XCO₂.

The TCCON XCH₄ and XCO₂ data used in this publication are publicly available from <http://tccon.ornl.gov>. The following data have been used: Sodankylä (Kivi et al., 2014), Bialystok (Deutscher et al., 2014), Karlsruhe (Hase et al., 2014), Orleans (Warneke et al., 2014), Garmisch (Sussmann and Rettinger, 2014), Park Falls (Wennberg et al., 2014a), Lamont (Wennberg et al., 2014b), Saga (Kawakami et al., 2014), Darwin (Griffith et al., 2014a), Wollongong (Griffith et al., 2014b), Lauder120 (Sherlock et al., 2014a), and Lauder125 (Sherlock et al., 2014b).

Table A1. Table showing the comparison statistics between each XCO₂ model (sampled as per the GOSAT measurements) within $\pm 2^\circ$ of each TCCON site against the TCCON validation data. The difference (model–TCCON), the standard deviation of the difference, and the correlation coefficient are all provided as is the total number of measurements for each site, N , and the percentage “share” of the median for each model, %. For each of the three models, GEOS-Chem, MACC-II, and CarbonTracker, the best (bold) and worst (italic) value for each metric is highlighted. For the ensemble median data, all values which are better than the best individual model value are highlighted in bold-italic. The lower panel provides overall statistics across all sites. These include the relative accuracy (the standard deviation of the individual site biases), the overall precision (the standard deviation of the GOSAT–TCCON differences), and the overall share that each model contributes to the median ensemble.

$2^\circ \times 2^\circ$																
Coincident criteria		GEOS-Chem					MACC-II					CarbonTracker				
TCCON Site	N	Diff (ppm)	SD (ppm)	r	%	Diff (ppm)	SD (ppm)	r	%	Diff (ppm)	SD (ppm)	r	%	Diff (ppm)	SD (ppm)	r
Sodankylä	584	1.1	1.1	0.97	20	0.9	0.9	0.98	37	1.2	0.9	0.99	42	1.1	0.8	0.99
Bialystok	1429	0.6	1.5	0.95	25	0.4	1.1	0.97	33	0.6	1.0	0.98	44	0.6	1.0	0.97
Karlsruhe	1569	−0.2	1.4	0.92	22	−0.6	1.1	0.95	33	−0.4	1.1	0.95	45	−0.4	1.1	0.95
Orleans	1650	0.3	1.2	0.95	22	0.3	0.9	0.98	33	0.4	0.9	0.97	46	0.3	0.8	0.98
Garmisch	1527	0.8	1.3	0.93	22	0.6	1.3	0.94	34	0.8	1.2	0.95	43	0.7	1.1	0.95
Park Falls	2434	0.4	1.1	0.97	23	0.1	1.0	0.98	38	0.5	1.0	0.98	40	0.3	0.9	0.98
Lamont	7464	−0.2	1.6	0.92	20	−0.1	0.9	0.98	39	0.0	0.9	0.98	41	−0.1	0.9	0.98
Saga	379	−0.6	1.1	0.93	27	−1.0	0.9	0.95	33	−0.3	0.9	0.95	40	−0.6	0.9	0.96
Darwin	2491	0.0	0.8	0.97	12	0.5	0.7	0.97	27	0.4	0.6	0.98	60	0.3	0.6	0.98
Wollongong	2601	−0.1	0.8	0.96	36	−0.1	0.8	0.96	44	0.2	0.9	0.95	21	0.0	0.8	0.96
Lauder_120	124	−0.1	0.9	0.82	27	−0.3	0.7	0.86	44	−0.2	0.8	0.84	29	−0.2	0.8	0.84
Lauder_125	368	0.3	0.4	0.99	30	0.2	0.3	0.99	40	0.4	0.4	0.99	30	0.3	0.3	0.99
Statistics for all sites																
Relative accuracy (ppm)		0.48				0.53				0.47				0.48		
Overall precision (ppm)		1.3				0.97				0.96				0.92		
Total median share (%)		22				36				41						

Table A2. Table showing the seasonal averages of the data plotted in Fig. 7 for each of the Transcom regions. The retrieved a posteriori error, the uncertainty related to the model XCO₂, their combined total, and the mean and standard deviation of the GOSAT–MACC difference are all provided for each season and for each Transcom region.

Region	Season	A posteriori (ppb)	Model (ppb)	Total (ppb)	$\mu_{\Delta\text{XCH}_4}$ (ppb)	$\sigma_{\Delta\text{XCH}_4}$ (ppb)	Region	Season	A posteriori (ppb)	Model (ppb)	Total (ppb)	$\mu_{\Delta\text{XCH}_4}$ (ppb)	$\sigma_{\Delta\text{XCH}_4}$ (ppb)
North America boreal	spring	7.4	2.2	7.8	6.3	13.0	Eurasian boreal	spring	7.1	3.2	8.0	4.5	13.1
	summer	6.6	4.9	8.3	2.9	9.6		summer	6.3	7.0	9.5	4.4	9.9
	autumn	10.0	2.8	10.5	6.2	11.4		autumn	9.0	3.1	9.6	5.4	11.1
	winter	10.8	2.3	11.1	11.5	14.5		winter	9.9	2.5	10.3	5.7	13.9
North America temperate	spring	5.1	3.0	6.0	11.2	9.2	Eurasian temperate	spring	4.4	3.5	5.7	11.7	9.7
	summer	4.6	4.7	6.6	7.4	8.6		summer	4.3	5.8	7.3	5.5	15.0
	autumn	5.3	3.1	6.3	7.7	8.4		autumn	4.4	4.1	6.2	6.5	10.2
	winter	6.5	2.4	6.9	11.6	10.0		winter	5.2	2.6	5.8	9.9	9.6
South America tropical	spring	6.4	4.3	7.8	8.6	11.2	Tropical Asia	spring	6.0	6.0	8.5	8.9	12.2
	summer	5.3	4.3	6.9	10.8	8.6		summer	6.9	5.0	8.7	10.9	16.1
	autumn	5.8	5.2	7.9	10.7	12.6		autumn	6.2	6.0	8.7	11.7	16.8
	winter	6.5	4.1	7.8	7.4	15.5		winter	5.6	4.4	7.2	10.9	9.0
South America temperate	spring	4.7	3.7	6.1	6.0	10.7	Australia	spring	3.9	2.5	4.7	11.4	4.7
	summer	4.3	3.4	5.5	9.1	7.4		summer	3.7	1.8	4.1	11.7	4.5
	autumn	4.2	3.7	5.7	9.1	9.7		autumn	3.7	2.1	4.3	13.9	5.2
	winter	4.7	3.8	6.1	7.2	13.5		winter	4.2	2.5	5.0	15.0	5.2
Northern Africa	spring	3.6	3.6	5.2	8.8	7.3	Europe	spring	6.7	3.5	7.6	9.2	12.1
	summer	3.6	4.6	5.9	7.2	7.4		summer	5.9	5.3	8.0	10.7	8.7
	autumn	3.4	3.5	4.9	12.3	8.4		autumn	7.5	3.0	8.2	8.1	8.9
	winter	3.3	2.8	4.3	8.0	4.9		winter	9.4	3.1	10.0	9.3	13.2
Southern Africa	spring	4.7	4.6	6.7	15.5	9.9		spring					
	summer	3.7	3.4	5.1	12.6	5.7		summer					
	autumn	4.8	3.7	6.1	13.5	9.8		autumn					
	winter	5.4	4.8	7.3	19.8	10.7		winter					

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