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Atmospheric Chemistry and Physics Discussions



# TransCom model simulations of $CH_4$ and related species: linking transport, surface flux and chemical loss with $CH_4$ variability in the troposphere and lower stratosphere

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## Abstract

A transport model intercomparison experiment (TransCom-$CH_4$) has been designed to investigate the roles of surface emissions, transport and chemical loss in simulating the global methane distribution. Model simulations were conducted using twelve models and four model variants and results were archived for the period of 1990–2007. The transport and removal of six $CH_4$ tracers with different emission scenarios were simulated, with net global emissions of $513 \pm 9$ and $514 \pm 14$ Tg $CH_4$ $yr^{-1}$ for the 1990s and 2000s, respectively. Additionally, sulfur hexafluoride ($SF_6$) was simulated to check the interhemispheric transport, radon ($^{222}Rn$) to check the subgrid scale transport, and methyl chloroform ($CH_3CCl_3$) to check the chemical removal by the tropospheric hydroxyl radical (OH). The results are compared to monthly or annual mean time series of $CH_4$, $SF_6$ and $CH_3CCl_3$ measurements from 8 selected background sites, and to satellite observations of $CH_4$ in the upper troposphere and stratosphere. Most models adequately capture the vertical gradients in the stratosphere, the average long-term trends, seasonal cycles, interannual variations and interhemispheric gradients at the surface sites for $SF_6$, $CH_3CCl_3$ and $CH_4$. The vertical gradients of all tracers between the surface and the upper troposphere are consistent within the models, revealing vertical transport differences between models. We find that the interhemispheric exchange rate ($1.39 \pm 0.18$ yr) derived from $SF_6$ is faster by about 11 % in the 2000s compared to the 1990s. Up to 60 % of the interannual variations in the forward $CH_4$ simulations can be explained by accounting for the interannual variations in emissions from biomass burning and wetlands. We also show that the decadal average growth rate likely reached equilibrium in the early 2000s due to the flattening of anthropogenic emission growth since the late 1990s. The modeled $CH_4$ budget is shown to depend strongly on the troposphere-stratosphere exchange rate and thus to the model's vertical grid structure and circulation in the lower stratosphere. The 15-model median $CH_4$ and $CH_3CCl_3$ atmospheric lifetimes are estimated to be $9.99 \pm 0.08$ and $4.61 \pm 0.13$ yr,





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respectively, with little interannual variability due to transport and temperature as noted by the $\pm 1\,\sigma$.

## 1 Introduction

The variability of atmospheric $CH_4$ depends on the spatio-temporal variations of the surface fluxes, atmospheric transport and destruction due to OH, Cl and $O^1D$ chemistry. In recent years, measurements of $CH_4$ and related species are being conducted at an increasingly large number of sites at hourly or daily time intervals and with high instrumental precision (Rasmussen and Khalil, 1984; Aoki et al., 1990; Dlugokencky et al., 1998; Cunnold et al., 2002; WDCGG, 2010 for a complete list of observational programs). Satellite $CH_4$ observations from SCanning Imaging Absorption SpectroMeter for Atmospheric CartograpHY (SCIAMACHY), Atmospheric Infrared Sounder (AIRS), and Greenhouse Gases Observing SATellite (GOSAT) are also becoming available, albeit at a lower precision (Frankenberg et al., 2008; Xiong et al., 2008; Yoshida et al., 2011). Significant developments in understanding the $CH_4$ emission distributions have been achieved in the past few years through forward modeling (e.g., Fung et al., 1991; Gupta et al., 1996; Houweling et al., 2000; Dentener et al., 2003; Wang et al., 2004; Patra et al., 2009a). Inverse model results show the ability of the models to reproduce the observed atmospheric $CH_4$ trends and variabilities within the uncertainty of the processes involved (Hein et al., 1997; Houweling et al., 1999; Mikaloff-Fletcher et al., 2004; Chen and Prinn, 2006; Bousquet et al., 2006; Bergamaschi et al., 2009). However, further improvements (reduction in the posterior emission uncertainty) of inverse modeling results depend on a better quantification of (the errors in) the prior emissions and sinks, and on error reductions in forward model transport. Presently, inverse estimates of global $CH_4$ emissions range between 500 to 600 $Tg\,yr^1$, depending on the transport properties and the chemical loss parameterization in the forward models. Individual flux components vary by even greater percentages (e.g., Matthews and Fung, 1987; Yan et al., 2009).





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The performance of atmospheric transport models has been investigated within the TransCom project since the early 1990s for the non-reactive tropospheric species, such as sulfur hexafluoride ($SF_6$) and carbon dioxide ($CO_2$) (Law et al., 1996, 2008; Denning et al., 1999). Convective parameterizations in CTMs have been tested through simulation of Radon ($^{222}Rn$), which has a radioactive decay half-life of 3.8 days (e.g., Jacob et al., 1997). The full chemistry model simulations of reactive species with a focus on ozone ($O_3$) chemistry have also been tested using multiple CTMs, where $CH_4$ is treated as a tracer with a prescribed concentration evolution (Stevenson et al., 2006 and references therein). Note that most, if not all, full chemistry models do not treat $CH_4$ in an interactive manner, because of its long lifetime. The $CH_4$ lifetime ranged from 6.3 to 12.5 yr due to large range of simulated OH concentrations in the participating models (Stevenson et al., 2006). A more conservative estimate of $CH_4$ lifetime is required for calculating the global warming potential (GWP) for $CH_4$ and its impact on climate change, or developing effective emission mitigation policies. For example, a 26 % decrease in global burden of OH can lead to an increase in GWP for $CH_4$ up to 40 %, when the radiative forcing due to $CH_4$, relative to carbon dioxide ($CO_2$) on per kilogram basis, is integrated over 100 yr (Shindell et al., 2009).

The aim of the TransCom-$CH_4$ experiment described in this paper is to quantify the role of transport, emission distribution and chemical loss in simulating the interhemispheric (IH) gradient, seasonal cycle, synoptic variation and diurnal cycle of $CH_4$. The dependence of the $CH_4$ budget on model vertical transport in the stratosphere will be further analysed. We setup long simulation period (1988–2007, including two years of spin-up) for the following reasons: (1) in the 1990s and 2000s methane growth rates have fluctuated between 15 ppb $yr^{-1}$ to −5 ppb $yr^{-1}$ (Dlugokencky et al., 1998; Simpson et al., 2006; Rigby et al., 2008), and (2) we would like to obtain a better understanding of the role of emissions (using a set of six $CH_4$ emissions scenarios), chemical loss, and transport model characteristics, such as the stratosphere-troposphere exchange (STE) and the IH exchange rates on $CH_4$ concentration variations in the troposphere. The proposed 18-yr simulation period allows a proper quantification of the removal





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fluxes in the troposphere and stratosphere and of the influence of transport processes on these removal rates. Since the previous TransCom intercomparison experiments spanned only a few years, the 18 yr of $SF_6$ simulation allows us to track the interannual variability (IAV) in the IH exchange rate for the first time. We also discuss the dependence of $CH_4$ and $CH_3CCl_3$ lifetimes on the model grid structure and transport, as well as the transport and temperature as drivers of IAVs in lifetimes.

In Sect. 2, we describe the experimental protocol, followed by the key information on the participating models and analysis methodology. We focus this analysis (Sect. 3) on the comparison of model results with atmospheric observations of $SF_6$, $CH_3CCl_3$ and $CH_4$ at 8 surface sites and the salient differences in model properties. An attempt is also made to understand possible implications of (1) inert tracer ($SF_6$) transport or short-lived radioactive tracer ($^{222}Rn$) transport on chemically active species ($CH_3CCl_3$, $CH_4$), and (2) the effect of the OH abundance, as constrained by $CH_3CCl_3$ on $CH_4$. Scope for further analysis using the TransCom-$CH_4$ database and conclusions are given in Sects. 4 and 5, respectively.

## 2 Models, measurements and methods

Previous TransCom experiments focused on chemically non-reactive species ($SF_6$, $CO_2$, $^{222}Rn$). A $CH_4$ intercomparison requires the introduction of atmospheric chemistry. Additionally, the sources and atmospheric lifetime of $CH_4$ are distinctly different from $CO_2$, which may provide a different view on transport model differences. Detailed documentation of the requested simulation is available in the TransCom-$CH_4$ protocol (Patra et al., 2010). Transport model simulations are requested for the period of 1 January 1990 to 31 December 2007, after a spin-up of 2-yr (1988–1989) using analyzed or atmospheric general circulation model (AGCM) meteorology or a combination of both (referred here as AGCM-nudged). A schematic diagram of TransCom-$CH_4$ model intercomparison set-up is shown in Fig. 1.





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## 2.1 Photochemical and surface loss processes

The following chemical removal reactions for $CH_4$ (R1–R3) and $CH_3CCl_3$ (R4–R6) are prescribed in the forward simulations.

$$CH_4 + OH \xrightarrow{k_{OH}} CH_3 + H_2O \qquad \left[k_{OH} = 2.45 \times 10^{-12} \exp(-1775/T)\right] \tag{R1}$$

$$CH_4 + O^1D \xrightarrow{k_{O^1D}} \text{Products} \qquad \left[k_{O^1D} = 1.5 \times 10^{-10}\right] \tag{R2}$$

$$CH_4 + Cl \xrightarrow{k_{Cl}} CH_3 + HCl \qquad \left[k_{Cl} = 7.3 \times 10^{-12} \exp(-1280/T)\right] \tag{R3}$$

$$CH_3CCl_3 + OH \xrightarrow{k_{OH}} \text{Products} \qquad \left[k_{OH} = 1.64 \times 10^{-12} \exp(-1520/T)\right] \tag{R4}$$

$$CH_3CCl_3 \xrightarrow{J_{CH_3CCl_3}} \text{Products} \tag{R5}$$

$$CH_3CCl_3 + OH \xrightarrow{\text{deposition}_{CH_3CCl_3}} \text{Oceanic} \qquad CH_3CCl_3 \tag{R6}$$

The temperature-dependent reaction rates ($k$) are taken from the JPL synthesis of chemical kinetics (Sander et al., 2006). The monthly-mean OH fields are provided here for online calculation in the model by combining the semi-empirically calculated tropospheric (Spivakovsky et al., 2000) and 2-dimensional (2-D) model simulated stratospheric distributions. For $CH_4$ reactions with OH, Cl and $O^1D$ radicals in the stratosphere, parameterized loss rates $[k_{O^1_D} \times O^1D + k_{Cl} \times Cl]$ are provided, which are based on the Cambridge 2-D model (Velders, 1995).

The tropospheric OH field is reduced by 8 %, an amount that was required to optimize the agreement between the TM5 simulated and observed $CH_3CCl_3$ decline since 2000 (Huijnen et al., 2010). The model simulations performed here allow us to verify whether observations of $CH_3CCl_3$ can also be reproduced for a longer simulation period, i.e., 1990–2007, by TM5 and a variety of other models. The supplied OH field has





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about equal OH abundance in the NH and the SH (Spivakovsky et al., 2000). Since the NH/SH OH-ratio in full chemistry model simulations varies between 1.1 and 1.5 (Krol et al., unpublished data, 2008, based on the model intercomparison described in Shindell et al., 2006) we encouraged modelers to submit another set of simulations using their preferred OH field, e.g. obtained by a full chemistry version of their model.

For $CH_3CCl_3$, the photolysis rates $J$ due to solar UV radiation are provided from ACTM (Patra et al., 2009a) Because the resolution in the stratosphere varies widely between models, it is necessary to scale the stratospheric loss of MCF to a common value. This value is calculated by mass-weighted averaging:

$$J_{\mathrm{av,CH_3CCl_3}} = \frac{\sum_{i,j,k} J_{\mathrm{CH_3CCl_3}}(i,j,k) \times M(i,j,k)}{\sum_{i,j,k} M(i,j,k)} \quad (1)$$

Here, $M(i,j,k)$ denotes the air mass in gridbox $(i,j,k)$ from ACTM. Modelers were required to scale their interpolated $J_{\mathrm{MCF}}$ field to match the $J_{\mathrm{av,MCF}}$ field of the mass-weighted annual and global mean $J_{\mathrm{av,MCF}}$ value of $7.959 \times 10^{-8}\,\mathrm{s}^{-1}$ (Similarly, the annual and global mean rate constant for $CH_4$ oxidation due to stratospheric Cl and $O^1D$ combined is rescaled to $2.069 \times 10^{-10}\,\mathrm{s}^{-1}$).

The monthly deposition velocities ($\mathrm{deposition}_{\mathrm{CH_3CCl_3}}$; units: $\mathrm{m\,s}^{-1}$) of $CH_3CCl_3$ to ocean surfaces are provided (Krol et al., 1998; Kanakidou et al., 1999). This sink should be applied in the model as:

$$\mathrm{CH_3CCl_3} = (\mathrm{CH_3CCl_3})_0 \times \exp\left(-\mathrm{deposition}_{\mathrm{CH_3CCl_3}} \times \frac{1}{dz} \times dt\right) \quad (2)$$

where, $dz$ = atmospheric lowest layer depth (m), $dt$ = timestep (s), and subscript 0 indicates initial concentration.

Radon decays in the atmosphere with a half-life of 3.8 days, and this decay is calculated in the model at each timestep, following

$$^{222}\mathrm{Rn} = \left(^{222}\mathrm{Rn}\right)_0 \times \exp(-dt \times 2.11 \times 10^{-6}) \quad (3)$$





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where $^{222}Rn$ is the radon mixing ratio at all gridpoints. This setup follows the recommendation of World Climate Research Programme (Jacob et al., 1997).

Due to long timescales of $CH_4$ and $CH_3CCl_3$ oxidation and vertical transport in the stratosphere (age-of-air $\sim$5 yr), several years of spin-up are required to establish realistic $CH_4$ and $CH_3CCl_3$ vertical profiles throughout the model atmosphere. A set of 3-D initial conditions, prepared following a 10-yr spin-up simulation by ACTM, is made available for 1 January 1988 for $CH_4$, $SF_6$ and $CH_3CCl_3$. $CH_4$, $SF_6$ and $CH_3CCl_3$ concentrations at South Pole (SPO) are 1655 ppb, 1.95 ppt and 130 ppt, respectively, for January 1988. Radon will be spun-up quickly due to its half-life of several days. Hence, its initial concentration is set to zero.

## 2.2 Fluxes

The typical seasonal variations of the six $CH_4$ emission scenarios are shown in Fig. 2a. Annual total emissions time series are depicted in Fig. 2b for $CH_4$, and in Fig. 2c for $SF_6$ and $CH_3CCl_3$. The following source and sink components of $CH_4$ were considered for the emissions of the six $CH_4$ tracers listed in Table 1:

1. Interannually varying anthropogenic emissions (IAV ANT) are based on annual mean $1^\circ \times 1^\circ$ maps from the Emission Database for Global Atmospheric Research (EDGAR; version 3.2/FT) (Olivier and Berdowski, 2001). The combination of different emission categories and the inter-/extra-polation of EDGAR emission maps for the years 1990, 1995, 2000 are described elsewhere (Patra et al., 2009a).

2. Anthropogenic emissions (IAV ANT E4) are based on the more advanced EDGAR database (version 4.0) (http://edgar.jrc.ec.europa.eu), where $1 \times 1^\circ$ emission maps are available for each year until 2005. The 2005 emissions were also used for the 2006–2008 period.

3. Cyclostationary natural emissions (CYC NAT), such as those from all types of natural wetlands, domestic and large-scale biomass burning, and termites are





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based on the GISS inventory (Matthews and Fung, 1987; Fung et al., 1991), and the emissions due to rice paddies are taken from Yan et al. (2009). All these emissions are scaled as in Patra et al. (2009a). Though predominantly anthropogenic, emissions from rice cultivation are included in this category because its seasonal cycle is controlled by seasonal rainfall and temperature. The emissions due to oceanic exchange ($\sim 10\,\mathrm{Tg\,CH_4\,yr^{-1}}$) are distributed over the coastal region (Lambert and Schmidt, 1993; Houweling et al., 1999) and mud volcano emissions are based upon Etiope and Milkov (2004).

4. Wetland emissions with interannual variation (IAV WL) have been derived from the wetland emission module of the ORganizing Carbon and Hydrology in Dynamic EcosystEms (ORCHIDEE) terrestrial ecosystem model (Ringeval et al., 2010). This model uses satellite-derived area of inundation for the period of 1994–2000 (Prigent et al., 2007). The emission is scaled by a multiplication factor of 0.76 to match the wetland emission component in CYC NAT. An average seasonal cycle is used for the rest of the simulation periods (1988–1993 and 2001–2008).

5. Second set of wetland and rice emissions (IAV WLe) is obtained from the Vegetation Integrative Simulator for Trace gases (VISIT) terrestrial ecosystem model (Ito, 2010), which calculates inundated area based on analyzed rainfall, temperature (Mitchell and Jones, 2005). The rice and wetland emissions are scaled by 0.895 and 0.69, respectively, to match with CYC NAT.

6. Biomass burning emissions (IAV BB) are taken from the Global Fire Emission Database (GFED version 2), representing mainly forest and savannah burning (van der Werf et al., 2006). Since this dataset is available only after 1997, an average seasonal cycle is used for the 1988–1996 period. Unlike the wetland emissions, global total IAV BB emissions are lower than that those incorporated in CYC NAT. Thus the CYC BB emission is only partially replaced by IAV BB (see Table 1). This methodology is likely to double count some of the open burning and to underestimate the emissions from closed burning.





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7. Inversion-derived emissions (IAV INV) are obtained by optimizing surface fluxes to reproduce the measured $CH_4$ concentrations using the LMDZ model for the period of 1988–2005 (Bousquet et al., 2006). An average seasonal cycle is repeated for 2006–2008.

8. The soil sink represents a climatological average year, accounting for seasonality, derived from the LMDZ atmospheric $CH_4$ inversion (Bousquet et al., 2006). The global total removal amounts to 27.21 Tg $CH_4$ $yr^{-1}$.

The integrated $CH_4$ emissions for the different combinations of emission fields (tracer 1 to 6 in Table 1) agree within 3 Tg $CH_4$ over 1990–2005 (8675 Tg $CH_4$ for $CH_4$_CTL, which is 542 Tg $yr^{-1}$ on average). Only the IAV INV tracer total emissions are slightly lower (8641 Tg $CH_4$).

Three other tracers ($SF_6$, $^{222}$Radon and $CH_3CCl_3$) are simulated using the following fluxes:

1. Annual mean $SF_6$ emission distributions at $1 \times 1^\circ$ are taken from EDGAR 4.0 (2009) for the period 1988–2005, and the global totals are scaled to Levin et al. (2010). The 2005 distribution is used from 2006 onwards. $SF_6$ emissions increased from 4.77 Tg $yr^{-1}$ in 1990 to 6.79 Tg $yr^{-1}$ in 2007.

2. Radon emissions are constructed based on the surface type in each model grid-cell; 0 poleward of 70°, $8.23 \times 10^{-23}$ mol $m^{-2}$ $s^{-1}$ for 60–70°, and $1.66 \times 10^{-20}$ and $8.30 \times 10^{-23}$ mol $m^{-2}$ $s^{-1}$ for land and ocean grids, respectively, within 60° S–60° N (Jacob et al., 1997). Radon emission fields were not to be rescaled to match a global total source, but are expected to produce a global radon source of approximately $2.2 \times 10^{-6}$ mol $s^{-1}$.

3. The annual mean $CH_3CCl_3$ emission distribution is taken from EDGAR3.2 and linearly corrected for the global totals following McCulloch and Midgley (2001) for the period 1988–1998. Emissions for 1999 to 2002 are taken as 27.5, 26.0, 17.7,





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and 16.1 Gg $yr^{-1}$, respectively. After 2002, the regional emission trends follow an exponential decay with a timescale of 5 yr (Krol et al., 2003; updated).

### 2.3 Participating models and output

Twelve chemistry-transport models and four of their variants (2 at higher horizontal resolution and 2 using different OH, Cl and O($^1$D) fields) have submitted simulation results for the period 1990–2007 (Table 2). Half of these models (ACTM, CCAM, IMPACT, LMDZ, PCTM, TM5) also participated in the previous TransCom continuous experiment, where they were tested for interhemispheric transport using $SF_6$, and synoptic and diurnal scale variability using continuous $CO_2$ measurements at surface sites (Law et al., 2008; Patra et al., 2008). Six other models (ACCESS, CAM, GEOS-Chem, MOZART, NIES08i, TOMCAT) participated for the first time in a TransCom experiment. The model horizontal resolution variedfrom 1 × 1° longitude × latitude to 6 × 4°. In the vertical, 19 to 67 levels were employed. Salient features of each model configurations (resolutions, meteorological fields) are given in Table 2 for guidance purpose only, and do not automatically linked with model performance as evaluated for various features in this study.

Concerning the wind field and other meteorology, all models, except ACCESS, used meteorological fields from weather forecast models either by interpolation (offline models) or by nudging towards horizontal winds (U, V) and temperature (online models). Most models generated output as 1-hourly averages, except LMDZ and MOZART, which provided output as 3-hourly averages.

Details of individual transport models can be found in the following references; ACCESS (Corbin and Law, 2011), ACTM and ACTM_OH (Patra et al., 2009a,b), CAM (Gent et al., 2009), CCAM (Law et al., 2006), GEOS-Chem and GEOS-Chem_DOH (Pickett-Heaps et al., 2011; Fraser et al., 2011), IMPACT and IMPACT_1 × 1.25 (Rotman et al., 2004), LMDZ (version 4; Hourdin et al., 2006), MOZART (version 4;





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Emmons et al., 2010), NIES08i (Belikov et al., 2011), PCTM (Kawa et al., 2004), TM5 and TM5_1 × 1 (Krol et al., 2005), TOMCAT (Chipperfield, 2006).

The TransCom-$CH_4$ experiment archived model simulations for 18 yr and 9 tracers, sampled at 280 surface sites and 115 vertical profile sites (at all model levels within troposphere) at hourly time intervals as well as 3-D output at 17 standard pressure levels for monthly-means for 1990–2007, and noon-time daily values for 2001–2007.

### 2.4 Observational data sources and processing

Selected sites from the Advanced Global Atmospheric Gases Experiment (AGAGE; http://agage.eas.gatech.edu) and the NOAA Earth System Research Laboratory, Global Monitoring Division (http://www.esrl.noaa.gov/gmd) networks are used in this study. These sites all have simultaneous measurements of $CH_4$, $SF_6$, and $CH_3CCl_3$ covering the 1990s and 2000s (Table 4). Unfortunately, radon measurements are not available for most of these sites. Monthly or annual mean observations have been calculated from continuous (hourly averages) or flask sampling (events) measurements data available from the World Data Center for Greenhouse Gases website (WDCGG, 2010). NOAA flasks are usually sampled under clean air (or baseline) conditions; this is usually onshore flow at coastal sites. The AGAGE continuous records have been flagged to remove local and regional pollution events.

The model outputs are extracted for the corresponding sites and sampling time from the hourly surface data files. For the ACCESS and CCAM models, we have chosen the BRWOCN and CGOOCN sites, the nearest ocean grid to the site, to better represent baseline conditions. The results of HBA (75.6° S, 26.5° W, 10 m) site are used as a replacement for the SPO site for PCTM. These selections are made as per the modeler's advice.

For $CH_4$, the NOAA4 calibration scale (Dlugokencky et al., 2005) agrees within 5 ppb with the AGAGE/Tohoku University calibration scale (Aoki et al., 1990). The calibration scales for $CH_3CCl_3$ are given in Prinn et al. (2005) for AGAGE and NOAA2003 (http://www.esrl.noaa.gov/gmd/ccl/scales/CH3CCl3_scale.html) for NOAA sites. Those





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for $SF_6$ are based on Rigby et al. (2010) and NOAA6 (http://www.esrl.noaa.gov/gmd/ccl/sf6_scale.html). The AGAGE and NOAA scales for $SF_6$ are in excellent agreement (NOAA–AGAGE = 0.02 ± 0.01 ppt, compared to repeatability of the analytical system of 0.04 ppt) (Rigby et al., 2010). The systematic NOAA–AGAGE difference of ~ 4 ppt in 1992 reduces linearly to around 0 ppt in 2001 for the $CH_3CCl_3$ concentrations as determined from co-samples measurements at CGO, SMO and MHD by both the networks (Paul Krummel and Tim Arnold, personal communication, 2010).

As a check for the stratospheric $CH_4$ distribution simulated by the models, a climatology of $CH_4$ vertical profiles measured by the ACE-FTS (Atmospheric Chemistry Experiment - Fourier Transform Spectrometer) instrument onboard the SCISAT-1 satellite in the upper troposphere and stratosphere altitudes has been used (De Mazière et al., 2008). Please note that the data coverage of the ACE-FTS instrument in the tropical region is sparse. The HALOE/UARS (Halogen Occultation Experiment onboard the Upper Atmosphere Research Satellite) (Park et al., 1996) had a denser coverage and measurements of stratospheric $CH_4$ from this are also used for validating simulated vertical gradients in the tropical stratosphere (100–10 mb).

For this analysis, the models were sampled within 1 to 3 h of the sampling times of the measurements. Time series were constructed of monthly or annual mean samples for verification of model simulated seasonal cycles and interannual variability, respectively. The Pearson's moment correlation analysis is performed to evaluate the agreements between the model and observed time series for seasonal cycles and interannual variations at the 8 selected sites.

### 2.5 Calculation of IH gradients, IH exchange rate, and atmospheric lifetimes

The IH exchange rate ($\tau_{ex}$) is estimated from the $SF_6$ annual mean concentration time series and the ratio of emission in the NH ($E_n$) and SH ($E_s$) using (Patra et al., 2009b and references therein):





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$$\tau_{ex} = \left[\Delta c_{n-s}\left(\frac{E_n}{E_s}+1\right)\right] / \left[\frac{E_n}{E_s}\frac{dc_s}{dt} - \frac{dc_n}{dt}\right] \quad (4)$$

$c_s$ and $c_n$ are the average concentrations of SH and NH sites, and $\Delta c_{n-s}$ is IH concentration gradient. We used two sites in the NH (BRW, MLO) and SH (CGO, SPO) to estimate the hemispherical average concentrations $c_n$ and $c_s$, respectively, at yearly time intervals ($dt$). Because $SF_6$ measurements at ALT are not available after 2005, BRW is chosen for this analysis. The values $c_n - c_s$ are shown as IH gradients of each species.

We also estimated $CH_4$ and $CH_3CCl_3$ lifetimes ($\tau$) using the mass balance equation

$$\frac{dB}{dt} = E - L = E - \frac{B}{\tau} \quad (5)$$

$B$, $E$ and $L$ are the annual total atmospheric burden, surface emission and photochemical loss, respectively. As approximate estimates of $B$, average concentrations of $CH_4$ and $CH_3CCl_3$ for 8 sites are multiplied by the concentration-to-mass conversion factors of 2.845 Tg $CH_4$ ppb$^{-1}$ and 23.689 Gg $CH_3CCl_3$ ppb$^{-1}$, respectively, (ideally vertical distribution properties of each species should be accounted for with appropriate air mass factors), for this calculation. However, we note that the average lifetimes over the period of 2000–2007 calculated using ACTM simulated gridded loss rates agree within 0.01 yr for $CH_3CCl_3$ with the lifetimes calculated using Eq. (5) suggesting that the simplification is acceptable.

## 3 Results and discussions

### 3.1 Zonal mean concentrations

Figure 3 compares the latitude-pressure variations of the zonal mean $CH_4$_CTL tracer in the troposphere and lower stratosphere to representative ACE-FTS measurements (please refer to Figs. S1–S17 for individual model comparison plots of $^{222}Rn$, $SF_6$ and





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$CH_4$ along the 70° E and 180° E longitudes, and zonal mean $CH_3CCl_3$ distributions corresponding to the years 1994 and 2005). Generally, all models exhibit similar large-scale features, equator-pole latitudinal gradients, and vertical gradients in the lower stratosphere. Most significantly different, however, is the transition between the troposphere and stratosphere and the heights at which the vertical gradients maximize in the models (the green-to-blue shaded regions). The ACCESS/GEOS-Chem/NIES-08i and CAM/MOZART/CCAM models show the maximum and minimum decrease rates of $CH_4$ with increasing height, respectively, in the height range of 100–50 mb. CAM, CCAM and MOZART models have only few vertical layers above 100 mb, and the NIES model employs isentropic coordinate system in the stratosphere. The formulation of models is known to affect the simulation of tracer gradients in the upper troposphere/lower stratosphere region. A version of the TOMCAT/SLIMCAT model, which uses isentropic coordinates in the stratosphere, produces stronger tracer gradients and a more realistic Brewer Dobson circulation than the p-coordinate version used here (e.g., Hossaini et al., 2010 and references therein). The "tropical pipe" (Plumb, 1996) along the upward transport branch of the Brewer-Dobson circulation appears "leaky" in the models than in the limb-viewing remote sensing observations by HALOE and ACE-FTS. As a result, the simulated concentration isopleths appear flatter compared to the observations with increasing latitudes in both hemispheres. The $CH_4$ meridional gradients between tropics (Eq–10° N) and northern extratropics (20° N–30° N) are calculated to be 140 ± 72 ppb for the models and 225 ppb for HALOE in the height range of 70–30 mb.

In the troposphere, vertical transport of the NH emission varies between the models, most prominently in the tropical region, where deep cumulus convection is prevalent. Table 3 shows mean vertical gradients and between model variabilities in three broad latitude ranges. Model to model differences are much less distinct at the mid and high latitudes than in the tropics for $SF_6$, $CH_4$ and $CH_3CCl_3$. However, two main categories of models can be identified based on the density of $CH_4$ isopleths in the height range between 350 and 200 mb. The position and concentration gradient across the





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tropopause differs considerably among models. These model features are also present in the $SF_6$ simulations suggesting the predominant role of transport in the simulation of the $CH_4$ vertical distributions. The penetrative mass flux due to deep cumulus convection in tropical latitudes is strongest in ACTM and GEOS-Chem (low $^{222}$Rn concentration difference of $\sim 5.63 \times 10^{-21}$ between 850 and 200 mb), and relatively weaker in NIES08i, PCTM and TOMCAT (high $^{222}$Rn concentration difference $> 12 \times 10^{-21}$). This is clearer from the simulated $^{222}$Rn distributions over the South Asian monsoon region at 70° E during boreal summer (Figs. S2–S5). It has been shown in Patra et al. (2009b), based on ACTM simulations with and without cumulus parameterization, that deep cumulus convective transport is the main cause for rapid vertical transport of tracers to the upper troposphere (seen as higher $^{222}$Rn concentrations compared to the lower troposphere). The higher horizontal resolution versions of both IMPACT and TM5 resulted in higher $^{222}$Rn concentrations in the middle-upper troposphere (i.e., smaller difference between 850 mb and 400/200 mb) compared to their respective lower resolution simulations. This is suggesting that some convective processes are being resolved in higher resolution models that are not present in the lower resolution models.

### 3.2 Model-observation comparison of $CH_4$, $SF_6$ and $CH_3CCl_3$ Annual means: 1990–2007

Figure 4 shows the time series of annual mean concentration differences between simulated and observed $CH_4$, $SF_6$ and $CH_3CCl_3$ at two selected sites (MLO and CGO). First $SF_6$ is considered, which has no chemical loss (middle row). Typical model behaviour is similar at all 8 sites. For most models, the simulated concentrations divert from the measurements by 0.2 ppt in 1995, after which differences remain at that level. The offsets between models can be explained by initial values assumed by each models. Only ACCESS shows increasing differences in time until 2006, and ACTM_OH, which uses EDGAR4.0 emissions without scaling between 1988–2005, and 2005 emission for 2006 and 2007 and produces a slower increase in the model concentration compared to observations after 2000. This suggests that the global total emissions





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estimated by Levin et al. (2010) and later confirmed by Rigby et al. (2010) are adequate also for independent state-of-the-art transport models.

The $CH_3CCl_3$ simulations are of an intermediate level of uncertainty, as this species is emitted to the atmosphere by a relatively well quantified industrial use as a solvent. However, uncertainties remain for its loss by reaction with OH in the troposphere, photolysis in the stratosphere and the rate of STE. Until 1990, $CH_3CCl_3$ emissions were increasing, followed by near-exponential decrease due to stringent restriction of its production/use by the Montreal protocol (WMO/SAOD, 2003). Because the lifetimes of $CH_3CCl_3$ due to photochemical removal is much longer in the stratosphere ($\sim$ 28.6 yr) than in the troposphere ($\sim$ 5.8 yr) (estimates from ACTM simulated loss rates at model grids), the troposphere to stratosphere transport plays a minor role in the global total budget of $CH_3CCl_3$ after the late 1990s because the concentration gradients across the tropopause reduced to less than 10 ppt (ref. Table 3). These lifetimes due to stratosphere and tropospheric losses are although within the range of independent estimates, $38^{+15}_{-11}$ and $6^{+0.5}_{-0.4}$, respectively, both the mean values are lower for ACTM (Prinn et al., 2005). Despite the fact that the models were initialized using the same concentration at 1 January 1988, significant differences in the simulated concentrations are found already for 1990, after two years of simulation. The established model-measurement differences in 1990 persist until the end of the simulations (LMDZ MCF being the only outlier) even though the $CH_3CCl_3$ concentrations become very small towards 2007. Given the $CH_3CCl_3$ lifetime of less than 5 yr, these differences are caused by differences in transport and removal rather than the initialization.

$CH_4$ is the most complicated species considered in the TransCom-$CH_4$ experiment, because the $CH_4$ surface emissions and STE are less certain than for $CH_3CCl_3$. Generally, the models that simulate lower $CH_3CCl_3$ concentrations compared to the multi-model mean, also yield lower $CH_4$ concentrations (such as MOZART, CCAM). For example, the high and low-resolution TM5 simulations show the highest concentrations of both $CH_4$ and $CH_3CCl_3$ at MLO. However, it is interesting to note that GEOS-Chem (with TransCom OH) calculates the lowest $CH_4$ concentrations among all models, while





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the simulated $CH_3CCl_3$ levels are not distinctly different. LMDZ is among the models that most strongly underestimate the observed level of $CH_3CCl_3$, whereas the opposite is true for $CH_4$ (Fig. 4). These contrasting behaviors among models for various $CH_4$ and $CH_3CCl_3$ simulations clearly suggest that $CH_4$ loss due to the reaction with OH in the troposphere is not the only control on the $CH_4$ budget differences between models and that other factors such as transport differences also play a role.

### 3.2.1 Seasonal cycles

Figure 5 shows model to measurement comparisons of the seasonal cycles of $CH_4$, $SF_6$ and $CH_3CCl_3$ at three selected sites (MLO, SMO and CGO) for the period 2002–2003. These sites have been selected because of their location at large distance of the continental emissions for each of these species. To highlight differences in seasonality, approximate linear trends and offsets corresponding to the period 2002–2003 have been subtracted from the monthly-mean values. All models capture the salient features in the seasonal cycles at very high statistical significance (ref. Table 4 for correlation coefficients at 8 sites), except for $SF_6$ at MLO, where the measurements show unusual fluctuations and a large data gap during 2002. Even for the years with dense data coverage (2005–2006), low average correlation coefficients ($r$) of 0.3 for the $SF_6$ seasonal cycles at MLO are obtained due to a very small seasonal cycle of less than 0.04 ppt. In contrast, the clear seasonalities (amplitude $> 0.04$ ppt) at ALT, MHD, SMO, CGO and SPO are well reproduced by the models ($r > 0.4$; significant at $P = 0.05$ in two tailed Student's $t$-test for 24 data points). The fact that the models are able to reproduce the observed seasonal cycles indicates that even though the signals are weak, they nevertheless provide useful information for model validation. All models reproduce the $CH_3CCl_3$ seasonal cycle fairly well at all 8 sites, both in phase and amplitude ($r > 0.8$).

For $CH_4$, the influence of the surface flux on the simulated seasonal cycles can be studied using the 6 different $CH_4$ tracers. The corresponding correlation coefficients, listed in Table 4, suggest that the $CH_4$ seasonal cycle depends strongly on the implemented wetland and biomass burning fluxes. The $CH_4$_INV simulations consistently





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produce higher correlation coefficients at all the NH sites (ALT, BRW, MHD, MLO and RPB), which was expected because the atmospheric-$CH_4$ inversion used data from these sites for flux optimization. $CH_4$_EXTRA results are next best in comparison with measured seasonal cycles for the NH high latitude sites. For any given tracer, the correlation coefficients are highest at remote SH sites, CGO and SPO, compared to all other sites. The use of ACTM_OH and GEOS-Chem_DOH or the higher horizontal resolution versions of IMPACT_1 × 1.25 and TM5_1 × 1 do not always improve the agreement between model and observations compared to the default implementation. These results suggest a need for improving our understanding of the $CH_4$ flux seasonality in the Northern Hemisphere land regions, noting that the OH loss is realistically represented as seen in the simulated $CH_3CCl_3$ seasonal cycles. Because ACCESS (blue line) was not run with analyzed winds and temperature, the simulated seasonal cycles are not as good as other models for $CH_4$ and $SF_6$ highlighting the role of meteorology in simulating tracer concentrations. The role of meteorology is less pronounced for $CH_3CCl_3$ because the emissions are weak in the 2000s. A more detailed analysis of seasonality at a larger number of sites will be conducted in a future study.

### 3.2.2 Interannual variability (IAV)

We calculated growth rates for all tracers as the difference between annual mean concentrations for two adjacent years. The growth rate at January 2000 is shown as the difference between 1999 and 2000 mean concentrations. The simulated and observed $SF_6$ growth rates (not shown) decreased from $\sim 0.25$ ppt yr$^{-1}$ in 1997 to $\sim 0.2$ ppt yr$^{-1}$ in 2000. Afterwards, the growth rate steadily increased to $\sim 0.25$ ppt yr$^{-1}$ in 2006 (please refer to Table 4 for correlation coefficients; $r \sim 0.7$ for 5 sites). The length of time series considered for the correlation calculation of the IAVs is 1990–2007 for $CH_4$ (except for $CH_4$_WL_BB, which has 7 yr of IAV, 1994–2000) and $CH_3CCl_3$, and 1996–2007 for $SF_6$ (the period when observations are available). The correlation coefficients greater than 0.44 and 0.53 are statistically significant at $P = 0.05$ for 18 and 12 data points, respectively. Average $CH_3CCl_3$ growth rates (not shown) hovered around 0 to 5, −12 to −17





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and $\sim -2.5\,\mathrm{ppt\,yr^{-1}}$ during 1991, 1997 and 2007, respectively, with gradual changes in between. These temporal variations are well simulated by all models ($r > 0.9$). Because both $SF_6$ and $CH_3CCl_3$ emissions are of purely anthropogenic origin, their production, consumption and release to the atmosphere vary relatively smoothly in comparison to the natural components of $CH_4$ emissions.

Figure 6 shows the model-observation comparison of the IAV in $CH_4$ growth rates for three broad latitude regions: the NH, tropics and SH corresponding to the CTL and EXTRA emissions (refer to Figs. S20 and S21 for others). Averaged observed $CH_4$ growth rates for the 1990s are 5.25, 5.06 and 7.01 $\mathrm{ppb\,yr^{-1}}$ in the NH, tropics and SH, respectively. Almost no increase in concentration is observed for the 2000s (except for 2007). Additionally, it can be seen that the IAVs in the growth rate are higher at the NH sites compared to the tropical and SH sites. Figure 6 shows that although the $CH_4$_CTL simulations capture the observed reduction in the decadal average growth rates, the IAV is not well reproduced. Most prominent is the 1997/1998 El Niño event (Langenfelds et al., 2002). During this event the observations show an increase, while the simulations show a decrease in the growth rate. Interestingly, the $CH_4$_CTL emission and OH concentration (both without IAV) cannot explain this model behavior, which is therefore attributed to the increased $CH_4 + OH$ reaction rate as modified by $CH_4$ transport and temperature in the model. This, in turn, is caused by the El Niño induced higher air temperatures (Reaction R1), resulting in faster removal of $CH_4$ from the troposphere and thus a decrease in the growth rate. Indeed, the 1998 $CH_4$_CTL lifetime (9.82 yr) is estimated to be the shortest among all simulation years.

As seen from Table 4, inclusion of biomass burning emission IAV ($CH_4$_BB) improves the IAV model-observation agreement at all sites compared to $CH_4$_CTL. However, when wetland emissions are included ($CH_4$_WL_BB), the correlations tend to deteriorate. Compared to $CH_4$_BB, only $CH_4$_EXTRA produces better model-observation agreement for growth rates (Fig. 6, right panels). The wetland $CH_4$ emission simulated by the VISIT ecosystem model included in $CH_4$_EXTRA displays a large positive anomaly on top of the emissions from biomass burning during 1997/1998. Combined,





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these emissions compensate for the extra $CH_4$ loss due to higher air temperatures. As a consequence, these emissions result in an excellent agreement with the growth rates observed at the SH and NH sites (except that the simulated tropical signal is a bit too strong in 1998). The IAV of the multi-model average $CH_4$ growth rate at BRW did not correlate significantly with the observed IAV because this site is located close to the Alaskan wetland region, and the site representation error is large for the coastal sites in coarse resolution global models (Patra et al., 2008).

The decreasing growth rate in the 1990s, near zero growth rates in early 2000s and the reappearance of positive $CH_4$ growth in the late 2000s have drawn considerable interests for developing emission inventories. For example, Lamarque et al. (2010) suggested a decrease of $CH_4$ emissions by about 40 Tg $CH_4$ from 1990 to 2000 for simulating the zero $CH_4$ growth rate in the early 2000s using the CAM-Chem model. Their estimate is largely inconsistent with our results, which is suggesting that a steady state is achieved between $CH_4$ chemical destruction and emissions during the early 2000s (Dlugokencky et al., 2003). The EDGAR4.0 anthropogenic $CH_4$ emission increase of $\sim 4$ Tg $CH_4$ $yr^{-1}$ during 2001–2007, synchronized with the Chinese economic growth, produces inconsistencies between observed and simulated growth rates during 2003–2007 (thus the lowest correlation coefficients for the growth rate IAVs in Table 4; see also Fig. S20). This indicates that forward simulations using multiple forward transport models are useful for the verification of emission inventories.

### 3.3 Interhemispheric gradients and exchange rates

Figure 7 shows the concentration gradients between two NH sites and two SH sites obtained using annual mean observed and modeled time series. All models except for the TM5s, simulate the observed $SF_6$ IH gradient within the measurement accuracy of $\pm 0.057$ ppt ($\sqrt{2} \times$ measurement precision). These gradients translate to an average IH exchange time ($\tau_{ex}$) of $1.39 \pm 0.18$ yr (for all models and years), which is an indication of close model-model agreement Fig. 8. This model-model spread is much smaller compared to the model pool of the 1990s (Denning et al., 1999), which gave





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$\tau_{ex}$ range of 0.8–2.0 yr. In this intercomparison, the $\tau_{ex}$ differences between models is 0.62 ± 0.06 yr compared 1.2 yr in the TransCom experiment during the 1990s. The average $\tau_{ex}$ of 1.39 yr is in excellent agreement with the observation-based estimates of 1.3 yr (Geller et al., 1997) and 1.5 yr (Levin and Hesshaimer, 1996). The underestimation by ACTM_OH version is due to smaller $SF_6$ emissions (note: this version used EDGAR4.0 without scaling), highlighting the role of the emission strength in the forward model simulations. A tendency towards faster IH exchange rates is seen for both the observations and the simulations (a decrease in exchange time by about 0.2 and 0.15 yr, respectively) between 1996–1999 and 2004–2007. This behavior might be linked to the widening of the tropical belt during the last three decades (Seidel et al., 2008), and the associated weakening of the tropical/subtropical mixing barrier for long-lived tracer transport between the two hemispheres (e.g., Miyazaki et al., 2008).

All three long-lived species show a similar relationship for the IH gradient: the model that produces a larger (smaller) $SF_6$ IH gradient generally also produces a larger (smaller) IH gradients for $CH_3CCl_3$ and $CH_4$ in comparison with the observations (Fig. 9). The intriguing exception is MOZART, which exhibits an excellent match for $SF_6$ IH gradient, but produces one of the largest $CH_4$ IH gradients (127 ppb compared to an observed value of 101 ppb) and one of the smallest $CH_3CCl_3$ gradients (0.13 ppt) during 2003–2007. Similar contrasting behaviour is also seen for several other models at lesser distinction, e.g., GEOS-Chem_DOH, NIES-08i lie above the fitted line for $CH_4$, but lie below the fitted line for $CH_3CCl_3$. The $CH_4$ IH gradients are best reproduced using the $CH_4$_INV emissions: deviations are within 5 ppb for 7 models (Fig. S17). Taking into account the IH gradient of all three species, TM5/CCAM, ACTM/GEOS-Chem/IMPACT/PCTM and LMDZ/NIES-08 showed systematically higher, similar and lower IH gradients, respectively, compared to the observations.





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### 3.4 Photochemical removal of $CH_4$ and the role of transport

The calculated photochemical loss of $CH_4$ varies between 490 and 509 Tg $CH_4$ $yr^{-1}$ during the first eight years (1992–1999), and between 497 and 513 Tg $CH_4$ $yr^{-1}$ during the last eight years (2000–2007) of the simulation. Figure 10a suggests that the eight-years averaged growth rates at the surface sites for the different models are, as expected, inversely proportional to the calculated photochemical destruction. However, this relationship appears loose, particularly for 2000–2007, when the models approach steady state (correlation coefficient, $r = -0.42$ for all models, but increase to $-0.82$ and $-0.61$ for 1990s and 2000s, respectively, by excluding NIES-08i). Possible explanations are investigated in Fig. 10b, using the vertical gradients in the equatorial lower stratosphere ($CH_4$ at 100 mb – $CH_4$ at 10 mb; zonal average for 5° S–5° N latitudes). Models showing greater gradients have slower Brewer-Dobson circulation, and thus stronger trapping of $CH_4$ in the troposphere, resulting in faster $CH_4$ destruction and smaller growth rate because $CH_4$ lifetime in the troposphere is an order of magnitude shorter than that in the stratosphere. Although these relationships are again quite loose during the 1990s, but as the models attain their steady state, statistically significant correlations ($r = -0.69$) are found for the period of 2000–2007 between the $CH_4$ growth rates and vertical gradients in the lower stratosphere. While the growth rates decrease from the 1992–1999 period to the 2000–2007 period in all models, the modeled gradients remain remarkably constant. Based on this analysis we suggest that the simulated concentration growth rates at the surface sites is linked to the troposphere to stratosphere transport rate of $CH_4$.

Figure 11 shows the temporal variability in the estimated lifetimes (Eq. 5) for $CH_4$_CTL and $CH_3CCl_3$ (ref. Table 2 for time-averaged model specific lifetimes). The median $CH_4$ lifetime due to atmospheric loss processes (Reactions R1–R3) is $9.99 \pm 0.08$ ($1\sigma$ for interannual variability) years and ranges from $9.50 \pm 0.10$ and $10.27 \pm 0.14$ yr for the different models. Here it should be remembered that all, but ACTM_OH and GEOS-Chem_DOH, models used the same OH distribution. The





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median $CH_4$ lifetime for 16 models or model variants agrees well with the lifetime (10.0 ± 0.17) estimated from the measured mean concentrations at 8-sites. This is because a close balance is achieved between the modeled atmospheric loss and net surface emissions (513 ± 9 and 514 ± 14 Tg $CH_4$ $yr^{-1}$ for the 1990s and 2000s, respectively). The median $CH_3CCl_3$ lifetime due to all loss processes (Reactions R4–R6) is estimated to be 4.61 ± 0.13, 4.59 ± 0.18 and 4.62 ± 0.02 during the 1992–2007, 1992–1999 and 2000–2007, respectively. TM5 simulates a lifetime of 4.92 ± 0.50 yr, which is close to the estimates using the measured $CH_3CCl_3$ concentrations of 4.9 ± 0.3 yr (Prinn et al., 2005), 5.0 yr (WMO/SAOD, 2003) or 4.94 ± 0.15 yr (this study based on 8 sites). All other models calculated a shorter lifetime, by an average value of 0.3 yr. CAM, CCAM, LMDZ and MOZART calculate lifetimes of 4 yr or shorter. The interannual variation in the estimated $CH_3CCl_3$ lifetimes during the 1990s is an order of magnitude higher than that in the 2000s. We find up to 5 % variability in the modeled $CH_3CCl_3$ lifetimes during 1992–1999, a period with substantial emissions. During 2000–2007, when the emissions of $CH_3CCl_3$ dropped significantly, less than 0.4 % variability is simulated. This implies that inversions to estimate OH from $CH_3CCl_3$ observations are less uncertain since 2000, a finding in good agreement with Montzka et al. (2011).

## 4 Further work and data accessibility

For this analysis we used results of chemical tracer simulations at only 8 selected sites, which is a very small subset of the 280 surface sites for which output is available. In addition vertical profiles of chemical tracers and several meteorological parameters have been archived at 115 sites. More analyses on the basis of the TransCom-$CH_4$ simulations are planned focusing on (1) $CH_4$ vertical profiles measured using aircraft, (2) analysis of vertical column averaged $CH_4$ concentrations using TCCON observations, (3) using increase, decrease and exponential decay of $CH_3CCl_3$ for optimizing tropospheric OH abundance. We also welcome use of this data set by the measurement community. In an effort towards ease of access, time series at a subset





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of surface sites are archived at the WMO World Data Centre for Greenhouse Gases (http://gaw.kishou.go.jp/). Information on how to access the full dataset is available on this website (http://transcom.project.asu.edu/T4_methane.php).

In addition to the site-specific data, gridded output at monthly intervals at the model horizontal resolution at standard pressure levels are archived for the period 1990–2007. Afternoon averages (12:00–15:00 LT – Local Time) at daily intervals are also archived for the period 2001–2007. We believe these sets of model output and their extension to recent years will be useful for comparing the model simulations with satellite observations (SCIAMACHY, AIRS, GOSAT) and aircraft observations (e.g., HIPPO – HIAPER Pole-to-Pole Observations of Carbon Cycle and Greenhouse Gases Study, Wofsy et al., 2011).

## 5 Summary and conclusions

We analyzed concentration time series of $CH_4$, $CH_3CCl_3$, $SF_6$ and $^{222}Rn$ simulated by 16 chemistry-transport models as part of the TransCom-$CH_4$ intercomparison experiment. We focused the analysis on the model-to-model differences in

1. The vertical redistribution of tracers, based on $^{222}Rn$ simulations and comparisons to $CH_4$ satellite observations in the upper troposphere and lower stratosphere.
2. Large-scale interhemispheric (IH) transport, by comparing modeled and observed IH gradients of $SF_6$, $CH_4$ and $CH_3CCl_3$.
3. Simulated seasonal cycles, by comparing to observed seasonal cycles at remote background stations.
4. Inter-annual variations in the simulated $CH_4$ growth rate, by focusing on the results of six different $CH_4$ emission time-lines.





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5. The role of removal by OH on the simulated $CH_4$ and $CH_3CCl_3$ concentrations.

The main conclusions can be summarized as follows:

i. Although the simulated zonal mean $^{222}Rn$ concentrations agree between models, significant differences are observed in regions of deep cumulus convection, e.g., the south Asian summer monsoon domain. Unfortunately, observational evidence to check the model behavior is lacking. Models also differ in the simulated height of large troposphere-stratosphere concentration gradients of $CH_4$. Compared to $CH_4$ satellite observations in the upper troposphere, most models appear to be too diffusive around the tropical tropopause.

ii. The IH exchange time, calculated from the simulated $SF_6$ distributions, ranges from 1.79 to 1.17 yr (average over 1996–2007) for the different models. The model-average value of 1.39 yr is in close agreement with earlier studies and observational evidence. Models that show faster IH exchange for $SF_6$, also exhibit faster exchange (smaller IH gradients) for $CH_4$ and $CH_3CCl_3$. This multi-tracer evidence provides clear directions for the improvement of specific models. Both the exchange rate calculated from the models and the rate calculated from the observations suggest an acceleration of the IH exchange in the 2000s compared to the 1990s.

iii. All models reproduce the observed seasonal cycles of $CH_4$ and $CH_3CCl_3$ at background sites very well. The simulated $CH_4$ seasonal cycles depend on the destruction by reaction with OH, suppressing the seasonality of the underlying emissions. Two of the six simulated $CH_4$ tracers show a higher correlation with the observed seasonal cycle.

iv. A set of six $CH_4$ global flux representations was used to investigate the role of specific processes in reproducing the observed interannual variations in the $CH_4$ growth rate. The control emission case ($CH_4$_CTL) without a significant increase in anthropogenic emissions and no interannual variability in natural emissions





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for the period 1990–2007 reproduces the declining growth rate in the 1990s, followed by the stabilization in the 2000s. Inclusion of interannual variation in emissions from the wetland and forest fires ($CH_4$_EXTRA) most successfully simulates the observed interannual variations in $CH_4$. It was also found that the higher tropospheric temperatures during the 1997/1998 El Niño resulted in larger $CH_4$ destruction, whereas the observations clearly show a rise in $CH_4$ concentration. To match the observations, either enhanced emissions are required (as in $CH_4$_EXTRA), or less than average OH should have been present in this period.

v. The simulation of $CH_3CCl_3$ is used to check the consistency of the employed OH abundance and distribution. Two models used an alternative OH field next to the prescribed field. Generally, models that simulate a low abundance of $CH_3CCl_3$ also simulate a low abundance of $CH_4$. However, there are exceptions, which indicates that $CH_4$ loss due to OH in the troposphere is not the only cause of the modeled $CH_4$ differences. Thus, horizontal and vertical transport differences may also be important.

Further analysis reveals that the simulated $CH_4$ growth rate shows (weak) correlations with the modeled vertical gradient in the equatorial lower stratosphere. This suggests that differences in vertical mixing of the emissions and in stratosphere-troposphere exchange are the main causes of the model-to-model differences. Next to the inter-hemispheric transport in models, this issue requires further analysis, e.g. based on the archived model output.

Finally, the multi-model lifetime estimates for $CH_4$ and $CH_3CCl_3$ were found to be fairly constant over the simulation period with median values of $9.99 \pm 0.08$ and $4.61 \pm 0.13$ for the period 1992 to 2007. This underscores the fact that OH (assumed constant in the simulations) is the driving factor in the budgets of these gases, and that transport and temperature (affecting the reaction rate) differences play a smaller role. We find net $CH_4$ emissions (soil sink subtracted) of $513 \pm 9$ and $514 \pm 14$ Tg $CH_4$ $yr^{-1}$ for the 1990s and 2000s, respectively, to the atmosphere are consistent with the





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atmospheric losses accounted for due to OH, $O^1D$ and Cl in order to simulate $CH_4$ concentrations and growth rates at the surface sites.

**Supplementary material related to this article is available online at: http://www.atmos-chem-phys-discuss.net/11/18767/2011/acpd-11-18767-2011-supplement.pdf.**

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## References


Aoki, S., Nakazawa, T., Murayama, S., and Kawaguchi S.: Measurements of atmospheric methane at the Japanese Antarctic Station, Syowa, Tellus B, 44, 273–281, 1992.

Bergamaschi, P., Frankenberg, C., Meirink, J. F., Krol, M., Villani, M. G., Houweling, S., Dentener, F., Dlugokencky, E. J., Miller, J. B., Gatti, L. V., Engel, A., and Levin, I.: Inverse modeling of global and regional $CH_4$ emissions using SCIAMACHY satellite retrievals, J. Geophys. Res., 114, D22301, doi:10.1029/2009JD012287, 2009.

Belikov, D., Maksyutov, S., Miyasaka, T., Saeki, T., Zhuravlev, R., and Kiryushov, B.: Mass-conserving tracer transport modelling on a reduced latitude-longitude grid with NIES-TM, Geosci. Model Dev., 4, 207–222, doi:10.5194/gmd-4-207-2011, 2011.

Bloom, S., da Silva, A., Dee, D., Bosilovich, M., Chern, J.-D., Pawson, S., Schubert, S., Sienkiewicz, M., Stajner, I., Tan, W.-W., and Wu, M.-L.: Documentation and Validation of the Goddard Earth Observing System (GEOS) Data Assimilation System – Version 4, Technical Report Series on Global Modeling and Data Assimilation 104606, NASA Goddard Space Flight Center, Maryland, 26, 2005.

Bousquet, P., Ciais, P., Miller, J. B., Dlugokencky, E. J., Hauglustaine, D. A., Prigent, C., van der Werf, G. R., Peylin, P., Brunke, E.-G., Carouge, C., Langenfelds, R. L., Lathiére, J., Papa, F., Ramonet, M., Schmidt, M., Steele, L. P., Tyler, S. C., and White, J.: Contribution of anthropogenic and natural sources to atmospheric methane variability, Nature, 443, 439–443, doi:10.1038/nature05132, 2006.

Butler, J. H., Daube, B. C., Dutton, G. S., Elkins, J. W., Hall, B. D., Hurst, D. F., King, D. B., Kline, E. S., Lafleur, B. G., Lind, J., Lovitz, S., Mondeel, D. J., Montzka, S. A., Moore, F. L., Nance, J. D., Neu, J. L., Romashkin, P. A., Scheffer, A., and Snible, W. J.: Halocarbons and other atmospheric trace species, Chapter 5, in: CMDL Summary Report No. 27, 2002–2003, edited by: Thompson, T. M., NOAA/US Department of Commerce, Boulder, USA, 115–135, 2004.

Cao, M., Gregson, K., and Marshall, S.: Global methane emission from wetlands and its sensitivity to climate change, Atmos. Environ., 32, 3293–3299, doi:10.1016/S1352-2310(98)00105-8, 1998.

Chen, Y.-H. and Prinn, R. G.: Estimation of atmospheric methane emissions between 1996–2001 using a 3-D global chemical transport model, J. Geophys. Res., 111, D10307, doi:10.1029/2005JD006058, 2006.






ACPD
11, 18767–18821, 2011



Chipperfield, M. P.: New version of the TOMCAT/SLIMCAT off-line chemical transport model: intercomparison of stratospheric tracer experiments, Q. J. Roy. Meteorol. Soc., 132, 1179–1203, doi:10.1256/qj.05.51, 2006.

Corbin, K. D. and Law, R. M.: Extending atmospheric $CO_2$ and tracer capabilities in ACCESS, CAWCR Technical Report No. 35, The Centre for Australian Weather and Climate Research, ISBN: 978-1-921826-177, Aspendale, 2011.

Cunnold, D. M., Steele, L. P., Fraser, P. J., Simmonds, P. G., Prinn, R. G., Weiss, R. F., Porter, L. W., Langenfelds, R. L., Wang, H. J., Emmons, L., Tie, X. X., and Dlugokencky, E. J.: In situ measurements of atmospheric methane at GAGE/AGAGE sites during 1985–2000 and resulting source inferences, J. Geophys. Res., 107, 4225, doi:10.1029/2001JD001226, 2002.

De Mazière, M., Vigouroux, C., Bernath, P. F., Baron, P., Blumenstock, T., Boone, C., Brogniez, C., Catoire, V., Coffey, M., Duchatelet, P., Griffith, D., Hannigan, J., Kasai, Y., Kramer, I., Jones, N., Mahieu, E., Manney, G. L., Piccolo, C., Randall, C., Robert, C., Senten, C., Strong, K., Taylor, J., Tétard, C., Walker, K. A., and Wood, S.: Validation of ACE-FTS v2.2 methane profiles from the upper troposphere to the lower mesosphere, Atmos. Chem. Phys., 8, 2421–2435, doi:10.5194/acp-8-2421-2008, 2008.

Denning, A. S., Holzer, M., Gurney, K. R., Heimann, M., Law, R. M., Rayner, P. J., Fung, I. Y., Fan, S., Taguchi, S., Friedlingstein, P., Balkanski, Y., Taylor, J., Maiss, M., and Levin, I.: Three-dimensional transport and concentration of $SF_6$: a model intercomparison study (TransCom-2), Tellus B, 51, 266–297, 1999.

Dentener, F., van Weele, M., Krol, M., Houweling, S., and van Velthoven, P.: Trends and inter-annual variability of methane emissions derived from 1979–1993 global CTM simulations, Atmos. Chem. Phys., 3, 73–88, doi:10.5194/acp-3-73-2003, 2003.

Dlugokencky, E. J., Masarie, K. A., Lang, P. M., and Tans, P. P.: Continuing decline in the growth rate of the atmospheric methane burden, Nature, 393(6684), 447–450, 1998.

Dlugokencky, E. J., Houweling, S., Bruhwiler, L., Masarie, K. A., Lang, P. M., Miller, J. B., and Tans, P. P.: Atmospheric methane levels off: temporary pause or a new steady-state?, Geophys. Res. Lett., 30(19), 1992, doi:10.1029/2003GL018126, 2003.

Dlugokencky, E. J., Myers, R. C., Lang, P. M., Masarie, K. A., Crotwell, A. M., Thoning, K. W., Hall, B. D., Elkins, J. W., and Steele, L. P.: Conversion of NOAA atmospheric dry air $CH_4$ mole fractions to a gravimetrically prepared standard scale, J. Geophys. Res., 110, D18306, doi:10.1029/2005JD006035, 2005.





ACPD
11, 18767–18821, 2011



Emmons, L. K., Walters, S., Hess, P. G., Lamarque, J.-F., Pfister, G. G., Fillmore, D., Granier, C., Guenther, A., Kinnison, D., Laepple, T., Orlando, J., Tie, X., Tyndall, G., Wiedinmyer, C., Baughcum, S. L., and Kloster, S.: Description and evaluation of the Model for Ozone and Related chemical Tracers, version 4 (MOZART-4), Geosci. Model Dev., 3, 43–67, doi:10.5194/gmd-3-43-2010, 2010.

Etiope, G. and Milkov, A. V.: A new estimate of global methane flux from onshore and shallow submarine mud volcanoes to the atmosphere, Environ. Geol., 46, 997–1002, doi:10.1007/s00254-004-1085-1, 2004.

Frankenberg, C., Warneke, T., Butz, A., Aben, I., Hase, F., Spietz, P., and Brown, L. R.: Pressure broadening in the $2\nu_3$ band of methane and its implication on atmospheric retrievals, Atmos. Chem. Phys., 8, 5061–5075, doi:10.5194/acp-8-5061-2008, 2008.

Fraser, A., Chan Miller, C., Palmer, P. I., Deutscher, N. M., Jones, N. B., and Griffith, D. W. T.: The Australian methane budget: interpreting surface and train-borne measurements using a chemistry transport model, J. Geophys. Res., submitted, 2011.

Fung, I., John, J., Lerner, J., Matthews, E., Prather, M., Steele, L. P., and Fraser, P. J.: Three-dimensional model synthesis of the global methane cycle, J. Geophys. Res., 96(D7), 13033–13065, 1991.

Geller, L. S., Elkins, J. W., Lobert, J. M., Clarke, A. D., Hurst, D. F., Butler, J. H., and Myers, R. C.: Tropospheric $SF_6$: observed latitudinal distribution and trends, derived emissions and interhemispheric exchange time, Geophys. Res. Lett., 24, 675–678, 1997.

Gent, P. R., Yeager, S. G., Neale, R. B., Levis, S., and Bailey, D. A.: Improvements in a half degree atmosphere/land version of the CCSM, Clim. Dynam., 79, 25–58, doi:10.1007/s00382-009-0614-8, 2009.

Gupta, M., Tyler, S., and Cicerone, R.: Modeling atmospheric $\delta^{13}CH_4$ and the causes of recent changes in atmospheric $CH_4$ amounts, J. Geophys. Res., 101(D17), 22923–22932, doi:10.1029/96JD02386, 1996.

Hein, R., Crutzen, P. J., and Heimann, M.: An inverse modeling approach to investigate the global atmospheric methane cycle, Global Biogeochem. Cy., 11, 43–76, 1997.

Hossaini, R., Chipperfield, M. P., Monge-Sanz, B. M., Richards, N. A. D., Atlas, E., and Blake, D. R.: Bromoform and dibromomethane in the tropics: a 3-D model study of chemistry and transport, Atmos. Chem. Phys., 10, 719–735, doi:10.5194/acp-10-719-2010, 2010.





ACPD
11, 18767–18821, 2011



Hourdin, F., Musat, I., Bony, S., Braconnot, P., Codron, F., Dufresne, J.-L., Fairhead, L., Filiberti, M.-A., Friedlingstein, P., Grandpeix, J.-Y., Krinner, G., Levan, P., and Lott, F.: The LMDZ4 general circulation model: climate performance and sensitivity to parametrized physics with emphasis on tropical convection, Clim. Dynam., 27, 787–813, doi:10.1007/s00382-006-0158-0, 2006.

Houweling, S., Kaminski, T., Dentener, F., Lelieveld, J., and Heimann, M.: Inverse modeling of methane sources and sinks using the adjoint of a global transport model, J. Geophys. Res., 104(D21), 26137–26160, 1999.

Houweling, S., Dentener, F., Lelieveld, J., Walter, B., and Dlugokencky, E.: The modeling of tropospheric methane: how well can point measurements be reproduced by a global model?, J. Geophys. Res., 105(D7), 8981–9002, 2000.

Huijnen, V., Williams, J., van Weele, M., van Noije, T., Krol, M., Dentener, F., Segers, A., Houweling, S., Peters, W., de Laat, J., Boersma, F., Bergamaschi, P., van Velthoven, P., Le Sager, P., Eskes, H., Alkemade, F., Scheele, R., Nédélec, P., and Pätz, H.-W.: The global chemistry transport model TM5: description and evaluation of the tropospheric chemistry version 3.0, Geosci. Model Dev., 3, 445–473, doi:10.5194/gmd-3-445-2010, 2010.

Ito, A.: Changing ecophysiological processes and carbon budget in East Asian ecosystems under near-future changes in climate: implications for long-term monitoring from a process-based model, J. Plant Res., 123, 577–588, 2010.

Jacob, D. J., Prather, M. J., Rasch, P. J., Shia, R.-L., Balkanski, Y. J., Beagley, S. R., Bergmann, D. J., Blackshear, W. T., Brown, M., Chiba, M., Chipperfield, M. P., Grandpré, J. D., Dignon, J. E., Feichter, J., Genthon, C., Grose, W. L., Kashibhatla, P. S., Kohler, I., Kritz, M. A., Law, K., Penner, J. E., Ramonet, M., Reeves, C. E., Rotman, D. A., Stockwell, D. Z., Velthoven, P. F. J. V., Verver, G., Wild, O., Yang, H., and Zimmermann, P.: Evaluation and intercomparison of global atmospheric transport models using $^{222}$Rn and other short-lived tracers, J. Geophys. Res., 102(D5), 5953–5970, 1997.

Kalnay, E., Kanamitsu, M., Kistler, R., Collins, W., Deaven, D., Gandin, L., Iredell, M., Saha, S., White, G., Woollen, J., Zhu, Y., Chelliah, M., Ebisuzaki, W., Higgins, W., Janowiak, J., Mo, K. C., Ropelewski, C., Wang, J., Leetmaa, A., Reynolds, R., Jenne, R., and Joseph, D.: The NCEP/NCAR 40-yr reanalysis project, B. Am. Meteorol. Soc., 77, 437–470, 1996.

Kanakidou, M., Dentener, F. J., Brasseur, G. P., Berntsen, T. K., Collins, W. J., Hauglustaine, D. A., Houweling, S., Isaksen, I. S. A., Krol, M., Lawrence, M. G., Muller, J. F. Poisson, N., Roelofs, G. J., Wang, Y., and Wauben, W. M. F.: 3-D global simulations of





ACPD
11, 18767–18821, 2011



tropospheric CO distributions – results of the GIM/IGAC intercomparison 1997 exercise, Chemosph. Global Change Sci., 1, 263–282, 1999.
Kanamitsu, M., Ebisuzaki, W., Woollen, J., Yang, S. K., Hnilo, J. J., Fiorino, M., and Potter, G. L.:, NCEP-DOE AMIP-II reanalysis (R-2), B. Am. Meteorol. Soc., 83, 1631–1643, 2002.
Kawa, S. R., Erickson III, D. J., Pawson, S., and Zhu, Z.: Global $CO_2$ transport simulations using meteorological data from the NASA data assimilation system, J. Geophys. Res., 109, D18312, doi:10.1029/2004JD004554, 2004.
Krol, M. C., van Leeuwen, P.-J., and Lelieveld, J.: Global OH trend inferred from methylchloroform measurements, J. Geophys. Res., 103, 10697–10711, 1998.
Krol, M. C., Lelieveld, J., Oram, D. E., Sturrock, G. A., Penkett, S. A., Brenninkmeijer, C. A. M., Gros, V., Williams, J., and Scheeren, H. A.: Continuing emissions of methyl chloroform from Europe, Nature, 421, 131–135, 2003.
Krol, M., Houweling, S., Bregman, B., van den Broek, M., Segers, A., van Velthoven, P., Peters, W., Dentener, F., and Bergamaschi, P.: The two-way nested global chemistry-transport zoom model TM5: algorithm and applications, Atmos. Chem. Phys., 5, 417–432, doi:10.5194/acp-5-417-2005, 2005.
Lamarque, J.-F., Bond, T. C., Eyring, V., Granier, C., Heil, A., Klimont, Z., Lee, D., Liousse, C., Mieville, A., Owen, B., Schultz, M. G., Shindell, D., Smith, S. J., Stehfest, E., Van Aardenne, J., Cooper, O. R., Kainuma, M., Mahowald, N., McConnell, J. R., Naik, V., Riahi, K., and van Vuuren, D. P.: Historical (1850–2000) gridded anthropogenic and biomass burning emissions of reactive gases and aerosols: methodology and application, Atmos. Chem. Phys., 10, 7017–7039, doi:10.5194/acp-10-7017-2010, 2010.
Lambert, G. and Schmidt, S.: Reevaluation of the oceanic flux of methane: uncertainties and long term variations, Chemosph. Global Change Sci., 26, 579–589, 1993.
Langenfelds, R. L., Francey, R. J., Pak, B. C., Steele, L. P., Lloyd, J., Trudinger, C. M., and Allison, C. E.: Interannual growth rate variations of atmospheric $CO_2$ and its $\delta^{13}C$, $H_2$, $CH_4$, and CO between 1992 and 1999 linked to biomass burning, Global Biogeochem. Cy., 16(3), 1048, 2002.
Law, R. M., Rayner, P. J., Denning, A. S., Erickson, D., Fung, I. Y., Heimann, M., Piper, S. C., Ramonet, M., Taguchi, S., Taylor, J. A., Trudinger, C. M., and Watterson, I. G.: Variations in modeled atmospheric transport of carbon dioxide and the consequences for $CO_2$ inversions, Global Biogeochem. Cy., 10, 783–796, doi:10.1029/96GB01892, 1996.





ACPD
11, 18767–18821, 2011



Law, R. M., Kowalczyk, E. A., and Wang, Y. P.: Using atmospheric $CO_2$ data to assess a simplified carbon-climate simulation for the 20th century, Tellus B, 53, 427–437, doi:10.1111/j.1600-0889.2006.00198.x, 2006.

Law, R. M., Peters, W., Rodenbeck, C., Aulagnier, C., Baker, I., Bergmann, D. J., Bousquet, P., Brandt, J., Bruhwiler, L., Cameron-Smith, P. J., Christensen, J. H., Delage, F., Denning, A. S., Fan, S., Geels, C., Houweling, S., Imasu, R., Karstens, U., Kawa, S. R., Kleist, J., Krol, M. C., Lin, S.-J., Lokupitiya, R., Maki, T., Maksyutov, S., Niwa, Y., Onishi, R., Parazoo, N., Patra, P. K., Pieterse, G., Rivier, L., Satoh, M., Serrar, S., Taguchi, S., Takigawa, M., Vautard, R., Vermeulen, A. T., and Zhu, Z.: TransCom model simulations of hourly atmospheric $CO_2$: experimental overview and diurnal cycle results for 2002, Global Biogeochem. Cy., 22, GB3009, doi:10.1029/2007GB003050, 2008.

Levin, I. and Hesshaimer, V.: Refining of atmospheric transport model entries by the globally observed passive tracer distributions of $^{85}$krypton and sulfur hexafluoride ($SF_6$), J. Geophys. Res., 101, 16745–16755, 1996.

Levin, I., Naegler, T., Heinz, R., Osusko, D., Cuevas, E., Engel, A., Ilmberger, J., Langenfelds, R. L., Neininger, B., Rohden, C. v., Steele, L. P., Weller, R., Worthy, D. E., and Zimov, S. A.: The global $SF_6$ source inferred from long-term high precision atmospheric measurements and its comparison with emission inventories, Atmos. Chem. Phys., 10, 2655–2662, doi:10.5194/acp-10-2655-2010, 2010.

Matthews, E. and Fung, I.: Methane emissions from natural wetlands: Global distribution, area, and ecology of sources, Global Biogeochem. Cy., 1, 61–86, doi:10.1029/GB001i001p00061, 1987.

McCulloch, A. and Midgley, P. M.: The history of methyl chloroform emissions: 1951–2000, Atmos. Environ., 35, 5311–5319, 2001.

Mikaloff-Fletcher, S. E., Tans, P. P., Bruhwiler, L. M., Miller, J. B., and Heimann, M.: $CH_4$ sources estimated from atmospheric observations of $CH_4$ and its $^{13}C/^{12}C$ isotopic ratios: 2. Inverse modeling of $CH_4$ fluxes from geographical regions, Global Biogeochem. Cy., 18, GB4005, doi:10.1029/2004GB002224, 2004.

Mitchell, T. D. and Jones, P. D.: An improved method of constructing a database of monthly climate observations and associated high-resolution grids, Int. J. Climatol., 25, 693–712, doi:10.1002/joc.1181, 2005.





ACPD
11, 18767–18821, 2011



Miyazaki, K., Patra, P. K., Takigawa, M., Iwasaki, T., and Nakazawa, T.: Global-scale transport of carbon dioxide in the troposphere, J. Geophys. Res., 113, D15301, doi:10.1029/2007JD009557, 2008.

Olivier, J. G. J. and Berdowski, J. J. M.: Global emissions sources and sinks, in: The Climate System, edited by: Berdowski, J., Guicherit, R., and Heij, B. J., ISBN 9058092550, A. A. Balkema Publishers/Swets & Zeitlinger Pub., Lisse, The Netherlands, 33–78, 2001.

Onogi, K., Tsutsui, J., Koide, H., Sakamoto, M., Kobayashi, S., Hatsushika, H., Matsumoto, T., Yamazaki, N., Kamahori, H., Takahashi, K., Kadokura, S., Wada, K., Kato, K., Oyama, R., Ose, T., Mannoji, N., and Taira, R.: The JRA-25 reanalysis, J. Meteorol. Soc. Jpn., 85, 369–432, 2007.

Park, J. H., Russell, J. M., Gordley, L. L., Drayson, S. R., Benner, D. C., McInerney, J. M., Gunson, M. R., Toon, G. C., Sen, B., Blavier, J. F., Webster, C. R., Zipf, E. C., Erdman, P., Schmidt, U., and Schiller, C.: Validation of halogen occultation experiment $CH_4$ measurements from the UARS, J. Geophys. Res., 101, 10183–10203, 1996.

Patra, P. K., Law, R. M., Peters, W., Rodenbeck, C., Takigawa, M., Aulagnier, C., Baker, I., Bergmann, D. J., Bousquet, P., Brandt, J., Bruhwiler, L., Cameron-Smith, P. J., Christensen, J. H., Delage, F., Denning, A. S., Fan, S., Geels, C., Houweling, S., Imasu, R., Karstens, U., Kawa, S. R., Kleist, J., Krol, M. C., Lin, S.-J., Lokupitiya, R., Maki, T., Maksyutov, S., Niwa, Y., Onishi, R., Parazoo, N., Pieterse, G., Rivier, L., Satoh, M., Serrar, S., Taguchi, S., Vautard, R., Vermeulen, A. T., and Zhu, Z.: TransCom model simulations of hourly atmospheric $CO_2$: analysis of synoptic-scale variations for the period 2002–2003, Global Biogeochem. Cy., 22, GB4013, doi:10.1029/2007GB003081, 2008.

Patra, P. K., Takigawa, M., Ishijima, K., Choi, B.-C., Cunnold, D., Dlugokencky, E. J., Fraser, P., Gomez-Pelaez, A. J., Goo, T.-Y., Kim, J.-S., Krummel, P., Langenfelds, R., Meinhardt, F., Mukai, H., O'Doherty, S., Prinn, R. G., Simmonds, P., Steele, P., Tohjima, Y., Tsuboi, K., Uhse, K., Weiss, R., Worthy, D., and Nakazawa, T.: Growth rate, seasonal, synoptic, diurnal variations and budget of methane in lower atmosphere, J. Meteorol. Soc. Jpn., 87(4), 635–663, 2009a.

Patra, P. K., Takigawa, M., Dutton, G. S., Uhse, K., Ishijima, K., Lintner, B. R., Miyazaki, K., and Elkins, J. W.: Transport mechanisms for synoptic, seasonal and interannual $SF_6$ variations and "age" of air in troposphere, Atmos. Chem. Phys., 9, 1209–1225, doi:10.5194/acp-9-1209-2009, 2009b.





ACPD
11, 18767–18821, 2011



Patra, P. K., Houweling, S., Krol, M., Bousquet, P., Bruhwiler, L., and Jacob, D.: Protocol for TransCom $CH_4$ intercomparison, Version 7, http://www.jamstec.go.jp/frcgc/research/d4/prabir/papers/TC_CH4protocol_v7.pdf (last access: 26 April 2010), 2010.

Pickett-Heaps, C. A., Jacob, D. J., Wecht, K. J., Kort, E. A., Wofsy, S. C., Diskin, G. S., Worthy, D. E. J., Kaplan, J. O., Bey, I., and Drevet, J.: Magnitude and seasonality of wetland methane emissions from the Hudson Bay Lowlands (Canada), Atmos. Chem. Phys., 11, 3773–3779, doi:10.5194/acp-11-3773-2011, 2011.

Plumb, R. A.: A "tropical pipe" model of stratospheric transport, J. Geophys. Res., 101, 3957–3972, doi:10.1029/95JD03002, 1996.

Prigent, C., Papa, F., Aires, F., Rossow, W. B., and Matthews, E.: Global inundation dynamics inferred from multiple satellite observations, 1993–2000, J. Geophys. Res., 112, D12107, doi:10.1029/2006JD007847, 2007.

Prinn, R. G., Weiss, R., Fraser, P., Simmonds, P., Cunnold, D., Alyea, F., O'Doherty, S., Salameh, P., Miller, B., Huang, J., Wang, R., Hartley, D., Harth, C., Steele, L., Sturrock, G., Midgley, P., and McCulloch, A.: A history of chemically and radiatively important gases in air deduced from ALE/GAGE/AGAGE, J. Geophys. Res., 105, 17751–17792, doi:10.1029/2000JD900141, 2000.

Prinn, R. G., Huang, J., Weiss, R. F., Cunnold, D. M., Fraser, P. J., Simmonds, P. G., McCulloch, A., Harth, C., Reimann, S., Salameh, P., O'Doherty, S., Wang, R. H. J., Porter, L. W., Miller, B. R., and Krummel, P. B.: Evidence for variability of atmospheric hydroxyl radicals over the past quarter century, Geophys. Res. Lett., 32, L07809, doi:10.1029/2004GL022228, 2005.

Rasmussen, R. A. and Khalil, M. A. K.: Atmospheric methane in recent and ancient atmospheres: concentrations, trends, and interhemispheric gradient, J. Geophys. Res., 89, 11599–11605, 1984.

Rigby, M., Prinn, R. G., Fraser, P. J., Simmonds, P. G., Langenfelds, R. L., Huang, J., Cunnold, D. M., Steele, L. P., Krummel, P. B., Weiss, R. F., O'Doherty, S., Salameh, P. K., Wang, H. J., Harth, C. M., Muhle, J., and Porter, L. W.: Renewed growth of atmospheric methane, Geophys. Res. Lett., 35, L22805, doi:10.1029/2008GL036037, 2008.

Rigby, M., Mühle, J., Miller, B. R., Prinn, R. G., Krummel, P. B., Steele, L. P., Fraser, P. J., Salameh, P. K., Harth, C. M., Weiss, R. F., Greally, B. R., O'Doherty, S., Simmonds, P. G., Vollmer, M. K., Reimann, S., Kim, J., Kim, K.-R., Wang, H. J., Olivier, J. G. J., Dlugokencky, E. J., Dutton, G. S., Hall, B. D., and Elkins, J. W.: History of atmospheric $SF_6$ from





ACPD
11, 18767–18821, 2011



1973 to 2008, Atmos. Chem. Phys., 10, 10305–10320, doi:10.5194/acp-10-10305-2010, 2010.

Ringeval, B., de Noblet-Ducoudré, N., Ciais, P., Bousquet, P., Prigent, C., Papa, F., and Rossow, W. B.: An attempt to quantify the impact of changes in wetland extent on methane emissions on the seasonal and interannual time scales, Global Biogeochem. Cy., 24, GB2003, doi:10.1029/2008GB003354, 2010.

Rotman, D. A., Atherton, C., Bergmann, D. J., Cameron-Smith, P. J., Chuang, C. C., Connell, P. S., Dignon, J. E., Franz, A., Grant, K. E., Kinnison, D. E., Molenkamp, C. R., Proctor, D. D., and Tannahill, J. R.: IMPACT, the LLNL 3-D global atmospheric chemical transport model for the combined troposphere and stratosphere: model description and analysis of ozone and other trace gases, J. Geophys. Res., 109, D04303, doi:10.1029/2002JD003155, 2004.

Sander, S. P., Golden, D. M., Kurylo, M. J., Moortgat, G. K., Keller-Rudek, H., Wine, P. H., Ravishankara, A. R., Kolb, C. E., Molina, M. J., Finlayson-Pitts, B. J., Huie, R. E., and Orkin, V. L.: Chemical Kinetics and Photochemical Data for Use in Atmospheric Studies, Evaluation Number 15, JPL Publication 06-2, Jet Propulsion Laboratory, California Institute of Technology, Pasadena, CA, 2006.

Seidel, D. J., Fu, Q., Randel, W. J., and Reichler, T. J.: Widening of the tropical belt in a changing climate, Nat. Geosci., 1, 21–24, doi:10.1038/ngeo.2007.38, 2008.

Shindell, D. T., Faluvegi, G., Stevenson, D. S., Krol, M. C., Emmons, L. K., Lamarque, J.-F., Petron, G., Dentener, F. J., Ellingsen, K., Schultz, M. G., Wild, O., Amann, M., Atherton, C. S., Bergmann, D. J., Bey, I., Butler, T., Cofala, J., Collins, W. J., Derwent, R. G., Doherty, R. M., Drevet, J., Eskes, H. J., Fiore, A. M., Gauss, M., Hauglustaine, D. A., Horowitz, L. W., Isaksen, I. S. A., Lawrence, M. G., Montanaro, V., Muller, J. F., Pitari, G., Prather, M. J., Pyle, J. A., Rast, S., Rodriguez, J. M., Sanderson, M. G., Savage, N. H., Strahan, S. E., Sudo, K., Szopa, S., Unger, N., van Noije, T. P. C., and Zeng, G.: Multi-model simulations of carbon monoxide: comparison with observations and projected near-future changes, J. Geophys. Res., 111, D19306, doi:10.1029/2006JD007100, 2006.

Shindell, D. T., Faluvegi, G., Koch, D. M., Schmidt, G. A., Unger, N., and Bauer, S. E.: Improved attribution of climate forcing to emissions, Science, 326, 716–718, doi:10.1126/Science.1174760, 2009.





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Simpson, I. J., Rowland, F. S., Meinardi, S., and Blake, D. R.: Influence of biomass burning during recent fluctuations in the slow growth of global tropospheric methane, Geophys. Res. Lett., 33, L22808, doi:10.1029/2006GL027330, 2006.

Spivakovsky, C., Logan, J. A., Montzka, S. A., Balkanski, Y. J., Foreman-Fowler, M., Jones, D. B. A., Horowitz, L. W., Fusco, A. C., Brenninkmeijer, C. A. M., Prather, M. J., Wofsy, S. C., McElroy, M. B.: Three-dimensional climatological distribution of tropospheric OH: update and evaluation, J. Geophys. Res., 105(D7), 8931–8980, 2000.

Stevenson, D. S., Dentener, F. J., Schultz, M. G., Ellingsen, K., van Noije, T. P. C., Wild, O., Zeng, G., Amann, M., Atherton, C. S., Bell, N., Bergmann, D. J., Bey, I., Butler, T., Cofala, J., Collins, W. J., Derwent, R. G., Doherty, R. M., Drevet, J., Eskes, H. J., Fiore, A. M., Gauss, M., Hauglustaine, D. A., Horowitz, L. W., Isaksen, I. S. A., Krol, M. C., Lamarque, J.-F., Lawrence, M. G., Montanaro, V., Muller, J.-F., Pitari, G., Prather, M. J., Pyle, J. A., Rast, S., Rodriguez, J. M., Sanderson, M. G., Savage, N. H., Shindell, D. T., Strahan, S. E., Sudo, K., and Szopa, S.: Multimodel ensemble simulations of present-day and near-future tropospheric ozone, J. Geophys. Res., 111, D08301, doi:10.1029/2005JD006338, 2006.

Sudo, K., Takahashi, M., Kurokawa, J., and Akimoto, H.: CHASER: a global chemical model of the troposphere 1. Model description, J. Geophys. Res., 107, 4339, doi:10.1029/2001JD001113, 2002.

Takigawa, M., Takahashi, M., and Akiyoshi, H.: Simulation of ozone and other chemical species using a Center for Climate System Research/National Institute for Environmental Studies atmospheric GCM with coupled stratospheric chemistry, J. Geophys. Res., 104, 14003–14018, 1999.

Uppala, S. M., Koallberg, P. W., Simmons, A. J., Andrae, U., da Costa Bechtold, V., Fiorino, M., Gibson, J. K., Haseler, J., Hernandez, A., Kelly, G., Li, X., Onogi, K., Saarinen, S., Sokka, N., Allan, R. P., Andersson, E., Arpe, K., Balmaseda, M. A., Beljaars, A. C. M., van de Berg, L., Bidlot, J., Bormann, N., Caires, S., Chevallier, F., Dethof, A., Dragosavac, M., Fisher, M., Fuentes, M., Hagemann, S., Holm, E., Hoskins, B. J., Isaksen, L., Janssen, P. A. E. M., Jenne, R., McNally, A. P. A., Mahfouf, J.-F., Morcrette, J.-J., Rayner, N. A., Saunders, R. W., Simon, P., Sterl, A., Trenberth, K. E., Untch, A., Vasiljevic, D., Viterbo, P., and Woollen, J.: The ERA-40 re-analysis, Q. J. Roy. Meteorol. Soc., 131, 2961–3012, doi:10.1256/qj.04.176, 2005.





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van der Werf, G. R., Randerson, J. T., Giglio, L., Collatz, G. J., Kasibhatla, P. S., and Arellano Jr., A. F.: Interannual variability in global biomass burning emissions from 1997 to 2004, Atmos. Chem. Phys., 6, 3423–3441, doi:10.5194/acp-6-3423-2006, 2006.

Velders, G.: Description of the RIVM 2-dimensional stratosphere model, RIVM Report 722201002, The Netherland, 1995.

Wang, J. S., Logan, J. A., McElroy, M. B., Duncan, B. N., Megretskaia, I. A., and Yantosca, R. M.: A 3-D model analysis of the slowdown and interannual variability in the methane growth rate from 1988 to 1997, Global Biogeochem. Cy., 18, GB3011, doi:10.1029/2003GB002180, 2004.

WDCGG – WMO World Data Centre for Greenhouse Gases: Japan Meteorological Agency, Tokyo, online available at: http://gaw.kishou.go.jp/wdcgg.html (last access: 19 April 2010), 2010.

WMO/SAOD, World Meteorological Organization, Scientific Assessment of Ozone Depletion: Global Ozone Research Monitoring Project – Report No. 47, WMO, Geneva, 498 pp., 2003.

Wofsy, S. C., the HIPPO Science Team and Cooperating Modellers and Satellite Teams: HIAPER pole-to-pole observations (HIPPO): fine grained, global scale measurements for determining rates for transport, surface emissions, and removal of climatologically important atmospheric gases and aerosols, Philos. T. Roy. Soc. A, 369, 2073–2086, 2011.

Xiong, X., Barnet, C. D., Maddy, E., Sweeney, C., Liu, X., Zhou, L., and Goldberg, M.: Characterization and validation of methane products from the atmospheric infrared sounder (AIRS), J. Geophys. Res., 113, G00A01, doi:10.1029/2007JG000500, 2008.

Yan, X., Akiyama, H., Yagi, K., and Akimoto, H.: Global estimations of the inventory and mitigation potential of methane emissions from rice cultivation conducted using the 2006 intergovernmental panel on climate change guidelines, Global Biogeochem. Cy., 23, GB2002, doi:10.1029/2008GB003299, 2009.

Yoshida, Y., Ota, Y., Eguchi, N., Kikuchi, N., Nobuta, K., Tran, H., Morino, I., and Yokota, T.: Retrieval algorithm for $CO_2$ and $CH_4$ column abundances from short-wavelength infrared spectral observations by the Greenhouse gases observing satellite, Atmos. Meas. Tech., 4, 717–734, doi:10.5194/amt-4-717-2011, 2011.





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**Table 1.** List of tracers simulated in the TransCom-$CH_4$ intercomparison project. See Sect. 2.2 for a description of the $CH_4$ flux components.

| PARAMETERS | DESCRIPTION | TIME RESOLUTION |
|---|---|---|
| **$CH_4$ tracers** | | |
| 1. $CH_4$_CTL | CYC NAT (CYC BB & CYC WL) + IAV ANT EDGAR 3.2 | Monthly; Partial IAV |
| 2. $CH_4$_CTL_E4 | CYC NAT + IAV ANT EDGAR4.0 | Monthly; Partial IAV |
| 3. $CH_4$_BB | $CH_4$_CTL – 0.35 CYC BB + IAV BB | Monthly; Partial IAV |
| 4. $CH_4$_WL_BB | $CH_4$_BB – CYC WL + 0.76 IAV WL | Monthly; Full IAV |
| 5. $CH_4$_INV | IPSL/LSCE inversion | Monthly; Full IAV |
| 6. $CH_4$_EXTRA* | $CH_4$_BB – CYC WL – Rice + IAV WLe (0.69 Wetland + 0.895 Rice) | Monthly; Full IAV |
| **Other tracers** | | |
| 7. $SF_6$ | EDGAR 4.0 (@JRC); Global totals modified | Annual; Full IAV |
| 8. Radon-222 ($^{222}Rn$) | 1.0 and 0.1 atom $m^{-2}$ $s^{-1}$ over land and ocean, respectively | Annual; No IAV |
| 9. $CH_3CCl_3$ (MCF) | EDGAR3.2 with trends and distributions modified | Annual; Full IAV |

* this tracer is called EXTRA because the VISIT terrestrial ecosystem model (Ito, 2010) fluxes are still under evaluation, but included here since no other bottom-up wetland emission scenario was available with IAV for the full simulation period at the time the intercomparison protocol was released. VISIT is driven by climate variables from the Climate Research Unit time series version 3.0 (CRU TS3.0) dataset (Mitchell and Jones, 2005; updated values) and NCEP/NCAR reanalysis (Kalnay et al., 1996) for the periods of 1988–2005 and 2006–2007, respectively, and $CH_4$ cycling in the inundated areas is modeled using the scheme of Cao et al. (1998 and references therein).





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**Table 2.** Overview of participating transport models and model variants, and average lifetimes of atmospheric $CH_4$ and $CH_3CCl_3$ are given.

| Sl. No. | Model Name[a] | Institution[b] | Resolution | | Meteorology[e] | Avg. lifetime (1992–07) | |
|---|---|---|---|---|---|---|---|
| | | | Horizontal[c] | Vertical[d] | | $CH_4$_CTL | $CH_3CCl_3$ |
| 1 | **ACCESS** | CSIRO | 3.75 × 2.5° | 38 | AGCM; SST | 9.93 ± 0.13 | 4.55 ± 0.15 |
| 2 | **ACTM** | RIGC | ∼2.8 × 2.8° | 67$\sigma$ | NCEP2; U, V, T; SST | 10.0 ± 0.10 | 4.60 ± 0.13 |
| 2a | ***ACTM_OH***[$] | RIGC | ∼2.8 × 2.8° | 67$\sigma$ | NCEP2; U, V, T; SST | 9.51 ± 0.10 | 4.84 ± 0.13 |
| 3 | CAM | CU | 2.5 × ∼1.9° | 28$\sigma$ | NCEP/NCAR | 10.2 ± 0.11 | 3.77 ± 0.13 |
| 4 | **CCAM** | CSIRO | ∼220 km | 18$\sigma$ | NCEP; U, V; SST | 9.94 ± 0.27 | 4.01 ± 0.15 |
| 5 | GEOS-Chem | UoE | 2.5 × 2.0° | 30/47$\eta$ | NASA/GSFC/GEOS4/5 | 9.60 ± 0.11 | 4.70 ± 0.13 |
| 5a | *GEOS-Chem_DOH* | UoE | 2.5 × 2.0° | 30/47$\eta$ | NASA/GSFC/GEOS4/5 | 9.95 ± 0.11 | 4.84 ± 0.13 |
| 6 | IMPACT | LLNL | 5.0 × 4.0° | 55$\eta$ | NASA/GSFC/GEOS4 | 10.1 ± 0.05 | 4.63 ± 0.3 |
| 6a | *IMPACT_1 × 1.25* | LLNL | 1.25 × 1.0° | 55$\eta$ | NASA/GSFC/GEOS4 | 9.99 ± 0.07 | 4.54 ± 0.16 |
| 7 | **LMDZ** | LSCE | 3.75 × 2.5° | 19$\eta$ | ECMWF; U, V, T; SST | 10.0 ± 0.09 | 3.90 ± 0.25 |
| 8 | MOZART | MIT | ∼1.8 × 1.8° | 28$\sigma$ | NCEP/NCAR | 9.88 ± 0.15 | 3.90 ± 0.15 |
| 9 | NIES08i | NIES | 2.5 × 2.5° | 32$\sigma$-$\theta$ | JCDAS, ERA-interim-PBL | 10.0 ± 0.06 | 4.75 ± 0.02 |
| 10 | PCTM | GSFC | 1.25 × 1.0° | 58$\eta$ | NASA/GSFC/GEOS5 | 10.1 ± 0.1 | 4.54 ± 0.21 |
| 11 | TM5 | SRON | 6.0 × 4.0° | 25$\eta$ | ECMWF, ERA-interim | 10.1 ± 0.12 | 4.92 ± 0.50 |
| 11a | *$TM_5$-1 × 1* | SRON | 1.0 × 1.0° | 25$\eta$ | ECMWF, ERA-interim | 10.1 ± 0.11 | 4.88 ± 0.14 |
| 12 | TOMCAT | UoL | ∼2.8 × 2.8° | 60$\eta$ | ECMWF, ERA-40/interim | 9.98 ± 0.12 | 4.71 ± 0.18 |

[a] CTMs driven by AGCM transport are identified in bold (nudging parameters in right-most column), and model variants are shown in italics. The model variants are indicated by post-fixed parameters, following a "_".

[b] CSIRO: Commonwealth Scientific and Industrial Research Organisation, Australia; GSFC: NASA Goddard Space Flight Center, USA; RIGC: Research Institute for Global Change, Japan; CU: Cornel University, USA; LLNL: Lawrence Livermore National Laboratory, USA; LSCE: Laboratoire des Sciences du Climat et de l'Environnement, France; NIES: National Institute for Environmental Studies, Japan; SRON: Netherlands Institute for Space Research; UoE: University of Edinburgh, UK; UoL: University of Leeds, UK.

[c] Longitude × latitude or distance or spectral resolution indicated by T (triangular) maximum wave number (T42 and T63 for ∼2.8 × 2.8° and ∼1.8 × 1.8°, respectively).

[d] Terrain-following (height) coordinate system for ACCESS, $\sigma$ vertical coordinates are pressure divided by surface pressure, $\eta$ vertical coordinates are a hybrid sigma-pressure coordinate (GEOS-Chem has 30 or 47 layers for 1990–2006 or 2007, respectively), NIES08i has a hybrid sigma-isentropic.

[e] The source of meteorology (NCEP2 (AMIP DOE II): Kanamitsu et al., 2002; NCEP: Kalnay et al., 1996; NASA/GSFC/GEOS4/5: Bloom et al., 2005; ECMWF: Uppala et al., 2005; JCDAS: Onogi et al., 2007) and parameters used in nudged AGCMs are given.

[$] The tropospheric OH field is taken from CHASER full chemistry model [Sudo et al., 2002] and scaled by × 0.88, and stratospheric OH is taken from AGCM (Takigawa et al., 1999) as discussed in Patra et al. (2009a). $SF_6$ emissions are used from EDGAR4.0 without scaling the global totals to match with Levin et al. (2010).





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**Table 3.** Multi-model averages ($\pm 1\sigma$; and between model variability defined by $1\sigma$/average, within parenthesis in %) of simulated $SF_6$ (ppt), $CH_4$_CTL (ppb), $CH_3CCl_3$ (ppt) and $^{222}Rn$ ($\times 10^{-21}$) gradients in the troposphere and UT/LS region for three broad latitude bands, namely, the SH midlatitude, tropics and NH midlatitude.

| Species | SH (60–30° S) | Tropics (15° S–15° N) | NH (30–60° N) |
|---|---|---|---|
| | | Tropospheric gradients | |
| Difference between: | 850–400 mb | 850–200 mb | 850–400 mb |
| $SF_6$ | −0.026 ± 0.005 (18 %) | 0.044 ± 0.017 (39 %) | 0.128 ± 0.018 (14 %) |
| $CH_4$ | −9.91 ± 2.57 (26 %) | 17.18 ± 7.69 (45 %) | 44.65 ± 5.91 (13 %) |
| $CH_3CCl_3$ | 0.473 ± 0.145 (31 %) | 0.295 ± 0.182 (62 %) | 1.01 ± 0.25 (25 %) |
| $^{222}Rn$ | 0.984 ± 0.527 (54 %) | 8.66 ± 2.73 (32 %) | 23.90 ± 6.73 (28 %) |
| | | UT/LS gradients | |
| Difference between: | 200–100 mb | 100–50 mb | 200–100 mb |
| $SF_6$ | 0.158 ± 0.074 (47 %) | 0.264 ± 0.148 (56 %) | 0.229 ± 0140 (61 %) |
| $CH_4$ | 84.45 ± 35.89 (43 %) | 124.04 ± 65.07 (52 %) | 128.77 ± 64.96 (50 %) |
| $CH_3CCl_3$ | 7.86 ± 1.37 (17 %) | 16.59 ± 4.48 (27 %) | 9.79 ± 3.75 (38 %) |
| $^{222}Rn$ | 1.084 ± 0.708 (65 %) | 0.415 ± 0.203 (49 %) | 0.353 ± 0445 (126 %) |





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**Table 4.** Average (across all models) correlation coefficient ($r$) between model simulated and observed seasonal cycles and interannual variation of $CH_4$, $SF_6$ and $CH_3CCl_3$.

| Tracer \ Site* | ALT | BRW | MHD | MLO | RPB | SMO | CGO | SPO |
|---|---|---|---|---|---|---|---|---|
| For seasonal cycles (2002–2003) | | | | | | | | |
| $CH_4$_CTL | .16 ± .3 | −.09 ± .2 | .73 ± .1 | .63 ± .1 | .86 ± .1 | .74 ± .2 | .98 ± .02 | .96 ± .1 |
| $CH_4$_CTL_E4 | .21 ± .3 | −.10 ± .3 | .72 ± .1 | .69 ± .1 | .88 ± .1 | .71 ± .2 | .97 ± .02 | .96 ± .1 |
| $CH_4$_BB | .10 ± .3 | −.11 ± .2 | .68 ± .1 | .66 ± .1 | .86 ± .1 | .72 ± .2 | .95 ± .1 | .96 ± .1 |
| $CH_4$_WL_BB | .38 ± .3 | .48 ± .2 | .42 ± .2 | .70 ± .1 | .86 ± .1 | .64 ± .3 | .94 ± .1 | .95 ± .1 |
| $CH_4$_INV | .88 ± .1 | .63 ± .2 | .92 ± .1 | .74 ± .1 | .88 ± .1 | .61 ± .2 | .95 ± .02 | .92 ± .1 |
| $CH_4$_EXTRA | .84 ± .1 | .37 ± .2 | .79 ± .1 | .50 ± .2 | .80 ± .1 | .69 ± .3 | .93 ± .1 | .95 ± .1 |
| $SF_6$ | .48 ± .2 | .23 ± .2 | .68 ± .1 | −.06 ± .1 | .13 ± .2 | .70 ± .4 | .43 ± .2 | .50 ± .2 |
| $CH_3CCl_3$ | .78 ± .1 | .93 ± .02 | .87 ± .1 | .91 ± .1 | .96 ± .02 | .89 ± .1 | .88 ± .1 | .81 ± 1 |
| For interannual variations ($CH_4$, $CH_3CCl_3$: 1991–2007; $SF_6$: 1996–2007) | | | | | | | | |
| CH4_CTL | .52 ± .2 | .14 ± .2 | .66 ± .1 | .32 ± .2 | .34 ± .2 | .38 ± .2 | .46 ± .2 | .43 ± .2 |
| CH4_CTL_E4 | .27 ± .2 | .01 ± .2 | .46 ± .2 | −.02 ± .2 | .05 ± .3 | .24 ± .2 | .03 ± .2 | −.06 ± .2 |
| $CH_4$_BB | .69 ± .2 | .40 ± .2 | .76 ± .1 | .47 ± .2 | .75 ± .1 | .54 ± 2 | .60 ± .2 | .57 ± .2 |
| $CH_4$_WL_BB | .55 ± .2 | .47 ± .2 | .51 ± .2 | .38 ± .2 | .68 ± .2 | .53 ± .2 | .48 ± .1 | .46 ± .2 |
| $CH_4$_INV | .64 ± .1 | .32 ± .2 | .68 ± .1 | .27 ± .2 | .42 ± .2 | .38 ± .2 | .63 ± .2 | .56 ± .2 |
| $CH_4$_EXTRA | .61 ± .2 | .36 ± .2 | .77 ± .1 | .51 ± .2 | .77 ± .2 | .73 ± .1 | .68 ± .1 | .69 ± .1 |
| $SF_6$ | .33 ± .2 | .67 ± .1 | .75 ± .2 | .70 ± .2 | – | .34 ± .1 | .83 ± .1 | .81 ± .1 |
| $CH_3CCl_3$ | .97 ± .03 | .97 ± .03 | .95 ± .04 | .96 ± .03 | .95 ± .04 | .96 ± .04 | .95 ± .03 | .94 ± .03 |

* ALT (Alert, Canada; 62° W, 82° N, 210 m), BRW (Point Barrow, Alaska, USA; 157° W, 71° N, 11 m), MLO (Mauna Loa Observatory, Hawaii, USA; 156° W, 20° N, 3397 m) and SPO (South Pole Observatory, Antarctica; 25° W, 90° S, 2810 m) are managed under the NOAA cooperative network by the Global Monitoring Division, Earth System Research Laboratory (GMD/ESRL) (Dlugokencky et al., 1998; Butler et al., 2004), and MHD (Mace Head, Ireland; 10° W, 53° N, 25 m), RPB (Ragged Point, Barbados; 59° W, 13° N, 45 m), SMO (Samoa, USA; 171° W, 14° S, 42 m) and CGO (Cape Grim, Australia; 145° E, 41° S, 94 m) sites are operated under the AGAGE network by the Massachusetts Institute of Technology (MIT, USA), Scripps Institutions of Oceanography, University of California, San Diego (SIO/UCSD, USA), Commonwealth Scientific and Industrial Research Organization (CSIRO, Australia), University of Bristol, UK and Georgia Institute of Technology, USA (Cunnold et al., 2002; Prinn et al., 2000)





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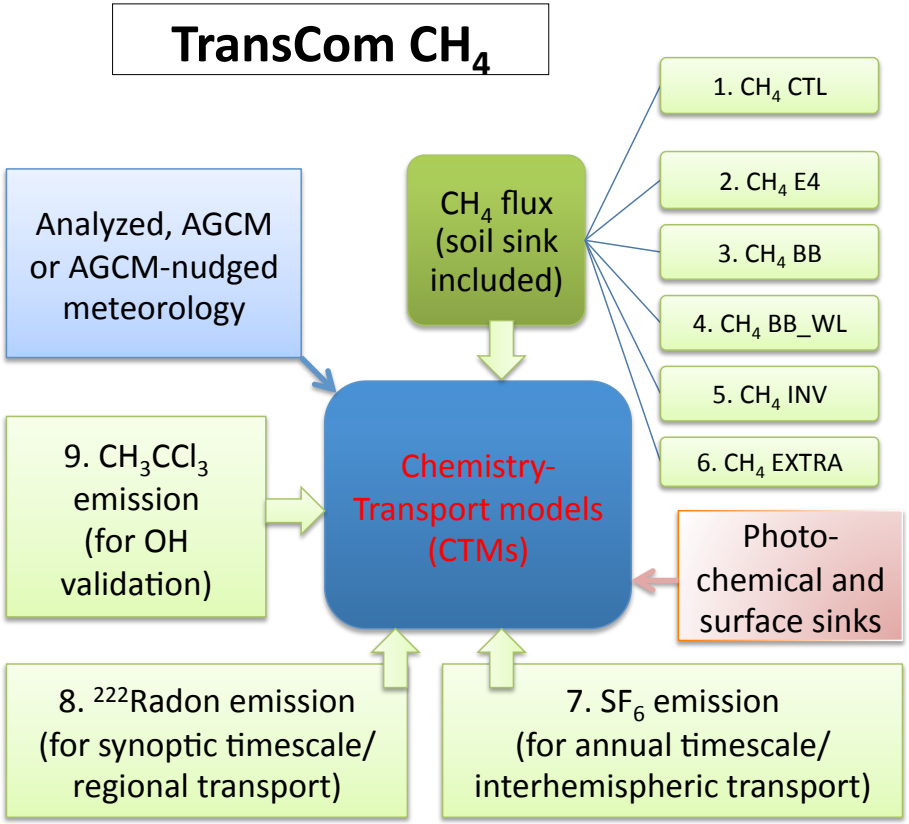


**Fig. 1.** Schematic diagram of TransCom-$CH_4$ model simulation experiment.





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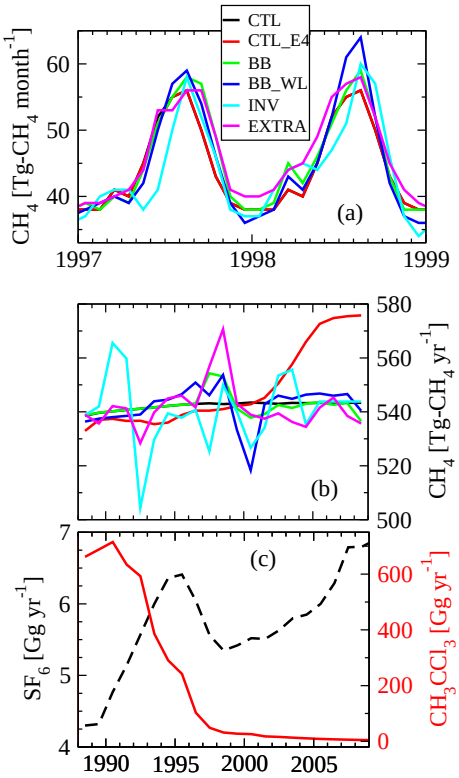


**Fig. 2.** The average $CH_4$ flux seasonalities are shown in **(a)**, and annual mean $CH_4$ fluxes for the period of 1988–2008 are depicted in **(b)**. The annual mean $SF_6$ and $CH_3CCl_3$ fluxes are shown in **(c)**.





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TransCom model simulations of $CH_4$ and related species

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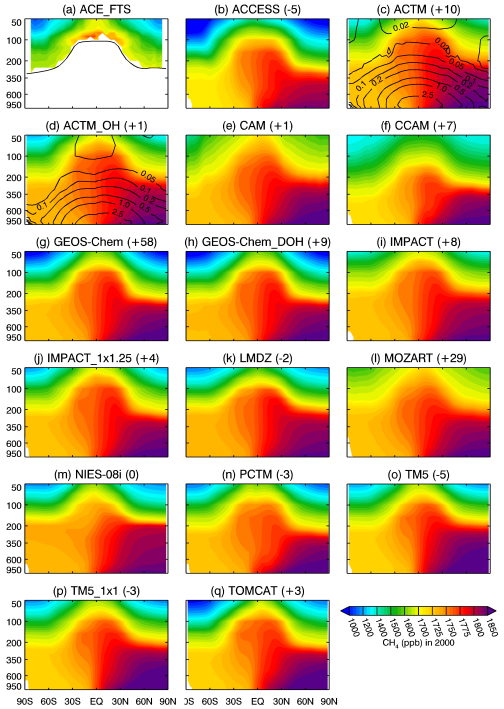


**Fig. 3.** Annual and zonal mean latitude-pressure (in mb) cross-sections of $CH_4$ for tropospheric and lower stratospheric altitudes as observed by the ACE-FTS instrument (**a**; climatology) and as simulated by the models in 2000 **(b–o)**. The black line in **(a)** shows the climatological tropopause height. The contour lines in (**c** and **d**) show $CH_4$ loss rate (units: molecule $cm^{-3}$ $s^{-1}$) as in the ACTM and ACTM_OH, respectively. An offset is added to the concentrations in each panel (given after the model name in ppb) that adjusts the model fields to a common average value of 1770 ppb between 950 mb and 500 mb. Detailed model-to-model comparisons for $^{222}Rn$, $SF_6$, $CH_4$ and $CH_3CCl_3$ for two seasons and over two longitudes are available in the Supplement (Figs. S1–S17).





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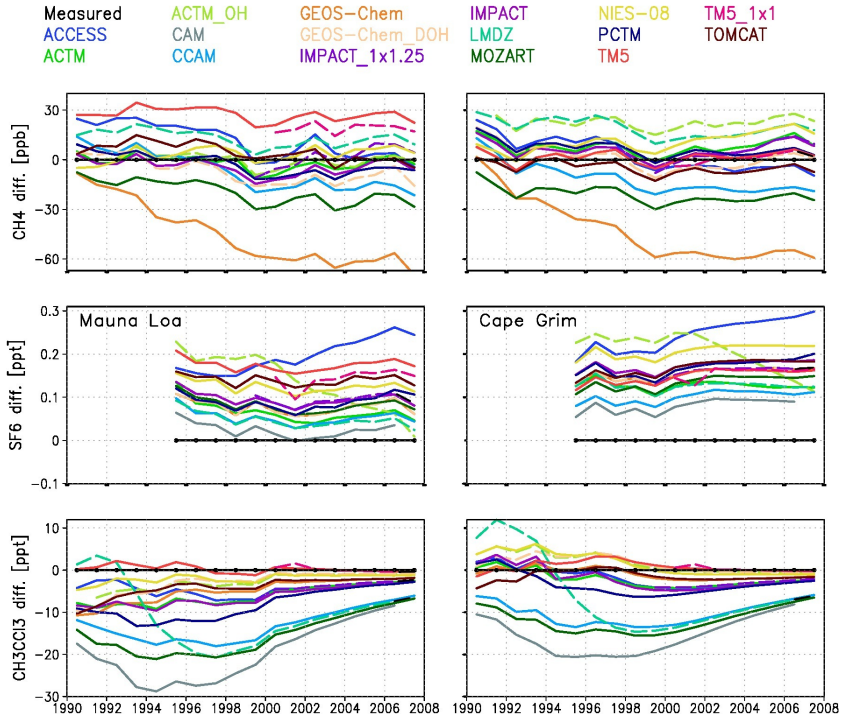


**Fig. 4.** Time series of differences between observed and simulated annual mean $CH_4$-CTL (top row), $SF_6$ (middle row) and $CH_3CCl_3$ (bottom row) at two selected sites, MLO (left column) and CGO (right column). CAM is excluded for $CH_4$-CTL plots, because soil sink was not accounted for during simulation. Time series of annual mean values and tropospheric model values, averaged over 1000–200 mb and all latitudes/longitudes, are shown in Figs. S18 and S19, respectively.





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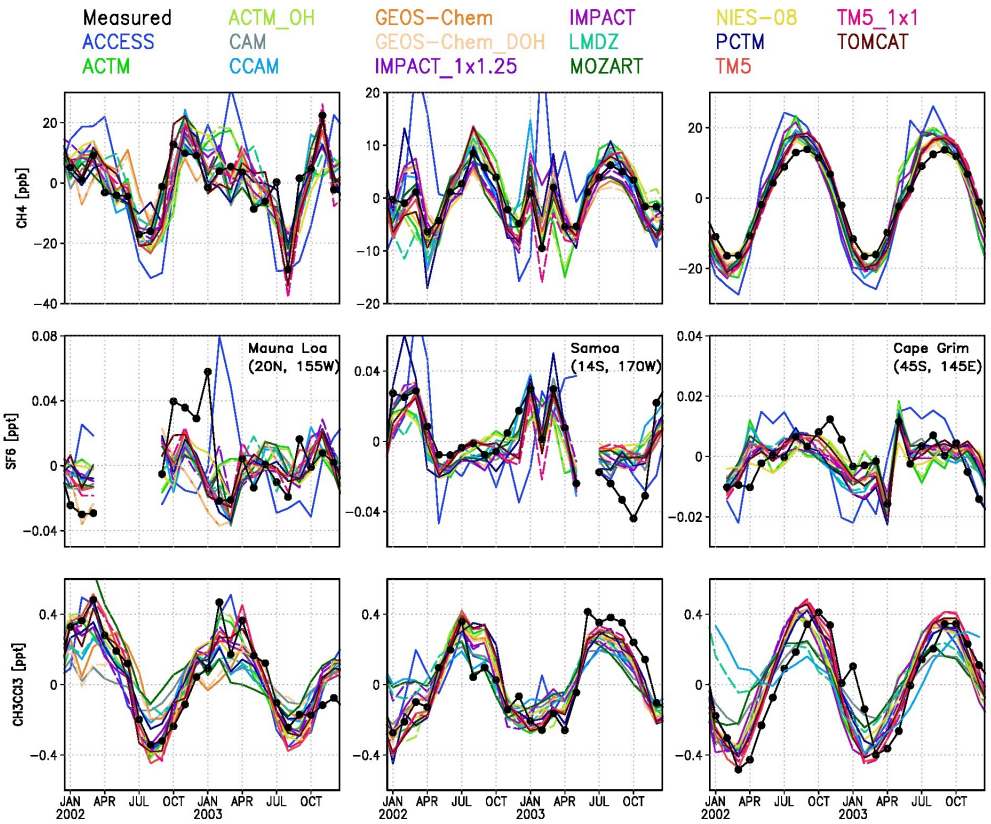


**Fig. 5.** Comparisons of observed and simulated seasonal cycles of $CH_4$_CTL (top row), $SF_6$ (middle row) and $CH_3CCl_3$ (bottom row) at 3 selected sites, MLO (left panel), SMO (middle panel) and CGO (right panel).





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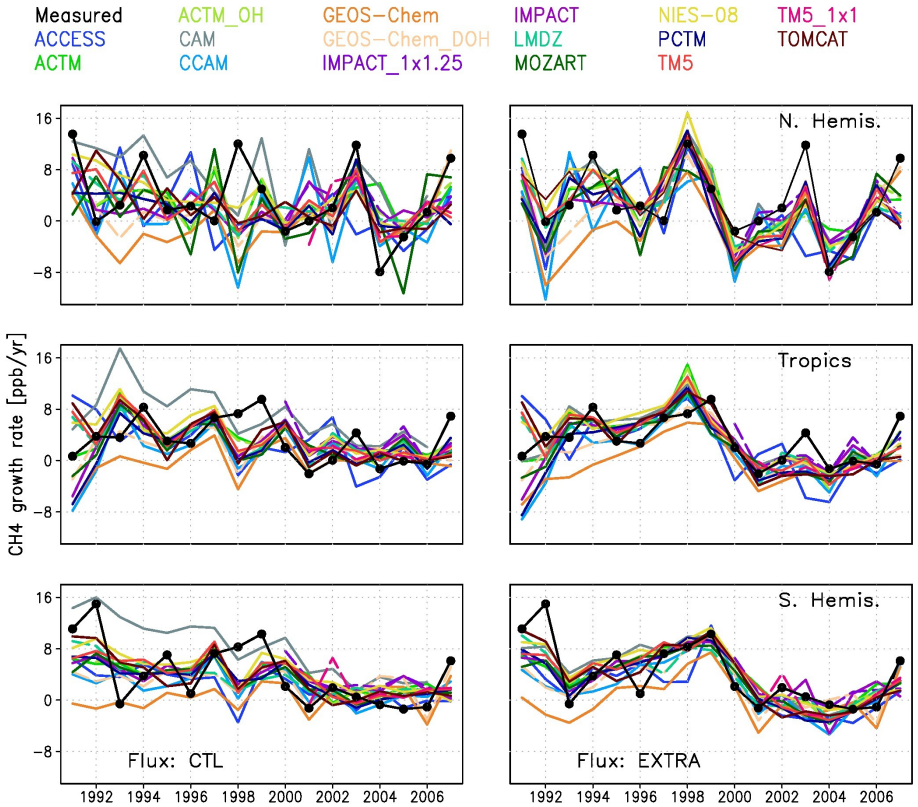


**Fig. 6.** Comparisons of observed and simulated annual mean growth rates of $CH_4$ in NH (top row; ALT, BRW and MHD average), tropics (middle row; MLO RPB and SMO average) and SH (bottom row; CGO and SPO average) for two selected fluxes, CTL (left column) and EXTRA (right column). Growth rate variabilities corresponding 4 other $CH_4$ tracers are given in Figs. S20 and S21.





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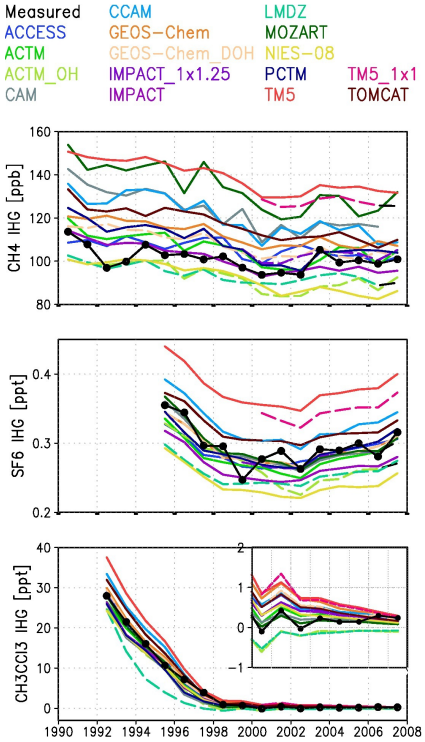


**Fig. 7.** Interhemispheric gradients (IHGs) for $CH_4$, $SF_6$ and $CH_3CCl_3$ concentrations between the NH (BRW, MLO) and SH (CGO, SPO) sites. The values at CGO (AGAGE) are adjusted to NOAA scales by adding an offset of 0.02 ppt for $SF_6$, and multiplied by 1.0003 and 1.0333 for $CH_4$ and MCF, respectively (see text in Sect. 2.4 for further details). Please note that adjustment of the AGAGE data to NOAA scale is made just for convenience. These 4 sites are chosen here because their data coverage is most complete during 1990–2007. Haley Bay (75.58° S, 26.5° W, 10 m) site is chosen for PCTM due to no SPO data in all files. Inset shows expanded *y*-axis view of MCF for the 2000–2007 period.





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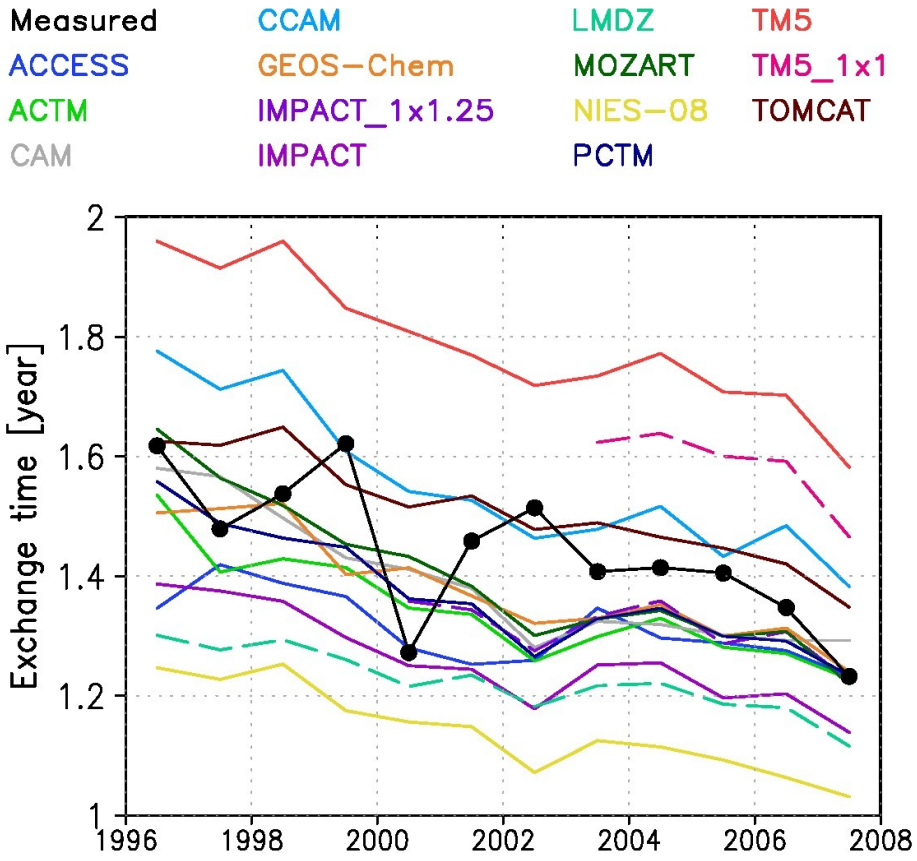


**Fig. 8.** IH exchange time ($\tau_{ex}$) estimated using the measured and simulated time series of average $SF_6$ in NH (BRW, MLO) and SH (CGO, SPO) by employing Eq. (4).





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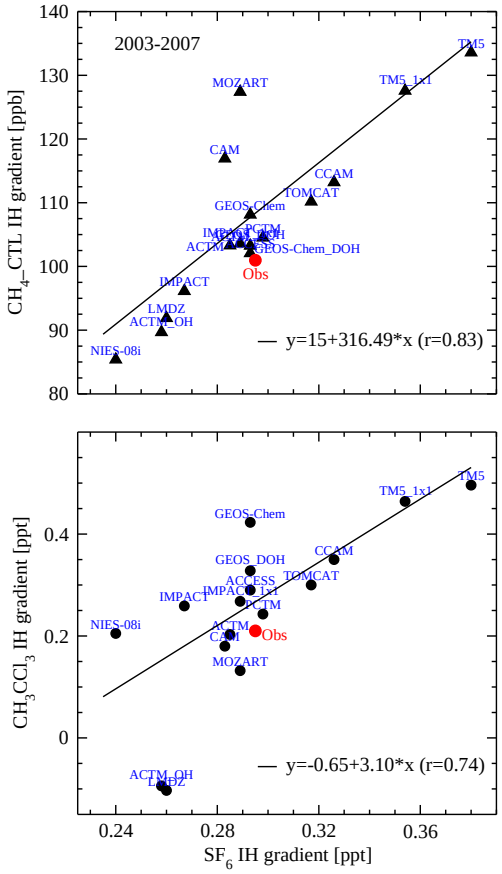


**Fig. 9.** Correlation between IH gradients of $SF_6$ with that of $CH_4$ and $CH_3CCl_3$ are depicted, using average values for the period 2003–2007 and for 4 sites as in Fig. 7. ACTM_OH is excluded here because the $SF_6$ emissions are not as per the protocol.





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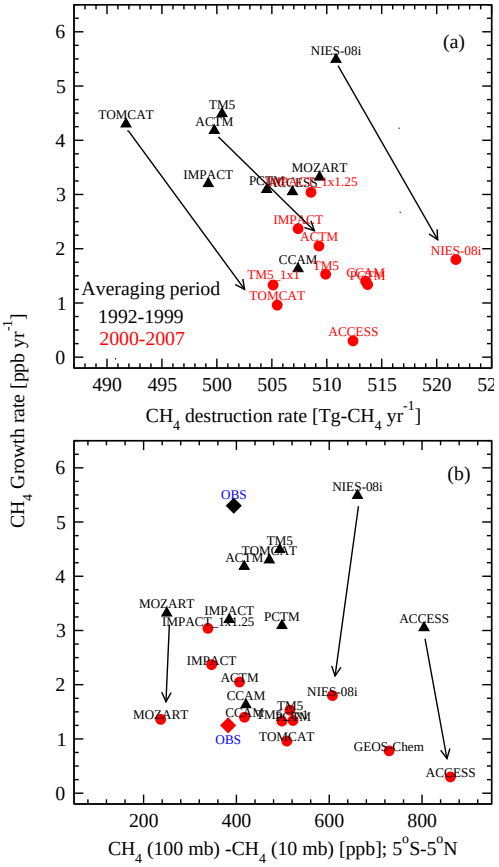


**Fig. 10.** Relationships of global loss rates for different models ($CH_4$_EXTRA) with their simulated growth rates at surface sites **(a)**, and the modeled vertical gradients in the lower stratosphere (100–10 mb layer; **b**) with growth rate. ACTM_OH and GEOS-Chem_DOH are excluded from this analysis as those models used different OH fields.





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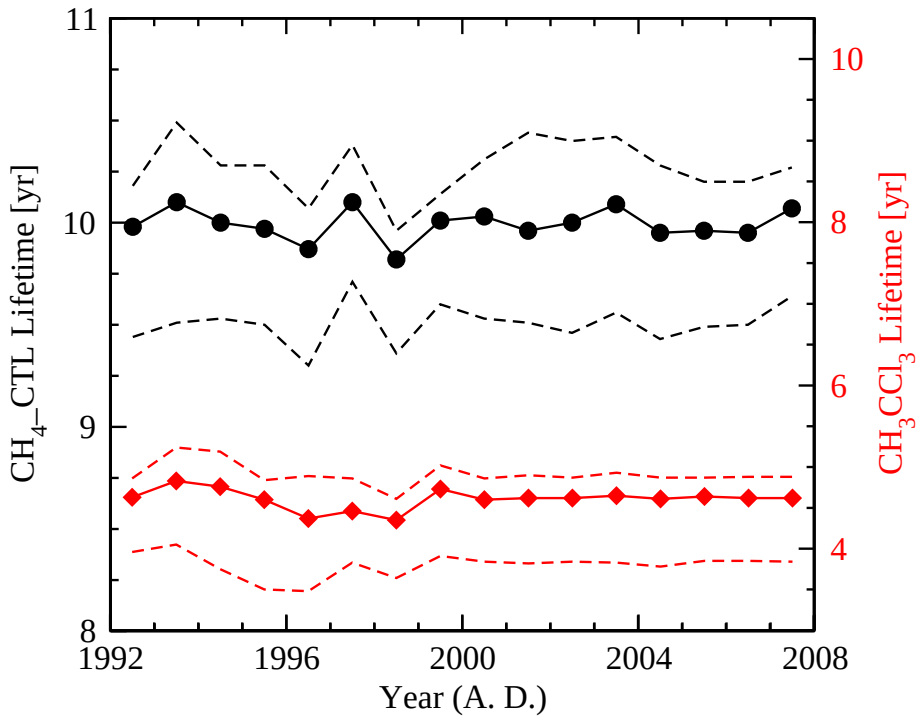


**Fig. 11.** Time series of median (symbols) and ranges (broken lines) of atmospheric lifetimes of $CH_4$ (black) and $CH_3CCl_3$ (red). Note that the mean $CH_4$ lifetimes for EXTRA ($9.96 \pm 0.08$ yr) and CTL ($9.99 \pm 0.08$ yr) fluxes agree within their interannual variability. A comparison of $CH_3CCl_3$ lifetimes calculated using Eq. (5) and gridded photochemical destruction rates is shown in Fig. S22 (ACTM only).





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