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Global high-resolution simulations of CO_2 and CH_4 using a NIES transport model to produce a priori concentrations for use in satellite data retrievals

T. Saeki[1], **R. Saito**[2], **D. Belikov**[1], **and S. Maksyutov**[1]

[1]Center for Global Environmental Research, National Institute for Environmental Studies, Tsukuba, 305-8506, Japan
[2]Research Institute for Global Change, JAMSTEC, Yokohama, 236-0001, Japan

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Correspondence to: T. Saeki (saeki.tazu@nies.go.jp)

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Abstract

The Greenhouse gases Observing SATellite (GOSAT) measures column-averaged dry air mole fractions of carbon dioxide and methane (XCO_2 and XCH_4, respectively). Since the launch of GOSAT, model-simulated three-dimensional concentrations from a National Institute for Environmental Studies offline tracer Transport Model (NIES TM) have been used as a priori concentration data for retrieving XCO_2 and XCH_4 from GOSAT short-wavelength infrared spectra at NIES. Though a priori concentrations for retrievals are optional, more reliable concentrations are desirable. In this paper we describe the newly developed NIES TM that has been adapted to provide global and near real-time concentrations of CO_2 and CH_4 using a high-resolution meteorological dataset, the Grid Point Value (GPV) prepared by the Japan Meteorological Agency. The spatial resolution of the NIES TM is set to $0.5^\circ \times 0.5^\circ$ in the horizontal in order to utilize GPV data, which have a resolution of $0.5^\circ \times 0.5^\circ$, 21 pressure levels, and a time interval of 3 h. GPV data are provided to the GOSAT processing system with a delay of several hours, and the near real-time model simulation produces a priori concentrations driven by diurnally varying meteorology. A priori variance–covariance matrices of CO_2 and CH_4 are also derived from the simulation outputs and observation-based reference data for each month of the year at a resolution of $0.5^\circ \times 0.5^\circ$ and 21 pressure levels. Model performance is assessed by comparing simulation results with the GLOBALVIEW dataset and other observational data. The overall root-mean-square differences between model predictions and GLOBALVIEW analysis are estimated to be 2.28 ppm and 12.68 ppb for CO_2 and CH_4, respectively, and the seasonal correlation coefficients are 0.86 for CO_2 and 0.61 for CH_4. The model showed good performance particularly at oceanic and free tropospheric sites. The model also performs well in reproducing both the observed synoptic variations at some sites, and stratospheric profiles over Japan. These results give us confidence that the performance of our GPV-forced high-resolution NIES TM is adequate for use in satellite retrievals.

GMDD

5, 2215–2258, 2012

1 Introduction

Global atmospheric transport models provide an effective means of quantifying the global cycle of long-lived atmospheric trace gases such as carbon dioxide (CO_2) and methane (CH_4). There is an increasing demand for high-resolution models that simulate global tracer transport over synoptic and sub-daily timescales to reproduce observed variations more accurately. For example, the Atmospheric Tracer Transport Model Intercomparison Project (TransCom) has initiated simulations of hourly and synoptic CO_2 concentration (Law et al., 2008; Patra et al., 2008), which will complement and make use of state-of-the-art measurements of greenhouse gases. In this TransCom continuous experiments, 25 transport models participated, and two of them ran at horizontal resolutions of $0.5^\circ \times 0.5^\circ$ and the others ran at $1^\circ \times 1^\circ$ to $3.8^\circ \times 5.0^\circ$ resolutions. Patra et al. (2008) concluded that increasing model horizontal resolution clearly improved the synoptic-scale variations in simulated CO_2. Maksyutov et al. (2008) compared model CO_2 results at horizontal resolutions of 2.0°, 1.0°, 0.5°, and 0.25° with continuous observations at a tower site in Japan and showed that increasing the model's horizontal resolution greatly improved the match with observations. But most of recent model simulations of these greenhouse gases still have carried out at horizontal resolutions of $1^\circ \times 1^\circ \sim 3.75^\circ \times 2.5^\circ$ (e.g. Allen et al., 2011; Saito et al., 2011; Patra et al., 2011).

Another demand for the high-resolution models stems from a new approach in which model-predicted CO_2 and CH_4 concentrations are used to give a priori concentrations for satellite spectroscopic data retrieval algorithms (e.g. Saitoh et al., 2009; Yoshida et al., 2011; O'Dell et al., 2012). Satellite observations cover most of the globe in several days to a few weeks, and retrievals based on satellite spectra require a priori concentrations of targeted gases. Global transport models can provide simulated a priori concentration profiles of those greenhouse gases in order to obtain optimal retrieval solutions and to physically interpret satellite-derived data. Instantaneous fields of view of satellite instruments are of the order of 10–100 km; e.g. $30 \times 120\,km^2$ for

GMDD
5, 2215–2258, 2012

the SCanning Imaging Absorption spectroMeter for Atmospheric CHartographY (SCIAMACHY) (Buchwitz et al., 2005), and a nadir circular footprint of about 10.5 km diameter for the Greenhouse gases Observing SATellite (GOSAT) (Yoshida et al., 2011). For GOSAT, the retrieval precisions are estimated to be smaller than 3.5 ppm and 15 ppb for column-averaged dry air mole fractions of both carbon dioxide and methane (XCO_2 and XCH_4), respectively (Yoshida et al., 2011). Furthermore, future satellites that observe greenhouse gases target higher precision with less bias. For example, the Orbiting Carbon Observatory-2 (OCO-2) is designed to retrieve XCO_2 theoretically with 1–2 ppm (0.3–0.5 %) precision for single-soundings with a small field of view with an area of 3 km^2 in nadir (Boesch et al., 2011). Therefore, much effort is currently being devoted to the development of global high-resolution transport models with less model error that meet the demands from satellite observations of greenhouse gases.

GOSAT is the first satellite to measure global distributions of XCO_2 and XCH_4 (Kuze et al., 2009; Yokota et al., 2009; Yoshida et al., 2011). At the National Institute for Environmental Studies (NIES), XCO_2 and XCH_4 are retrieved from the short-wavelength infrared (SWIR) spectra obtained by the Thermal And Near infrared Sensor for carbon Observation-Fourier Transform Spectrometer (TANSO-FTS) on board GOSAT. An optimal estimation method is used to retrieve XCO_2 and XCH_4 which minimizes the cost function

$$J(\boldsymbol{x}) = [\boldsymbol{y} - F(\boldsymbol{x})]^T \mathbf{S}_\varepsilon^{-1} [\boldsymbol{y} - F(\boldsymbol{x})] + [\boldsymbol{x} - \boldsymbol{x}_\mathrm{a}]^T \mathbf{S}_\mathrm{a}^{-1} [\boldsymbol{x} - \boldsymbol{x}_\mathrm{a}] \quad (1)$$

where $\boldsymbol{x}$ is the state vector to be retrieved, $\boldsymbol{y}$ is the vector containing the observed spectrum, $F(\boldsymbol{x})$ is the forward model that relates the state vector to the observed spectrum, $\mathbf{S}_\varepsilon$ is the error covariance matrix of the observed spectrum, $\boldsymbol{x}_\mathrm{a}$ is the a priori state of $\boldsymbol{x}$, and $\mathbf{S}_\mathrm{a}$ is the a priori variance–covariance matrix (VCM) (Yokota et al., 2009; Yoshida et al., 2011). A priori state $\boldsymbol{x}_\mathrm{a}$ includes a priori concentration profiles of CO_2 and CH_4. The retrieved XCO_2 and XCH_4 at NIES are available after April 2009 at GOSAT User Interface Gateway (GUIG; http://data.gosat.nies.go.jp/).

GMDD
5, 2215–2258, 2012

As a priori concentrations for NIES XCO_2 and XCH_4 retrievals, we use simulated data from a NIES atmospheric tracer transport model (hereafter NIES TM) (Japan Aerospace Exploration Agency, National Institute for Environmental Studies, and Ministry of the Environment, 2011 (hereafter JAXA/NIES/MoE, 2011); Yoshida et al., 2011). The a priori VCMs, $\mathbf{S}_a$, for CO_2 and CH_4 are derived from simulated NIES TM data and some reference data (Eguchi et al., 2010). A priori concentrations for satellite data retrievals are optional, and various a priori concentrations have been used to retrieve XCO_2 and XCH_4 from GOSAT SWIR spectra; e.g, constant a priori concentrations (Oshchepkov et al., 2011), monthly zonal means in 10° latitude bands for land and ocean from a forward model run (O'Dell et al., 2012), or model-simulated concentrations for the year 2007, 2008 (Butz et al., 2011) or 2009 that was extrapolated to 2010 (Schepers et al., 2012). However, as seen in Eq. (1), when the diagonal elements of $\mathbf{S}_a$ have small values, the a priori profiles largely constrain the retrieved results (e.g. Saitoh et al., 2009; Yoshida et al., 2011). We thus aim to reduce errors in a priori CO_2 and CH_4 concentrations for use in the NIES retrieval algorithm by using NIES TM at the relatively high horizontal resolution of 0.5° × 0.5°.

Another feature of the NIES a priori concentrations is that they are created by NIES TM driven by a near real-time high-resolution meteorological dataset at a horizontal resolution of 0.5° × 0.5° and 21 pressure levels. Accordingly, real-time SWIR retrieval processing is made available to create XCO_2 and XCH_4 products at every observed day by using a priori concentrations that reflect daily meteorological variations at the observed days. Use of high-resolution meteorological data would be expected to allow smaller-scale phenomena to be represented in the model.

In this paper, we describe a newly developed high-resolution NIES TM designed to provide near real-time global three-dimensional concentration fields that reflect daily meteorological conditions for satellite retrieval algorithms, which is currently implemented in the GOSAT Level 2 retrieval system to derive XCO_2 and XCH_4 at NIES (Sect. 2). In Sect. 3, the simulated CO_2 and CH_4 concentrations are compared with an analysis of GLOBALVIEW-CO_2 and CH_4 data products (hereafter GV-CO_2, GV-CH_4)

GMDD
5, 2215–2258, 2012

and other observations to evaluate the model performance and investigate potential biases arising from the model simulation. Our conclusions follow in Sect. 4.

2 NIES transport model and numerical experiments

A NIES off-line global transport model (Maksyutov et al., 2008) has been used to simulate seasonal and spatial distributions of long-lived atmospheric constituents in the lower and mid-troposphere. The advection scheme of the model was semi-Lagrangian, and a mass fixer was adopted. The vertical mixing in the model was represented by cumulus convection and turbulent diffusion with explicitly parameterized Planetary Boundary Layer (PBL) physical processes. The earlier version of the NIES transport model (denoted NIES-99) was developed to simulate the seasonal cycles of long-lived tracer species at a relatively coarse horizontal resolution (2.5°–5.0° longitude–latitude), and to perform source–sink inversions of atmospheric CO_2 (e.g. Gurney et al., 2002, 2004; Patra et al., 2002, 2003, 2005a). Improvements to NIES-99 led to a recent development of the model (NIES-05), which has a higher horizontal resolution (tested on $2^\circ \times 2^\circ$ to $0.25^\circ \times 0.25^\circ$ in Maksyutov et al., 2008). NIES-05 was driven by the ECMWF 3-hourly PBL height data and the vertical resolution was enhanced to 47 levels for better resolution of the mixing processes in the boundary layer. For more details of NIES TM, see Maksyutov et al. (2008). NIES-05 was able to simulate observed diurnal-synoptic scale variability of tracers of interest, and participated in the TransCom hourly CO_2 experiment (Law et al., 2008; Patra et al., 2008). The original version of NIES-05 used meteorology datasets from NCEP final analyses (http://dss.ucar.edu/datasets/ds083.2/).

We have recently upgraded NIES-05 to utilize a high-resolution meteorological dataset, the Japan Meteorological Agency (JMA) Grid Point Values (GPV) product (Belikov et al., 2011). GPV/JMA data is created from JMA Global Spectral Model, which is operated for short- and medium-range forecasts covering the entire globe with TL959 resolution and 60 vertical levels from the surface to 0.1 hPa and assimilated by 4D-Var within a framework of JMA's numerical weather prediction system

GMDD
5, 2215–2258, 2012

http://www.jma.go.jp/jma/en/Activities/nwp.html; JMA, 2007). Original GPV data is provided by 0.5° × 0.5° horizontal resolution from the surface to 100 hPa and 1.0° × 1.0° above 100 hPa to 10 hPa (http://www.jmbsc.or.jp/hp/online/f-online0a.html). For use in GOSAT project, GPV data is specially extended at resolution of 0.5° × 0.5° up to 10 hPa. Since November 2007, the extended GPV has been supplied for the GOSAT Data Handling Facility (DHF; JAXA/NIES/MoE, 2011) at NIES in near-real time (delay of several hours). The spatial resolution of the GOSAT version of the GPV dataset is 0.5° × 0.5° on 21 pressure levels (from 1000 to 10 hPa) and the time resolution is 3 h. We used both of GPV objective analysis and forecasts. The GPV data and the ECMWF 3-hourly PBL height data in GRIB2 format are automatically converted to direct access binary for NIES TM input on DHF everyday. Data size is about 1.1 GB per one model day.

We employed the NIES-05 model driven by GPV data to simulate atmospheric CO_2 and CH_4 concentrations for use as a priori concentrations for GOSAT SWIR Level 2 processing at NIES. The horizontal resolution was 0.5° with 47 sigma vertical levels from the surface to 0.01 sigma, and simulated CO_2 and CH_4 concentrations were calculated for every observed day. Flux climatologies were prepared because no real-time fluxes were available. The climatological CO_2 flux dataset was that prepared for the TransCom model inter-comparison studies (Gurney et al., 2004), which consisted of four components: (1) annual constant fossil fuel emissions with a spatial resolution of 0.5° × 0.5° (Brenkert, 1998); (2) monthly terrestrial biosphere flux obtained using the Carnegie-Ames-Stanford-Approach (CASA) model (Randerson et al., 1997); (3) monthly varying ocean flux (Takahashi et al., 2002); and (4) monthly flux corrections obtained using the cyclostationary inversion approach (Gurney et al., 2004) with NIES-99. For CH_4, we used the monthly varying flux for 2000, as obtained from Patra et al. (2009). This CH_4 flux is based on the Emission Database for Global Atmospheric Research (EDGAR) version 32FT2000 (Olivier and Berdowski, 2001) for anthropogenic CH_4, and on GISS emissions (Fung et al., 1991) for natural CH_4. The chemical destruction of CH_4 by OH radicals was calculated based on climatological monthly mean OH radical concentrations (Spivakovsky et al., 2000) and a temperature-dependent rate constant. All these

GMDD
5, 2215–2258, 2012

flux datasets were prepared at a spatial resolution of 1° × 1°, except for the CO_2 fossil fuel emission.

Stratospheric CO_2 and CH_4 variations are poorly understood due to a lack of precise observations over the globe. In addition, it is difficult to reproduce accurately transport and chemical processes in the stratosphere in NIES TM. This is a common problem in many transport models, where the model age of air in the stratosphere tends to be younger than that observed (Saito et al., 2011, and references therein). To reduce model biases in the stratosphere, the model stratospheric concentrations above the diagnosed tropopause are nudged towards the zonal-mean climatological concentrations based on observations. The tropopause in the model is determined from the gradient of potential temperature versus geopotential height at every model grid box at every time step. Zonal-mean climatologies for CO_2 and CH_4 were prepared at 2.5° resolution for every month. CO_2 monthly climatological concentrations in the stratosphere were constructed using the Gap-filled Ensemble Climatology Mean (GECM; Saito et al., 2011). GECM is a three-dimensional daily CO_2 concentration generated by combining information from in situ measurements and multi-model means, carried out in the framework of the TransCom satellite experiment in which six models participated. The mean age of air in the GECM stratosphere has been corrected using in situ profiles of SF_6. The latest version of GECM was used here with further corrections in the stratosphere; i.e., the vertical gradient in GECM CO_2 concentration at northern mid-latitudes from 30 hPa to 10 hPa was corrected to match CO_2 observations (Aoki et al., 2003; Engel et al., 2009). The climatological stratospheric CO_2 values were prepared from 2007 to 2015 with interannual variations for CO_2. For CH_4, monthly climatological values in the stratosphere were derived from satellite measurements by the Halogen Occultation Experiment (HALOE) (Russell et al., 1993), averaged over the period 1994–2005 to exclude the period of the Pinatubo eruption in 1991. HALOE provides a long time series of data, from October 1991 to November 2005 (Russell et al., 1993). Park et al. (1996) validated HALOE CH_4 data against the following correlative data from space-borne infrared spectroscopy: the Atmospheric Trace Molecule

GMDD
5, 2215–2258, 2012

Spectroscopy Experiment; the MARK IV balloon-borne Fourier transform spectrometer; rocket-based cryogenic whole air sampler, balloon-borne laser in situ sensor; and the Kernforschungsanlage cryogenic whole air sampler. They concluded that the total error for the 0.3 to 50 hPa region was less than 15 % and the precision was better than 7 %. Thus, we consider the HALOE CH_4 data set would be suitable for making climatological stratospheric CH_4 distributions. No clear trend had been seen in HALOE time series; thus, we repeatedly used the same monthly concentrations every year. These stratospheric corrections were about −0.5 ppm and −50 ppb for XCO_2 and XCH_4, respectively, in the northern mid latitudes.

The model was initialized with zonal-mean concentration fields on 1 January 2007 derived from GECM for CO_2 and GV-CH_4 (GLOBALVIEW-CH_4, 2009) for CH_4. As GPV data are only available after November 2007, the model for 2007 was forced with GPV 2008 data. CO_2 fossil fuel emission and CH_4 emissions were scaled to reproduce the 2007–2008 CH_4 trend at the South Pole (SPO) observed by the National Oceanic and Atmospheric Administration/Earth System Research Laboratory (NOAA/ESRL) in the WMO World Data Centre for Greenhouse Gases (WDCGG) database (http://ds.data.jma.go.jp/gmd/wdcgg/wdcgg.html). After 2 yr of spin-up with 2008 meteorological data, simulated concentrations at the model's southernmost grid box at 1 February 2009 were readjusted by offsets to the observed NOAA/ESRL CO_2 and CH_4 values at the South Pole from the WDCGG dataset. The model was then handed over for operational processing on the GOSAT DHF after February 2009. The JMA provides the GOSAT DHF with GPV data within a day and the near real-time model simulation has been performed for every observation day. The simulated CO_2 and CH_4 concentrations at 21 pressure levels have been provided as a priori concentrations to the GOSAT Level 2 data processing to retrieve XCO_2 and XCH_4 from SWIR spectra at NIES. Model integration time for one day for two tracers (CO_2 and CH_4) is about 530 s by wall-clock time on a single CPU in the NIES Supercomputer System (NEC SX-8R/128M16).

To produce a priori VCMs of CO_2 and CH_4 in Eq. (1), we used the simulated concentrations for the year 2008, GV-CO_2, GV-CH_4 and observational data. The details of

GMDD
5, 2215–2258, 2012

the procedure to produce VCMs were described in Eguchi et al. (2010). In brief, the VCM was defined as the sum of the bias and noise components, where the bias was obtained from the difference in seasonal cycle between simulated results of NIES TM and GV data, and the noise components consist of synoptic and interannual variations. The synoptic term was calculated from NIES TM results and interannual variations are derived from GV datasets over a few decades. The VCMs of CO_2 and CH_4 were prepared at each grid box of $0.5^\circ \times 0.5^\circ$, on 21 pressure levels over the globe for each month. In this study we modified the stratospheric part of the previous version of VCMs to obtain more realistic values; i.e., stratospheric CO_2 and CH_4 seasonal biases were set to the standard deviations of balloon-borne observations over Japan (Aoki et al., 2003; Nakazawa and Aoki, unpublished data, 2009), and HALOE, respectively. The resulting variances (diagonal elements) in the stratosphere were about 2–3 ppm^2 for CO_2 and about 500–10,000 ppb^2 for CH_4. Figure 1 shows VCMs for CO_2 and CH_4 over Sanriku, Japan (141.8° E, 39.2° E) in August. The values of diagonal elements are large near the surface and decrease with height. They show positive correlations, particularly near the surface. The off-diagonal elements are positive around the diagonal elements, while there are negative correlations between some pressure levels. Correlations in the stratosphere (200–10 hPa) are nearly the same zero, indicates a weak correlation between tropospheric and stratospheric time series.

3 Results and discussion

In the following subsections, we evaluate the model performance against the analyzed data from GV-CO_2 and GV-CH_4, and other observations. Annual mean and monthly biases of the simulated CO_2 and CH_4 are examined in Sects. 3.1 and 3.2, respectively. Synoptic variations that include annual trends are validated against some sites in the WDCGG dataset in Sect. 3.3. Balloon-borne observations in the stratosphere are used to validate the simulated vertical profiles in Sect. 3.4. Finally, the simulated surface CO_2 and CH_4 concentrations and their column-averaged dry air mole fractions are

GMDD
5, 2215–2258, 2012

presented in Sect. 3.5. For all the comparisons, the nearest horizontal and vertical model grid box to the observation location is selected.

3.1 Comparison with observations: annual-mean biases

Annual means of simulated CO_2 and CH_4 concentrations are compared with those from GV analysis (GLOBALVIEW-CO_2, 2009; GLOBALVIEW-CH_4, 2009) for the year 2008 at 260 (CO_2) and 206 (CH_4) GV sites (Fig. 2) because no GV-CH_4 dataset is opened for the years 2009 and 2010. Annual mean biases of the simulated concentrations against GV analysis are shown in Table 1 and Fig. 3. Simulated results averaged over 13:00–16:00 LT were used for the comparison. In general, the annual mean model biases (difference between simulations and GV analysis) are found to be less than ~ 1 ppm for CO_2 and ~ 10 ppb for CH_4 at oceanic and Southern Hemisphere sites. The model also reproduces the GV pole-to-pole gradients very well for both CO_2 and CH_4.

For CO_2, the average annual mean bias and its standard deviation are −0.01 ± 1.57 ppm and RMSD of 2.28 ppm (Table 1), which shows good agreement with less than 1 % between the model prediction and the GV data at most of the sites. Some inland or near-continental sites show significant positive biases (e.g. LJO, La Jolla, California, 32.90° N, 117.30° W, 10 m a.s.l; HUN, Hegyhatsal, Hungary, 46.95° N, 16.65° E, 248 m a.s.l.) or negative biases (BSC, Black Sea, Constanta, Romania, 44.17° N, 28.69° E, 3 m a.s.l). LJO is located on the west coast of California and the GV dataset shows a clear seasonal cycle with small synoptic variations, which only involve differing oceanic air masses. However a large point source with CO_2 15 ppm higher than the surroundings appears on the model grid close to LJO, and it is clear that in the model LJO is affected by plumes from this point source. This may explain the large model–observation mismatch at LJO. This might be caused by the high variability observed near source regions, which is not expected to be captured by global scale transport models. Even with the NIES TM 0.5° grids, such sub-grid scale dynamics could not be represented in the model. The BSC site is located in a coastal region of the Black Sea. Peérez-Landa et al. (2007a,b) studied the effect of regional and local meteorological

GMDD
5, 2215–2258, 2012

conditions on CO_2 transport in the coastal area of Valencia, Spain, and concluded that coastal circulation and strong local flux gradients introduced large biases against observations in a model. When a digital filter is applied to the BSC record in the GV data, large irregular seasonal variations of about −5 to +5 ppm are found, against regular seasonal peak-to-peak amplitudes of 14.8 ppm. BSC appears to be affected by local-scale circulation and/or local fluxes, and this may explain the failure of the model to reproduce BSC variations, as is the case also with HUN. The HUN site, a tall tower site, has the largest model–observation mismatch in the TransCom 3 seasonal experiment (Gurney et al., 2004). The model tends to overestimate GV data at most of the tower sites (e.g. ITN051, Grifton, North Carolina, United States, 35.35° N, 77.48° W, 9 + 51 m a.s.l.), mainly because of a failure to reproduce the large seasonal amplitudes and interannual variations, due to their location in areas affected by biogenic CO_2, such as forest or inland plain.

The simulated CH_4 mostly agree well with the GV data, with an average annual mean bias and standard deviation of 0.88 ± 14.16 ppb and RMSD of 12.68 ppb (Table 1). Significant positive or negative biases are found in Southeast Asia (SCS, South China Sea, shipboard observation), Eurasia (TAP, Tae-ahn Peninsula, Republic of Korea, 36.73° N, 126.13° E, 20 m a.s.l; KZD, Sary Taukum, Kazakhstan, 44.06° N, 76.82° E, 601 m a.s.l.), and North America (BNE010, Beaver Crossing, Nebraska, airborne observation, 40.80° N, 97.18° W, 0–2000 m; LEF030, Park Falls, Wisconsin, United States, tower observation, 45.95° N, 90.27° W, 472 + 30 m a.s.l.) with maximum 76 ppb and minimum −49 ppb. TAP, on the western edge of Korea, is influenced by seasonally varying wind direction, and local and remote CH_4 sources such as wetland (Dlugokencky et al., 1993), and this causes large interannual and seasonal variations, which the model finds difficult to reproduce. The SCS shipboard track is located downwind of Eastern and Southeastern Asia and is affected by the complex wind circulation of the Intertropical Convergence Zone (ITCZ). SCSN03 (3° N) and SCSN06 (6° N) in particular are points surrounded by Indochina, the Malay Peninsula, and Indonesian islands where CH_4 sources occur, such as wetlands and biomass burning. Located in the

GMDD
5, 2215–2258, 2012

American Prairie, CH_4 at BNE010 exhibits scattered and large interannual variations and seasonal cycles, which is also difficult to reproduce by the model.

3.2 Comparison with observations: monthly biases

To assess the ability of the model to reproduce seasonal variations, the simulated monthly mean concentrations of CO_2 and CH_4 for the year 2008 were compared with the analyzed seasonal amplitudes at GV sites (Fig. 4). Analyzed seasonal amplitudes at GV sites were taken from "seas" files stored in the dataset. The simulated results were detrended and monthly means of 13:00–16:00 LT concentrations used for the comparison.

The characteristics of the observed seasonal variability at oceanic sites and free troposphere sites (above 3000 m altitude) are generally reproduced fairly well by the model for both CO_2 and CH_4, but relatively large biases are found at tower sites and some of the land sites. Standard deviations over all GV sites are 1.58 ppm for CO_2 and 14.7 ppb for CH_4. Large seasonal biases over 10 ppm in CO_2 are seen at tower sites in mid-northern latitudes such as ITN (Grifton, North Carolina, United States, 35.35° N, 77.38° W, 9 m a.s.l.) and AMT (Argyle, Maine, United States, 45.03° N, 68.58° W, 50 m a.s.l.); both sites provide continuous measurements and are located near forested areas on the eastern coast of the United States. Seasonal biases at ITN and AMT show that the model seasonal amplitudes are smaller than those in the GV analysis; i.e., the model overestimates the observed summer minima in July and August and underestimates winter maxima. At continental sites with quasi-continuous measurements such as the towers, both the regional-local transport and daily flux variability, including temporal resolution of the biospheric fluxes, are found to be important in simulating such high-frequency CO_2 behavior (Geels et al., 2004, 2007; Patra et al., 2008; Wang et al., 2007). Patra et al. (2008) also found that at lower levels some models overestimated the magnitudes of synoptic variations at high-frequency observational sites at tall towers such as LEF (Park Falls, Wisconsin, United States, 45.95° N, 90.27° W, 472 + 11 ~ 396 m a.s.l.). These small-scale phenomena

GMDD
5, 2215–2258, 2012

may influence model–observation mismatches at tower sites even though the comparisons are on a monthly basis. Except for such tower sites and inland sites, the model succeeds in capturing the GV seasonal variations with biases smaller than 5 ppm at most sites and for most months.

For CH_4, the monthly model biases fall within some 20 ppb at oceanic and free troposphere sites (mostly airborne observational points). In particular, there appear to be no significant monthly model biases in the Southern Hemisphere. However, at continental sites such as TAP (Korea), HUN (Hungary) and ITN (North America), the agreement was poorer, mainly due to the influences of strong sources located near the sites and extreme climate conditions such as a strong inversion layer in a cold winter. Over near-equatorial regions, large biases over 50 ppb were also found at some sites such as DAA (north coast of Australia), SEY (the Indian Ocean), and CRI (west coast of India). These regions are influenced by nearby sources and the complex wind motion associated with the ITCZ and monsoon circulation, which affects transport of CH_4 from source regions. These conditions make it impossible for the model to reproduce the seasonal variations at these sites.

Statistics of monthly CO_2 and CH_4 biases against the GV analysis for the year 2008 are shown in Table 1 and Fig. 5. The overall correlation coefficients between the observed and modeled seasonal patterns at the GV sites are 0.86 for CO_2 and 0.61 for CH_4, which shows a high degree of consistency between the model and the GV analyses. High correlations are found particularly at oceanic sites: correlation coefficients are 0.97 and 0.70 for CO_2 and CH_4, respectively. Total model–observation differences (CRMSD in Table 1) are 0.51 and 1.02 for CO_2 and CH_4, respectively, and better performance are found at oceanic sites. Generally, the model underestimates the GV CO_2 amplitudes (total RSTD 0.77) and overestimates the GV CH_4 amplitudes (total RSTD 1.25). Simulated CO_2 and CH_4 at the tower sites show relatively poorer performance, particularly for the CH_4 case with correlations of 0.68 for CO_2 and 0.22 for CH_4. Tower sites are typically located near source regions on land. The use of the climatological flux dataset might make it difficult for the model to reproduce the large variations

GMDD
5, 2215–2258, 2012

observed at the tower sites. Model transport errors such as PBL height, vertical diffusion may also affect on the model's reproductivity of CO_2 concentration observed at tower sites. Except for the tower sites, the model shows statistically good performance at the oceanic, land, and free tropospheric sites.

3.3 Comparison with observations: synoptic variations

Daily averages from the simulated results and observations were compared at Mauna Loa, Hawaii (MLO; 19.5° N, 155.6° W, 3397 m a.s.l), South Pole, Antarctica (SPO; 89.98° S, 24.8° W, 2810 m a.s.l.), and Hateruma, Japan (HAT; 24.05° N, 124.8° E, 47 m a.s.l.). CO_2 and CH_4 data at MLO and SPO were provided by NOAA/ESRL in the WDCGG dataset. Daily CO_2 data at HAT were available from the Greenhouse Gases Trend Update (http://db.cger.nies.go.jp/g3db/ggtu/index.html) operated by the Center for Global Environmental Research (CGER), NIES. CH_4 data at HAT are hourly data (Tohjima et al., 2002, 2010; Tohjima, unpublished data, 2011). CH_4 observations at SPO are discrete observations and the other observations are continuous measurements. The continuous data and the 3-hourly model output were daily averaged based on local time for each site. As described in Sect. 2, GPV data are only available after December 2007, so the year 2008 meteorological data were used from January to November 2007 for the simulation.

Comparisons with continuous measurements of CO_2 and CH_4 show that the model was able to capture the observed synoptic and seasonal variations at each monitoring station (Fig. 6). Simulated CO_2 at MLO shows good agreement with the observed CO_2 though the model underestimates the observed spring maxima in 2008 and 2010, while simulated CH_4 at MLO slightly underestimates the observed CH_4 by about 20 ppb after 2009. MLO is remote from the large source regions, which are mainly on land, and transport is therefore a dominant factor for CO_2 variability. Unlike CO_2, CH_4 reacts with OH radicals during transport, which affects CH_4 variability. The interannual variability of atmospheric circulation is also important for the growth rate at MLO because transport determines the area the airmass come from, such as boreal Asia, the North Pacific,

GMDD
5, 2215–2258, 2012

or the tropical Atlantic (Higuchi et al., 2002; Patra et al., 2005b). In this mean, the model transport reproduces the overall features of observed CO_2 and CH_4 at MLO, but the use of climatological fluxes in this study gives some discrepancies. At SPO, another remote site away from strong source regions, the observed CO_2 has very small seasonal variations. A gap in the simulated CO_2 at the end of January 2009 is due to the offset correction described in Sect. 2. Differences between the model and the observed small variations in the first half of 2010 are slightly large at 2 ppm. Otherwise, the simulated CO_2 trend generally matches the observed one. In spite of the OH sink, the simulated CH_4 reproduces the observed CH_4 at SPO very well except for the second half of 2010, where there occurs a large increase in observed CH_4.

HAT is an island located on the East-Asian continental margin and is influenced by air masses transported from the Pacific Ocean in summer and from the continent in winter (Tohjima et al., 2002, 2010). Larger seasonal amplitudes and larger synoptic events are therefore observed here, relative to the background sites MLO and SPO. Though the CO_2 fluxes in the model are climatological, the model can simulate the observed seasonal pattern and occasional synoptic events such as the low concentrations in August 2008 and 2009. High-CO_2 events in winter are difficult to represent perfectly in the model, possibly due to transport of CO_2 from continental urban areas, but the model successfully simulates observed spring maxima. CH_4 at HAT is also well reproduced by the model; i.e., the model shows a clear seasonal pattern of summer and winter air mass exchanges. The model sometimes even captures sudden summer high CH_4. Basically HAT is covered by oceanic air mass and the CH_4 fluxes in the model are climatological; thus, these high-CH_4 events are thought to be CH_4 transported from continental CH_4 source regions.

Table 2 lists the statistics (ratio of standard deviations, correlation coefficients, overall biases, centered pattern root-mean-square differences) between daily averaged modeled and observed CO_2 and CH_4 at the three sites. As described above, only CH_4 at SPO is discrete data and the modeled CH_4 is taken from the same date and time as the observation. The statistics suggest that the model can simulate the observed daily

GMDD
5, 2215–2258, 2012

CO_2 and CH_4 variations fairly well with a correlation coefficient (r) > 0.8 at the three sites, except for CH_4 at MLO (r = 0.59). The calculated RSTDs are nearly 1 and overall biases are less than 1 ppm for CO_2 and 10 ppb for CH_4. CRMSDs are also below 1, indicating that the model performs reasonably well.

3.4 Comparison with observations: stratospheric profiles

There are few periodical high-precision observations of CO_2 and CH_4 in the stratosphere, but observations are made over Japan about once a year using a balloon-borne cryogenic sampler operated by Tohoku University, Japan (Nakazawa et al., 2002; Aoki et al., 2003). In this study we compare the simulated stratospheric profiles of CO_2 and CH_4 with the observed mean profiles from the balloon-borne data over Japan. The observed mean profiles and their standard deviations are obtained as follows: first the observed tracer concentrations over Sanriku (39.17° N, 141.8° E), Japan from 1985 to 2007 are averaged in each of five height bins: below 15 km, 15–20 km, 20–25 km, 25–30 km, above 30 km; then the concentrations at the highest level are shifted to match observations carried out on 22 August 2012 over Taiki-cho (42.48° N, 143.42° E), Japan (Nakazawa et al., 2002; Aoki et al., 2003; Nakazawa and Aoki, unpublished data, 2010). The simulated profiles on the same day over Sanriku are compared with the averaged observed profiles (Fig. 7). The simulated profile of CO_2 (corrected by age of air in the stratosphere, Sect. 2) is a close match to the observations with no bias on the day, and the difference between simulated and observed CO_2 profiles is within the standard deviation of the observed CO_2. The simulated CH_4 profile also shows very good agreement with the observed profiles within the observed standard deviation, though the simulated profile tends to be smaller than the observed one at a height of about 100–20 hPa. Recently, De Mazière et al. (2008) compared CH_4 profiles from the Atmospheric Chemistry Experiment-Fourier Transform Spectrometer (ACE-FTS) launched on August 2003 with those from HALOE. They found both instruments showed similar average profiles and variability from 15 km to 70 km, and though ACE-FTS showed slightly higher biases compared with HALOE, they concluded that their differences

GMDD
5, 2215–2258, 2012

were of the order of 5 % below 35 km (the target region of the present study) and were not significant because the error bars overlapped.

Though the model–observation comparison is for only one profile due to the lack of stratospheric observations, the agreement between the simulated and observed profiles shows that nudging the model stratosphere to the climatology field of CO_2 (corrected by age of air) and CH_4 (HALOE) works quite well with no bias on the observed day or within the observed standard deviations.

3.5 Global distributions

Overall, the model successfully reproduces the observed variations of CO_2 and CH_4 as described in the previous sections. In this section, we present the latitude–longitude distributions for both the model surface level and the column-averaged dry air mole fractions. Column-averaged dry air mole fractions, XCO_2, was obtained by weighting the concentration in each layer by the air mass in that layer from the model output at 21 pressure levels; thus,

$$XCO_2 = \sum_{n=L}^{21} {CO_2}^n \times \frac{\Delta P_n}{P_{\mathrm{srf}}}, \tag{2}$$

where ${CO_2}^n$ is the CO_2 concentration in layer n defined at a layer center in the pressure level coordinate, L is the lowest level of the grid (surface level), P_{srf} is surface pressure, and ΔP_n is the thickness of layer n in pressure. XCH_4 is calculated in the same way.

The temporal and spatial patterns of CO_2 and CH_4 are primarily governed by seasonal cycles of the sources, sinks, and atmospheric transport. Column abundances are also sensitive to changes in surface pressure and the tropopause height (e.g. Washenfelder et al., 2003). Figures 8 and 9 show the simulated surface CO_2 and CH_4 concentrations, column CO_2 and CH_4, and their differences for January and July 2010 at 13:00 LT, almost the same time as the GOSAT orbit descending node around 12:48 LT (Yoshida et al., 2011). Surface CO_2 concentrations in January are highest over

GMDD
5, 2215–2258, 2012

high-emission areas such as Siberia and Asia in the northern mid and high latitudes, and also over the equatorial regions (Africa, East South Asia, South America). Strong sinks in July are found in mid and high latitudes in Siberia and North America due to photosynthesis by land biomass in summer. Compared with the continental regions, the marine boundary layers exhibit low concentrations because of the absence of strong sources. In the Southern Hemisphere, the CO_2 concentrations are relatively homogenous, except for some limited continental areas such as South America. XCO_2 has less variation than surface CO_2, as expected. Strong sources over China, India, and Equatorial Africa and a strong sink in Siberia are still recognizable in XCO_2, particularly in July. Their difference (XCO_2 – surface CO_2) is generally smaller in equatorial regions due to the high tropopause height, and larger in northern high latitudes due to strong sources/sinks and low tropopause height. In July their difference is positive over mid and high northern latitudes due to large sinks at the surface. Nakazawa et al. (1993) found that the observed seasonal variation of CO_2 concentration showed a phase delay of about 1 month between the lower and upper troposphere by using long-term airborne observations over Japan. This fact supports the positive difference between XCO_2 and surface CO_2 over strong sink regions in mid and high northern latitudes.

High-CH_4 regions are simulated at the surface over land both in January and July, and CH_4 is higher in the Northern Hemisphere than in the Southern Hemisphere throughout the year due to large CH_4 emission in the Northern Hemisphere. XCH_4 exhibits the same trend as surface CH_4: relatively high in the Northern Hemisphere and low in the Southern Hemisphere. High-XCH_4 regions, which appear over South and Eastern Asia and Equatorial Africa, are associated with deep convection over these areas. Xiong et al. (2009) found a high-CH_4 plume over South Asia in the middle to upper troposphere associated with the monsoon season during July–September. These convective flows transport surface CH_4 to the upper tropopause and the model can capture such characteristics. While CO_2 is stable in the atmosphere, CH_4 reacts with $O(^1D)$ and Cl in the stratosphere in addition to chemical loss with OH radicals, creating a significant vertical decrease in the stratospheric concentration (Fig. 6). These

GMDD
5, 2215–2258, 2012

atmospheric sinks for CH_4 create a negative difference in (XCH_4 – surface CH_4) except for the equatorial region where the tropopause tends to be high due to strong convection.

Figure 10 shows the simulated surface CH_4 concentrations around Japan. The high-resolution model (0.5° × 0.5°; Fig. 9a) simulates a much clearer land–ocean contrast in CH_4 concentrations and synoptic-scale motions than the model with a resolution of 2° × 2° (Fig. 9b). An intrusion of air mass with low CH_4 concentrations from the Pacific is more sharply resolved in the 0.5° simulation than at 2° resolution. The 0.5° model appears to be able to resolve point sources of CH_4, such as highly populated urban areas (e.g. Tokyo), as shown by Maksyutov et al. (2008) for CO_2.

Figure 11 shows monthly zonal-mean latitudinal distributions of surface CO_2 and XCO_2, and their standard deviations at 13:00 LT, and Fig. 12 those of CH_4. Zonal-mean surface CO_2 shows a strong sink centered on about 60° N in July, which then spreads toward high and low latitudes in August. The seasonal amplitude for XCO_2 is about 9 ppm, which is about half of that for surface CO_2, 15 ppm. This tendency is consistent with previous modeling studies (e.g. Olsen and Randerson, 2004). Variances are large in northern mid and southern low latitudes, reflecting activity of land biosphere. The longitudinal variations in XCO_2 are about 2 ppm at maximum. The north-to-south gradient of the concentrations modeled at the surface level and that of XCH_4 concentrations simulated at 13:00 LT were similar in trend; i.e., the concentrations in the Northern Hemisphere are higher than those in the Southern Hemisphere. XCH_4 values at every latitude are shifted to lower concentrations due to chemical OH loss in the tropopause and the stratosphere. Small peaks in XCH_4 in equator are found in equatorial regions throughout the year, possibly due to strong vertical transport of surface CH_4 by deep cumulus convection (e.g. Patra et al., 2009; Terao et al., 2011). CH_4 variance at the surface is high over most of the latitude band, reflecting the various CH_4 sources over land. This tendency is the same for XCH_4 but the variance is less than half the surface value. Relatively large variances in southern high latitudes might be due to the high elevation of the Antarctic Continent at over 3000 m, which makes this region susceptible

GMDD
5, 2215–2258, 2012

to seasonal variations in tropopause height, as the stratospheric partial column of low CH_4 has a large weight.

4 Conclusions

We have developed the NIES transport model at a resolution of 0.5° × 0.5° × 47 sigma levels, driven by high-resolution meteorological data, GPV, with 0.5° × 0.5° resolution and 21 pressure levels (1000–10 hPa). This GPV-forced NIES TM has been designed to provide global high-resolution and near real-time a priori CO_2 and CH_4 concentrations for the GOSAT data retrieval algorithm to derive XCO_2 and XCH_4 at NIES. Since real-time fluxes of CO_2 and CH_4 are not available, the flux climatologies were used with the trend adjustment to the observed background concentrations. To overcome the problem of stratospheric transport in the model, the stratospheric part of the model was nudged to climatological values using three-dimensional CO_2 climatology that was adjusted to observed age of air for CO_2 and long-term satellite observations from HALOE for CH_4. We also updated the stratospheric part of the earlier version of a priori error variance–covariance matrices for CO_2 and CH_4 to give more realistic stratospheric values.

The model performance was assessed by comparing the model outputs with available observational records of atmospheric CO_2 and CH_4 concentrations. A large-scale, latitudinal distribution of the simulated annual mean CO_2 and CH_4 concentrations is found to be in good agreement with the analysis of GV sites, with overall annual biases and standard deviations of -0.14 ± 1.57 ppm and 0.88 ± 14.16 ppb, and with RMSDs of 2.28 ppm and 12.68 ppb at 260 CO_2 sites and 206 CH_4 sites, respectively. In particular, pole-to-pole gradients of CO_2 and CH_4 are reproduced exactly by the model with almost no biases. Comparison between monthly GV CO_2 and CH_4 and the model output showed that, despite large model–observation mismatch in monthly seasonal variations at some tower sites and some inland sites where large seasonal variations were observed, the model seasonal variations generally agreed well, particularly at

GMDD
5, 2215–2258, 2012

oceanic and free tropospheric sites, with GV values with averaged correction coefficients of 0.86 for CO_2 and 0.61 for CH_4 in terms of seasonal variations. The observed daily or discrete time series at MLO, SPO, and HAT are generally well reproduced by the model with statistically good performance, though some discrepancies were found, possibly due to the use of climatological fluxes. In the stratosphere, the simulated vertical profiles and growth rates agree well with the average profiles from balloon-borne observations over Japan within the observed standard deviations.

The global CO_2 and CH_4 distributions, and XCO_2 and XCH_4 obtained are in qualitative agreement with previous studies. The 0.5° model can resolves synoptic-scale motions and point sources than the 2.0° model does. Seasonal amplitudes in zonal-mean XCO_2 are found to be almost half those for surface CO_2. Zonal-mean XCH_4 shows different features from XCO_2 because of the chemical sinks in the atmosphere and its sensitivity to tropopause height.

These validations suggest that the model is able to reproduce fairly reasonable global concentrations as well as synoptic variations and give confidence in quantitative analysis of CO_2 and CH_4 cycles using the model, and its use in providing a priori concentrations for satellite retrievals. This model has been implemented on the GOSAT DHF system and has been run every observed day, and the simulated results have been used for a priori concentrations for GOSAT XCO_2 and XCH_4 retrievals. As future satellite instruments like OCO-2 are expected to have smaller footprints or higher precision to observe greenhouse gases more precisely, the ability to simulate a priori concentrations with a higher-resolution model would be useful in reducing error in a priori concentrations. Thus the updated high-resolution concentrations and VCMs provided by the developed model have the potential to be powerful tools for a priori of satellite data retrievals as well as for the high-resolution global modeling of greenhouse gases.

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GMDD
5, 2215–2258, 2012

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GMDD

5, 2215–2258, 2012

Global simulations of CO_2 and CH_4 for sattelite data retrievals

T. Saeki et al.

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GMDD
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5, 2215–2258, 2012

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GMDD

5, 2215–2258, 2012

Table 1. Statistics to show the model performance against GLOBALVIEW analyses of CO_2 and CH_4: annual-mean biases and seasonal variations.

Tracer	Site type[1]	Number of sites	Annual mean		Seasonal variation		
			Biases and standard deviations	RMSD[2]	RSTD[3]	Correlation coefficients[4]	CRMSD[5]
CO_2	Oceanic	74	−0.02 ± 1.03	0.17	0.83	0.97	0.27
	Land	93	−0.35 ± 1.46	3.36	0.83	0.91	0.41
	Tower	17	3.30 ± 2.56	13.60	0.64	0.68	0.74
	Free tropos.	76	−0.14 ± 0.61	6.76	0.78	0.93	0.39
	Total	260	−0.01 ± 1.57	2.28	0.77	0.86	0.51
CH_4	Oceanic	58	0.90 ± 7.88	6.89	1.14	0.70	0.84
	Land	83	4.33 ± 17.92	39.43	1.26	0.62	1.01
	Tower	8	7.53 ± 27.42	21.30	1.75	0.22	1.80
	Free tropos.	57	−5.09 ± 6.81	38.40	1.18	0.62	0.96
	Total	206	0.88 ± 14.16	12.68	1.25	0.61	1.02

[1] Site types: the same as those defined in Fig. 2.
[2] Root-mean-square differences (RMSD).
[3] Ratio of standard deviation (SD), which is calculated by dividing the model SD by the observed SD.
[4] Pearson's correlation coefficient.
[5] Centered pattern root-mean-square differences (CRMSD), which is a measure of the distance between model and observation.

GMDD
5, 2215–2258, 2012

Table 2. Statistics to show the model performance against continuous measurements for daily-mean CO_2 and CH_4 at HAT (Hateruma), MLO (Mauna Loa), and SPO (South Pole). Observed CH_4 data at SPO are discrete data, and the simulated results corresponding to the measurement date are used for the comparisons. Statistics are defined as in Table 1.

Tracers	Site	Time interval of comparisons	Period of comparisons	Number of data	RSTD	Correlation coefficients	Overall bias	CRMSD
CO_2	HAT	Daily	1 Jan 2007–30 Aug 2011	1638	0.82	0.90	−0.29	0.44
	MLO	Daily	1 Jan 2007–31 Dec 2010	1351	0.88	0.96	−0.41	0.28
	SPO	Daily	1 Jan 2007–31 Dec 2010	1439	1.12	0.98	0.73	0.25
CH_4	HAT	Daily	1 Jan 2007–31 Dec 2010	1421	0.95	0.83	2.44	0.57
	MLO	Daily	1 Jan 2007–31 Dec 2010	1377	0.82	0.59	−9.83	0.84
	SPO	Event	1 Jan 2007–31 Dec 2010	191	0.92	0.96	−1.19	0.30

GMDD
5, 2215–2258, 2012

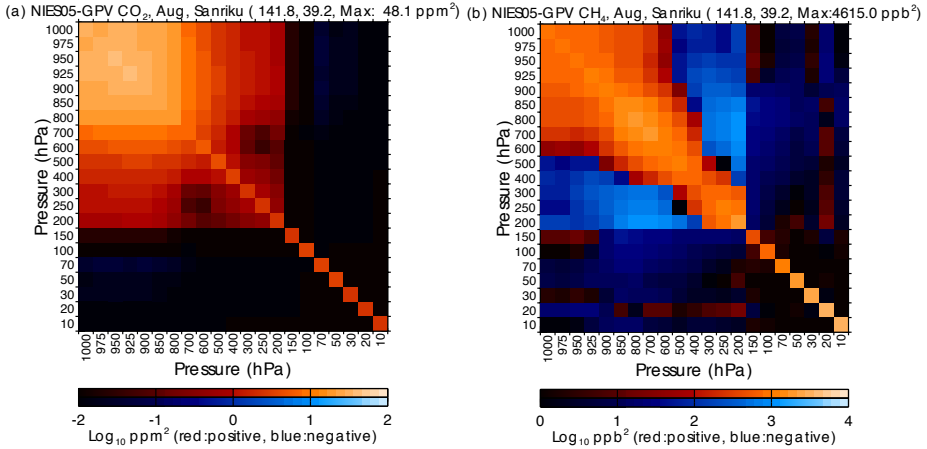

Fig. 1. A priori error variance–covariance matrices of **(a)** CO_2 and **(b)** CH_4 over Sanriku, Japan (141.8° E, 39.2° E) in August at 21 pressure levels from 1000 to 10 hPa. Color scales are logarithmic, with ranges of 0.01–100 ppm^2 for CO_2 and 1–10 000 ppb^2 for CH_4. Warm and cold colors indicate positive and negative correlations, respectively. Dark and light colors indicate small and large variance–covariance values, respectively.

GMDD
5, 2215–2258, 2012

Global simulations of CO_2 and CH_4 for sattelite data retrievals

T. Saeki et al.

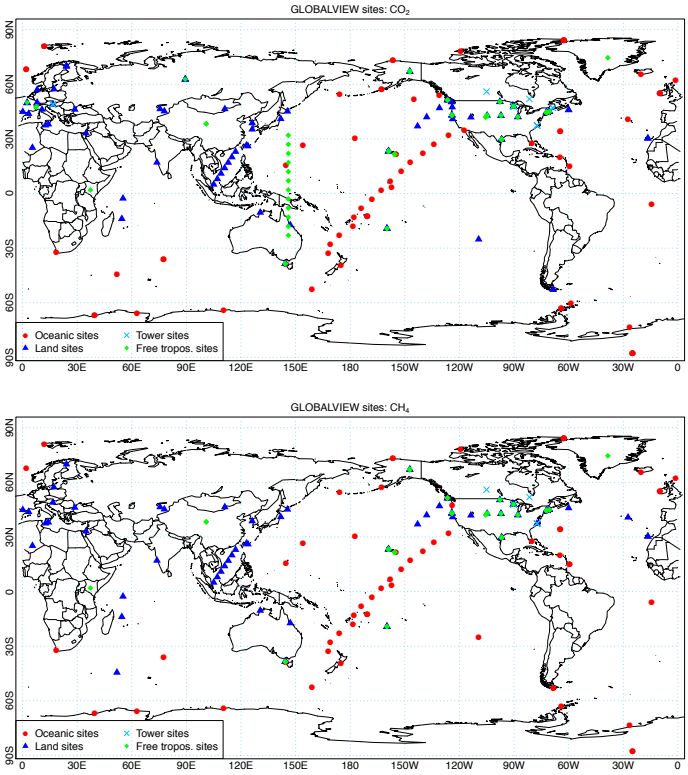

Fig. 2. GLOBALVIEW site locations used for the comparisons of CO_2 (top) and of CH_4 (bottom). "Oceanic"; sites with marine-boundary-layer (MBL) marks in the GV dataset (gv_table.co2 and gv_table.ch4); "land", sites below 3000 m which are neither MBL sites nor tower sites; "tower", tower sites with sampling platform code "3" in GV file names; "free tropos.", free troposphere sites located above 3000 m and marked as non-MBL sites, mostly airborne observational points.

GMDD
5, 2215–2258, 2012

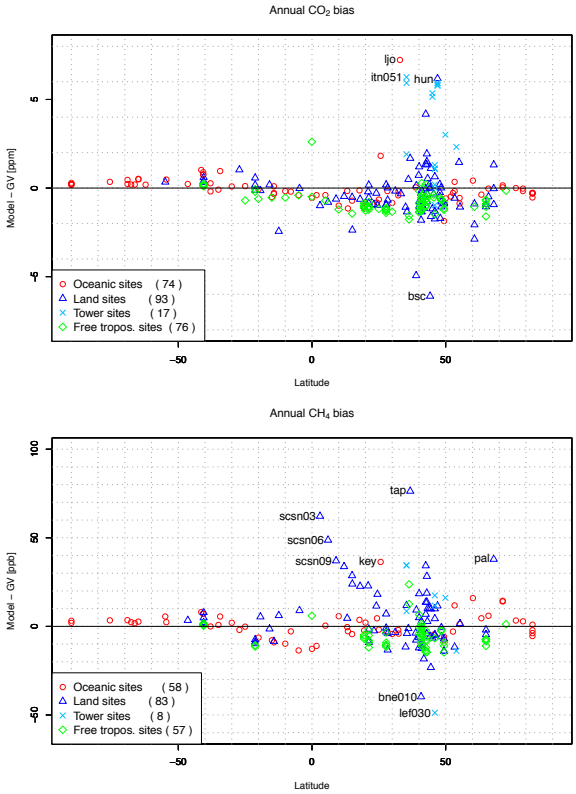

Fig. 3. Latitudinal distributions of differences in annual mean between the simulated and GV data for CO_2 (top) and CH_4 (bottom) at GV monitoring sites. The simulated results at 13:00–16:00 LT were used for the comparison. Letters in the plot represent the GV site code. Site types in the legend are the same as those defined in Fig. 2. Numbers in legend parentheses indicate the numbers of GV sites used for the analysis.

GMDD
5, 2215–2258, 2012

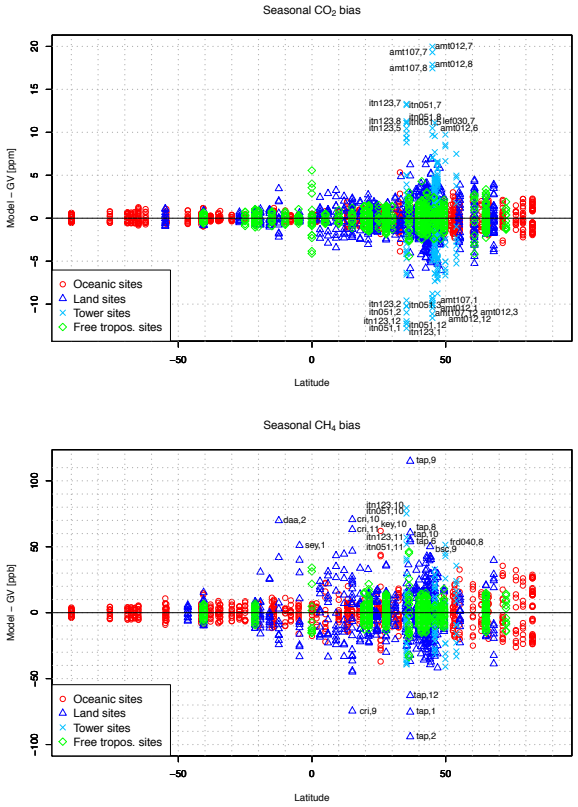

Fig. 4. Same as Fig. 3, but for monthly biases for CO_2 (top) and CH_4 (bottom). Letters and numbers in the plot represent GV site code and month, respectively.

GMDD
5, 2215–2258, 2012

Global simulations of CO_2 and CH_4 for sattelite data retrievals

T. Saeki et al.

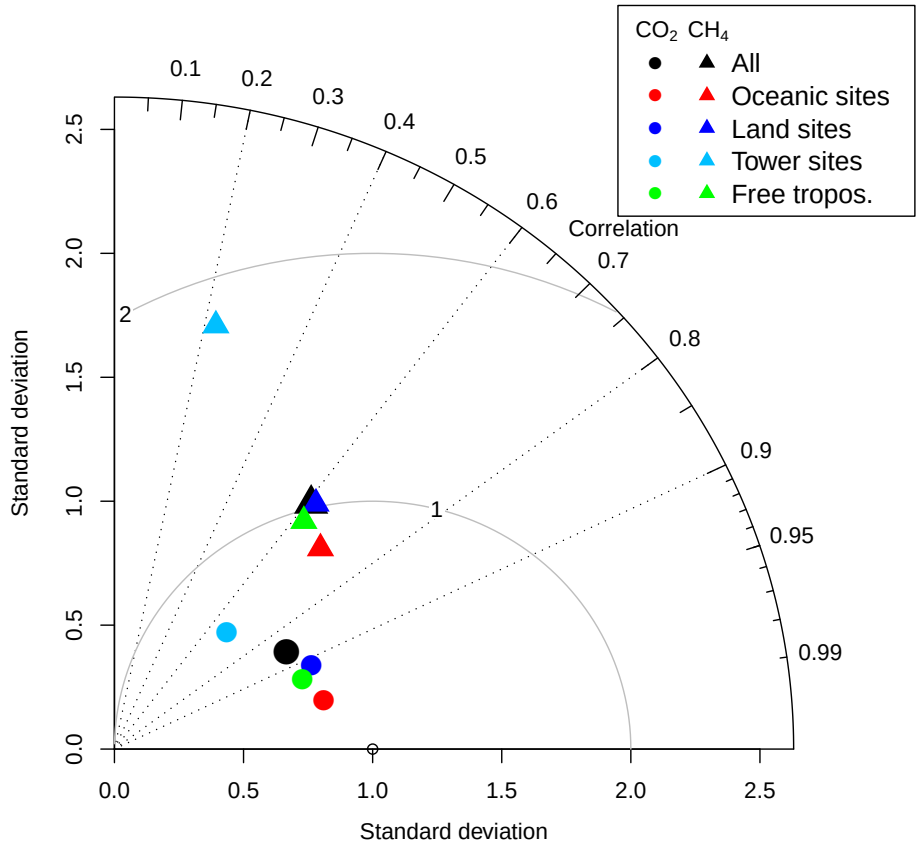

Fig. 5. Normalized Taylor diagram showing a statistical comparison (Pearson's correlation coefficient, normalized standard deviation, and centered root-mean-square difference) between simulated CO_2 (solid circles) and CH_4 (solid triangles), and GV analysis, for monthly seasonal variations.

GMDD
5, 2215–2258, 2012

Global simulations of CO_2 and CH_4 for sattelite data retrievals

T. Saeki et al.

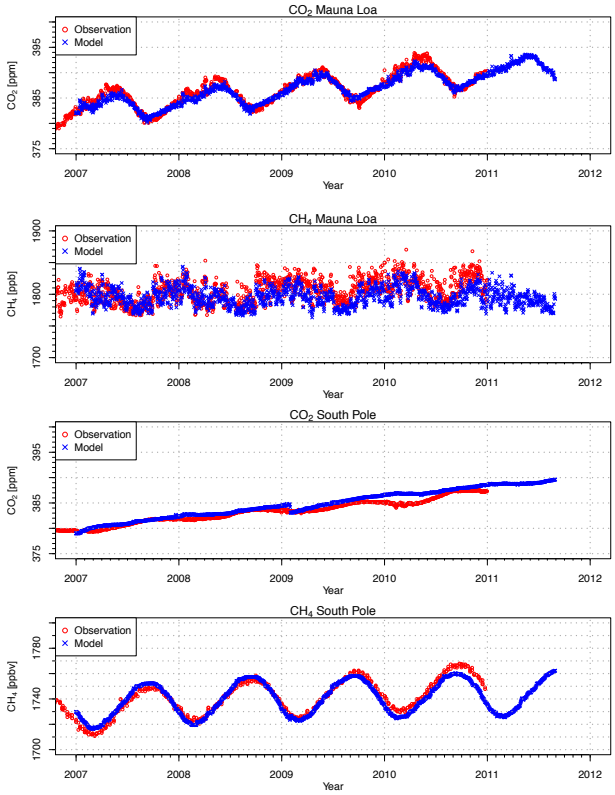

Fig. 6. Caption on next page.

GMDD
5, 2215–2258, 2012

Global simulations of CO_2 and CH_4 for sattelite data retrievals

T. Saeki et al.

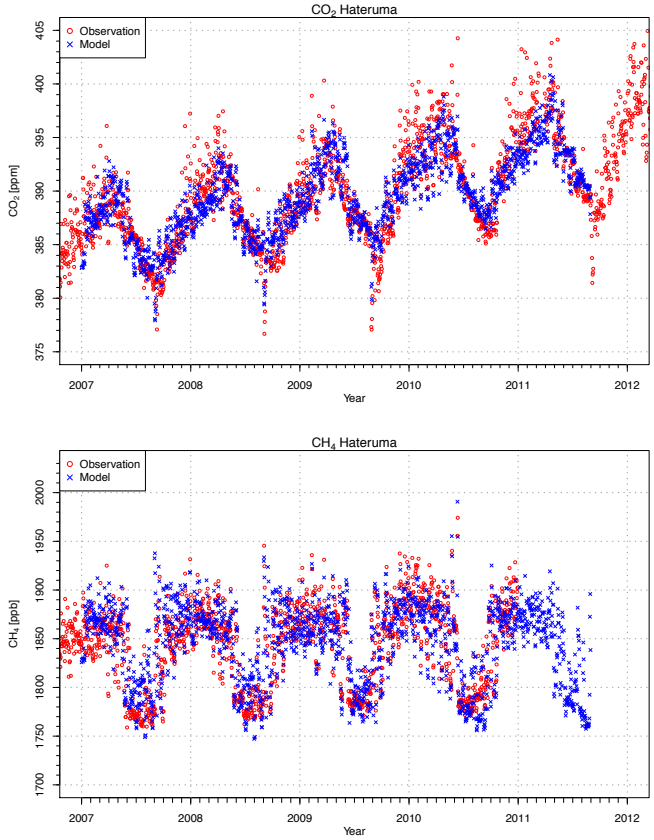

Fig. 6. Observed and simulated time series of daily mean CO_2 and CH_4 concentrations at Mauna Loa, South Pole, and Hateruma sites. CH_4 observations at South Pole are discrete data and are plotted directly; quasi-continuous data at the other sites and 3-hourly model output are daily averages. A gap in the simulated CO_2 at SPO at the end of January 2009 is due to the offset correction (see text).

GMDD
5, 2215–2258, 2012

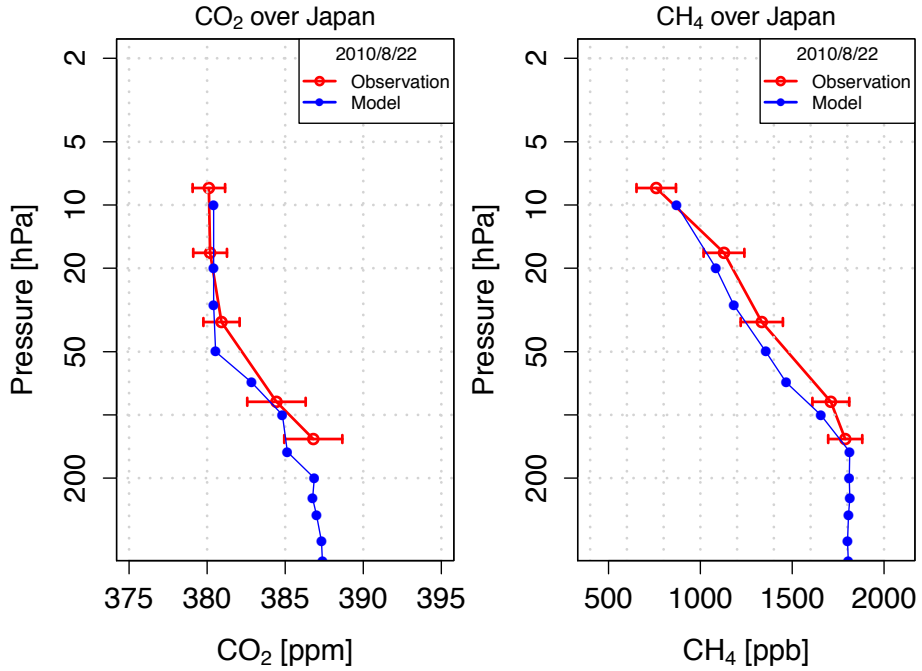

Fig. 7. Vertical profiles of CO_2 (left) and CH_4 (right) in the stratosphere over Japan. The blue lines indicate NIES TM simulated data on 22 August 2010. The red lines are observed average profiles and their standard deviations derived from balloon-borne measurements over Sanriku, Japan (141.8° E, 39.17° N).

GMDD
5, 2215–2258, 2012

Global simulations of CO_2 and CH_4 for sattelite data retrievals

T. Saeki et al.

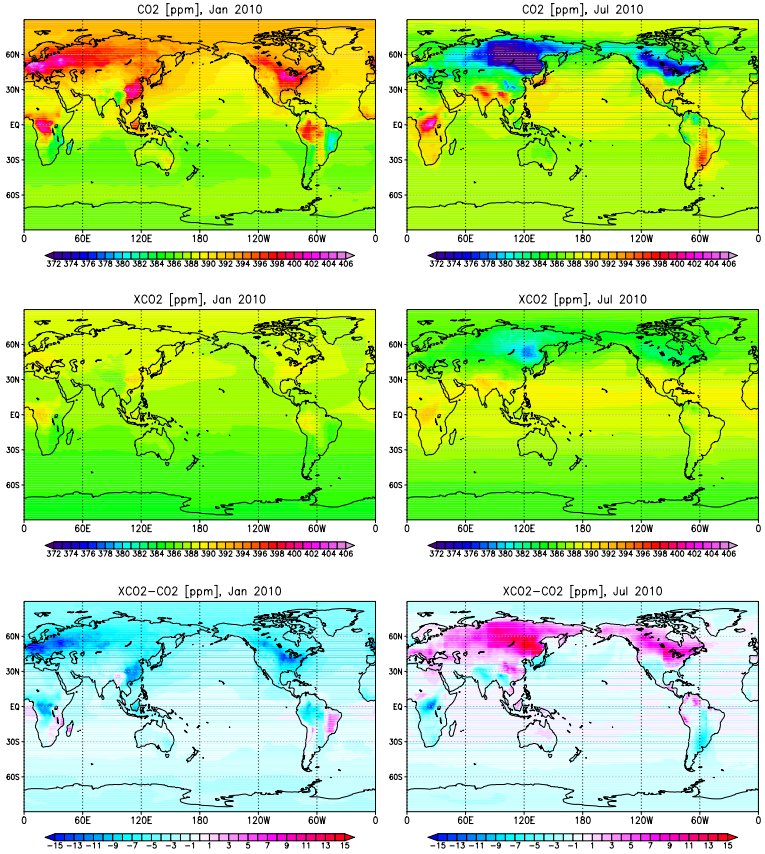

Fig. 8. Simulated monthly mean surface CO_2 (top), XCO_2 (middle), and their differences (bottom) at 13:00 LT in January (left column) and July (right column) 2010. Ranges of color scales are 372–406 ppm for CO_2 and XCO_2, and −15 to 15 ppm for the differences.

GMDD
5, 2215–2258, 2012

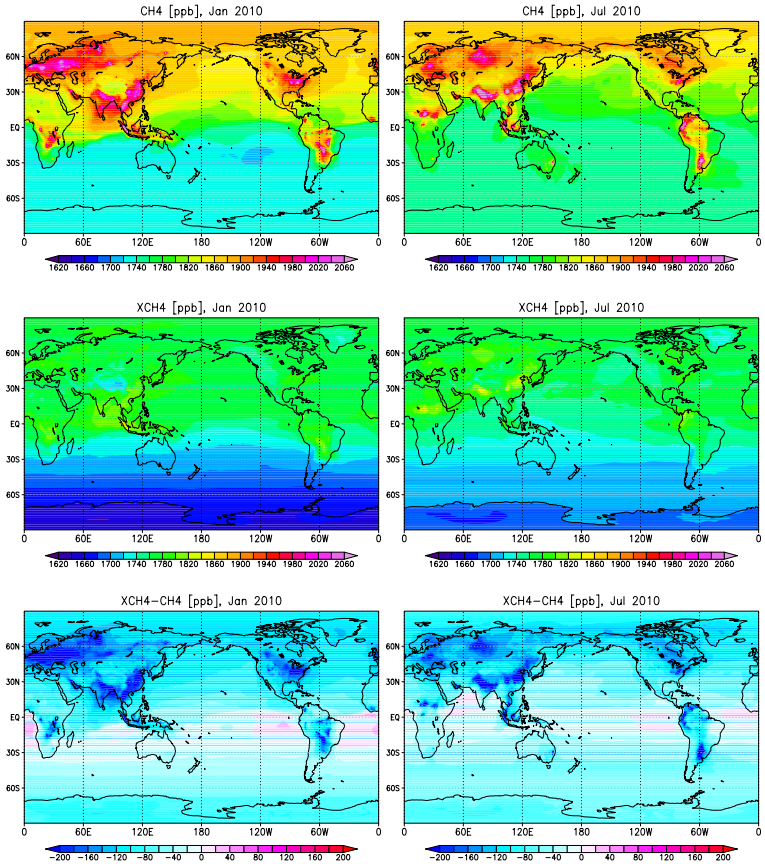

Fig. 9. Same as Fig. 7, but for CH_4, XCH_4, and their differences. Ranges of color scales are 1620–2060 ppb for CH_4 and XCH_4, and −200 to 200 ppb for the differences.

GMDD
5, 2215–2258, 2012

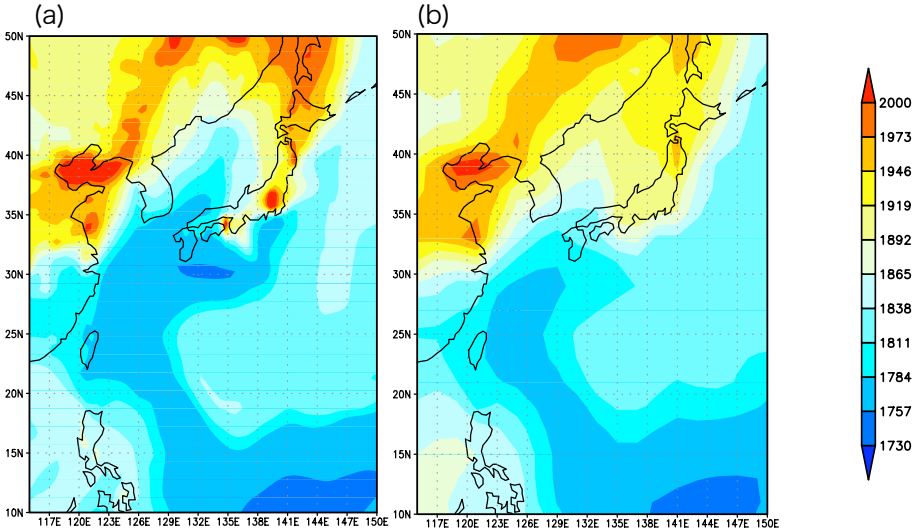

Fig. 10. Simulated surface CH_4 concentrations (ppb) around Japan at 13:00 JST on 6 July 2008 with a horizontal resolution of **(a)** 0.5° and **(b)** 2.0°.

GMDD
5, 2215–2258, 2012

Global simulations of CO_2 and CH_4 for sattelite data retrievals

T. Saeki et al.

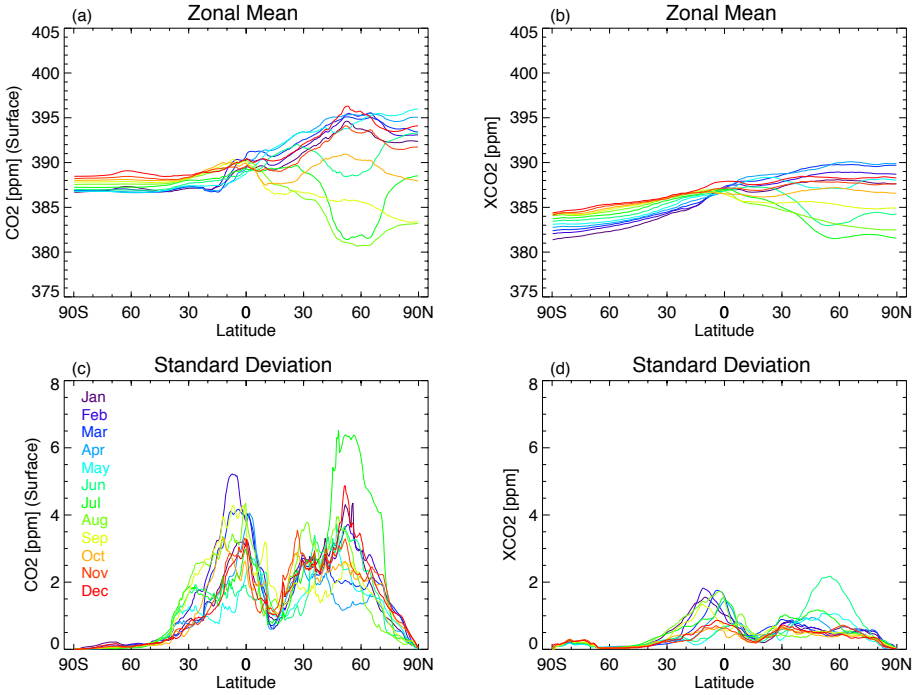

Fig. 11. Monthly zonal-mean latitudinal distributions of **(a)** CO_2 and **(b)** XCO_2 for 2010 and **(c, d)** their standard deviations against longitudinal variation.

GMDD
5, 2215–2258, 2012

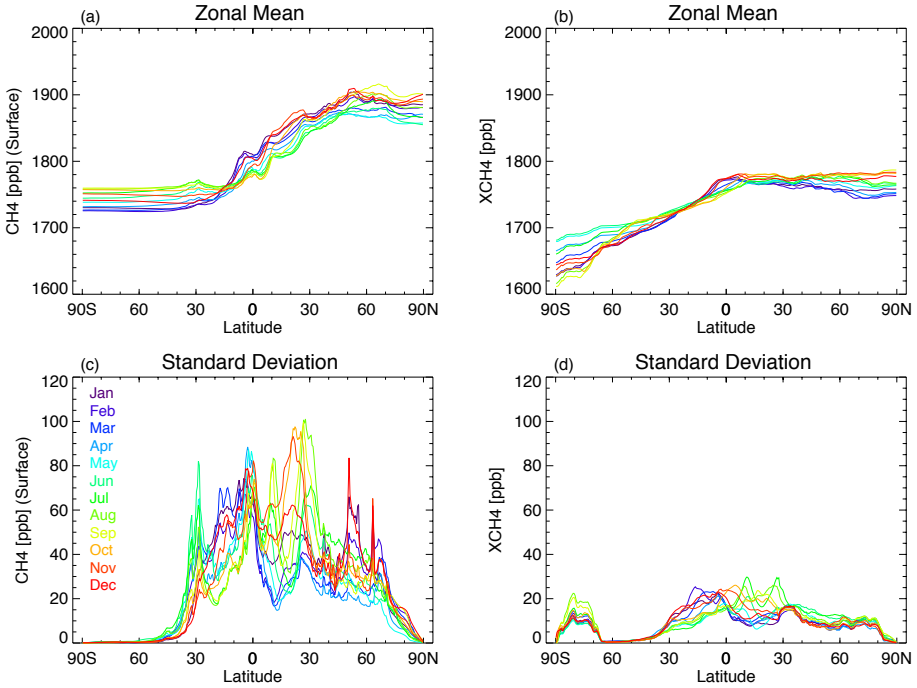

Fig. 12. Same as Fig. 11, but for CH_4 and XCH_4.

GMDD
5, 2215–2258, 2012

Global simulations of CO_2 and CH_4 for sattelite data retrievals

T. Saeki et al.