Technical note: The CAMS greenhouse gas reanalysis from 2003 to 2020

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Abstract. The Copernicus Atmosphere Monitoring Service has recently produced a greenhouse gases reanalysis (version egg4) that covers almost two decades from 2003 to 2020 and will be extended in the future. This reanalysis dataset includes carbon dioxide (CO₂) and methane (CH₄). The reanalysis procedure combines model data with satellite data into a globally complete and consistent dataset using the European Centre for Medium-range Weather Forecasts' Integrated Forecasting System (IFS). This dataset has been carefully evaluated against independent observations to ensure validity and point out deficiencies to the user. The greenhouse gas reanalysis can be used to examine the impact of atmospheric greenhouse gases concentrations on climate change, such as global and regional climate radiative forcing, assess intercontinental transport, and also serve as boundary conditions for regional simulations, among other applications and scientific studies. The caveats associated with changes in assimilated observations and fixed underlying emissions are highlighted, as well as their impact on the estimation of trends and annual growth rates of these long-lived greenhouse gases.

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1 Introduction

Atmospheric carbon dioxide (CO₂) and methane (CH₄) are the most abundant man-made greenhouse gases directly responsible for climate change (IPCC, 2021). Their long lifetime and increasing anthropogenic emissions near the surface account for their long-term trends (Friedlingstein et al., 2021). A lot of effort has been devoted to measuring the atmospheric concentrations from ground-based observatories (e.g. National Oceanic and Atmospheric Administration (NOAA), gml.noaa.gov; Integrated Carbon Observation System (ICOS), www.icos-cp.eu), which provide the gold standard for the estimation of trends, and more recently satellite data (Committee on Earth Observation Satellites (CEOS), Crisp et al., 2018) enhancing the spatial coverage of greenhouse gas observations at global scale. Atmospheric measurements also sample the variability of CO₂ and CH₄ coming from the weather and its associated atmospheric transport (e.g. Patra et al., 2008, 2011). For this reason, Numerical Weather Prediction (NWP) models have been extensively used to represent and reconstruct the variability of atmospheric concentrations of various tracers (e.g. Inness et al., 2019). Here we use the Integrated Forecasting System (IFS) of the European Centre for Medium-range Weather Forecasts (ECMWF) which has been adapted to include CO₂ and CH₄ in the weather forecast (Agustí-Panareda et al., 2017, 2019) to create a greenhouse gases (GHG) reanalysis. The reanalysis uses the data assimilation technique to combine CO₂ and CH₄ satellite data from the SCanning Imaging Absorption spectroMeter for Atmospheric CHartographY (SCIAMACHY, www.sciamachy.org), the Infrared Atmospheric Sounding Interferometer (IASI, www.eumetsat.int/iasi) and The Thermal and Near Infrared Sensor for Carbon Observation (TANSO, www.eorc.jaxa.jp/GOSAT/instrument 1.html) instruments with IFS model simulations of CO₂ and CH₄ (Agustí-Panareda et al., 2022). The dataset is based on a consistent and stable model version to provide a homogenous, continuous and gapless record of the CO₂ and CH₄ in the entire atmosphere since 2003.

The IFS includes a forecasting model and a data assimilation system combined. The data assimilation system also integrates meteorological observations as in the fifth generation of ECMWF meteorological reanalyses, ERA5 (Hersbach et al., 2020), to best constrain the atmospheric variability of greenhouse gases (Massart et al., 2014, 2016). The forecasting model provides a 3-dimensional representation and evolution of the atmospheric CO₂ and CH₄ and meteorological variables (Agustí-Panareda et al., 2019). At the model surface the greenhouse gases are forced by a set of surface fluxes and emissions. Such modelling configuration allows to produce a realistic representation of the spatio-temporal variability of greenhouse gases in the atmosphere over a wide range of scales from hours to seasons and from local to global (Agustí-Panareda et al., 2022).

Figure 1 showcases the global evolution of CO₂ and CH₄ represented by the CAMS GHG reanalysis data set over the period 2003-2020 and the span of the used satellite data. The seasonal averages illustrate the spatial and temporal variability information contained in the reanalysis dataset which can be exploited for a range of applications in atmospheric sciences. A key potential use of the CAMS GHG reanalysis is to assess the impact of greenhouse gases on climate change. The reanalysis 3-dimensional fields could be used to investigate global and regional climate radiative forcing (e.g. atmosphere.copernicus.eu/climate-forcing), serve as boundary conditions for regional simulations, assess intercontinental

- transport, and generally provide a reference for any other study focusing on atmospheric variability of CO₂ and CH₄. However, care should be taken when using the CAMS GHG reanalysis to estimate trends and annual growth rates of these long-lived greenhouse gases by considering the caveats associated with the changes in the satellite retrievals of CO₂ and CH₄ and the fact that neither anthropogenic emissions nor natural fluxes are adjusted by the data assimilation system, unlike atmospheric inversions (e.g. Chevallier et al., 2019).
- The objective of this technical report is to document the technical aspects of the method and input data used to produce the CAMS GHG reanalysis, and to provide guidance to potential users on the strengths and limitations of the dataset. Section 2 describes the processing chain to produce the reanalysis and its components. Section 3 focuses on the evaluation of the CAMS GHG reanalysis using independent observations from the TCCON and NDACC networks as well as surface in situ networks and AirCore profiles. A list of limitations and caveats of the CAMS GHG reanalysis associated with the changes in the assimilated data and the underlying model errors are compiled in Sect. 4. Finally, Sect. 5 provides a summary and outlook for future CAMS GHG reanalyses.

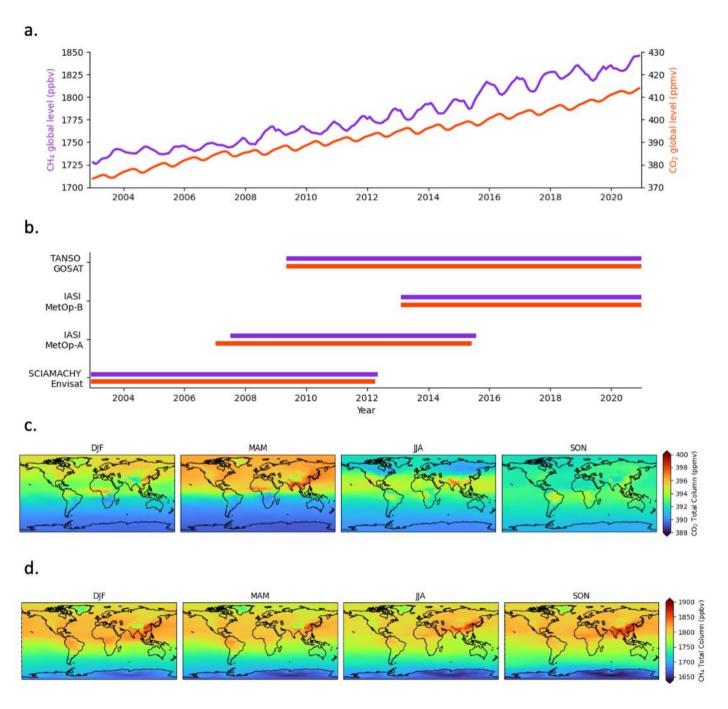


Figure 1. (a) Reanalysis timeseries of global column-averaged CO_2 (red) and CH_4 (purple) atmospheric mole fractions (global mean error ranges from -0.7 to +3.5ppm based on evaluation in section 3.3); (b) the span of the satellite data records for the corresponding species; (c) CO_2 and (d) CH_4 seasonal total column averages (DJF: December-January-February, MAM: March-April-May, JJA: June-July-August, SON: September, October, November) for the 2003-2020 period illustrate the typical seasonal cycle. Note that individual years can be affected by the large inter-annual variability of biogenic fluxes (e.g. during el Niño years).

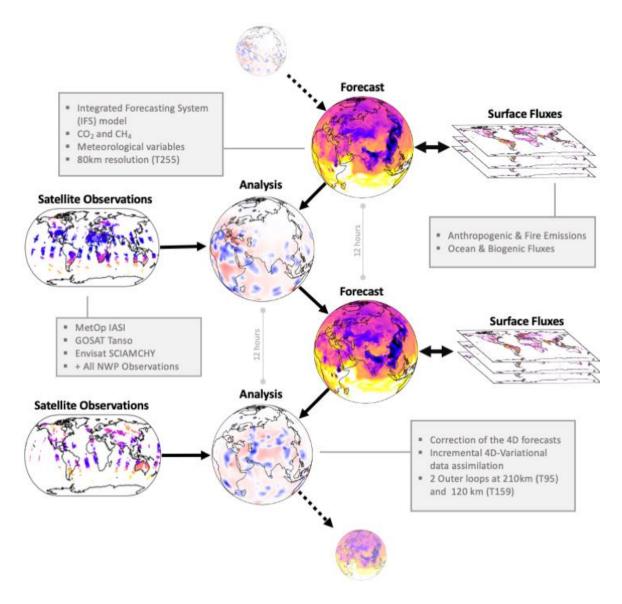


Figure 2. Schematic of the reanalysis cycling procedure. The flow diagram shows the steps and elements combined in the reanalysis. Surface fluxes are used as boundary condition for the atmospheric forecasts. Satellite data are combined with the forecast using data assimilation to produce an analysis (corrected 4D fields) to initialise the next forecast.

2 Methods

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This section gives an overview of the different building blocks of the CAMS GHG reanalysis and the processing chain that integrates the different components to produce the reanalysis dataset.

2.1 The reanalysis cycling chain

The reanalysis production chain is illustrated in Fig. 2. It is a cycling procedure based on a 12-hour data assimilation window that involves four main parts:

- Satellite retrievals of CO₂ and CH₄ (see section 2.2) as well as NWP observations (Hersbach et al., 2020).
- Surface fluxes (see section 2.3) that constitute the sources and sinks of CO₂ and CH₄ in the atmosphere are compiled from various sources. They provide the surface boundary condition for the tracer transport model.
- A model forecast (see section 2.4) that provides a 4-dimentional representation of the state of the greenhouse gases over space and time, along with other meteorological variables, during the 12-hour analysis window (from 09:00 to 21:00 and 21:00 to 09:00 UTC). The forecasts are initialised with the previous analysis, except for the first forecast for the initial date, which is initialised with atmospheric molar fractions from the CAMS inversion dataset (Chevallier et al., 2020; Segers et al., 2020a).
- The above elements are combined using a data assimilation system (see section 2.5) to produce an analysis (Massart et al., 2014, 2016). The analysis will serve to initialise the following forecast over the subsequent 12 hourly cycle.
- Details of these four different components of the reanalysis processing chain are provided in the subsections below, as well as the approach followed to monitor the assimilation of CO₂ and CH₄ satellite data.

2.2 Satellite GHG observations

The satellite measurements of radiances (L1 data) are processed by satellite retrievals developed by various data providers to derive information on the total and partial atmospheric column of CO₂ and CH₄ dry mole fraction (L2 data). In the CAMS GHG reanalysis only L2 products were used for CO₂ and CH₄. With nadir looking satellite instrument geometries the L2 data provide vertically integrated content with vertical sensitivity functions called either averaging kernel when an optimal estimation approach (Rodgers, 2000) is used or weighting functions, that provide information on where the retrieval sensitivity is located along the vertical. The satellite products assimilated in this reanalysis are all provided with averaging kernel and prior information or weighting functions (Massart et al, 2014, 2016). The rationale for selecting the CO₂ and CH₄ satellite products is based on the availability of operational data in near-real time as the strategy is to extend the CAMS GHG reanalysis to the present by eventually running it close to real time. Table 1 provides the specification for each of the assimilated satellite CO₂ and CH₄ products, selected as the state-of-the art retrievals at the beginning of 2017, when the CAMS GHG reanalysis production started. All of the L2 satellite products are freely available from the Copernicus Climate Change Service

- 125 (C3S) Copernicus Climate Data Store (Alos et al., 2019) at https://cds.climate.copernicus.eu/cdsapp#!/dataset/satellite-methane for CH₄. The GHG reanalysis integrate the L2 GHG data from the following satellite instruments:
- SCIAMACHY Envisat: The The SCanning Imaging Absorption spectroMeter for Atmospheric CartograpHY (SCIAMACHY) instrument onboard the Envisat satellite was launched by the European Space Agency (ESA) in 130 March 2002 and it was developed by a consortium involving the Netherland Space Office, the German Aerospace Centre and the Belgian Federal Science Policy Office. It measures radiances variations from the ultraviolet to the near visible infrared. The GHG L2 products use the nadir spectra of reflected and scattered solar radiation in the nearinfrared region. Satellite radiance observations in the near infrared spectral region with the nadir looking geometry are sensitive to changes in CO₂ and CH₄ down to the Earth's surface. The measurements provide total column information with sensitivity peaking near the surface. The ground pixel size is typically between 30 km and 60 km 135 and the swath width is about 960 km. There are no across-track gaps between the ground pixels but there are gaps along-track as SCIAMACHY operates only part of the time (approx. 50%) in nadir observation mode. The CO₂ and CH₄ column products are retrieved by the University of Bremen (Reuter et al., 2011) and the Netherland Institute for Space Research (SRON) (Frankenberg et al., 2011), respectively. Both of L2 products are delivered by the ESA GHG-140 Climate Change Initiative (Buchwitz et al, 2015) and the C3S Climate Data Store (https://cds.climate.copernicus.eu).
 - TANSO-FTS GOSAT: The Thermal And Near infrared Sensor for carbon Observations Fourier Transform Spectrometer (TANSO-FTS) instrument onboard the Greenhouse Gases Observing Satellite (GOSAT) satellite has been developed by the Japan Aerospace Exploration Agency (JAXA) and it was launched in January 2009. TANSO-FTS measures radiances in the short-wave infrared band that provide information of total-column CO₂ and CH₄ mole fractions. Similar to SCIAMACHY, the sensitivity of the total column information provided by L2 data is peaking near the surface due to the spectral band used. The ground pixel size is about 10 km, the swath is 750 km and it has a revisit time of 3 days. In contrast to SCIAMACHY, the GOSAT scan pattern consists of non-consecutive individual ground pixels, i.e., the scan pattern is not gap-free. For a general overview about GOSAT see also http://www.gosat.nies.go.jp/en/. The L2 retrieval product is engineered by the SRON (Schepers et al., 2012, 2016) and delivered by the ESA GHG-CCI and the C3S Climate Data Store (https://cds.climate.copernicus.eu).

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• IASI – Metop A and B: The Infrared Atmospheric Sounding Interferometer (IASI) instruments are onboard the Meteorological Operational satellites (Metop-A and Metop-B) launched in October 2006 and September 2012 respectively. The French National Centre for Space Studies (CNES) lead the design and developments of the instruments in collaboration with the European Organisation for the Exploitation of Meteorological Satellites (EUMETSAT). The IASI instruments measure the thermal infrared band with high spectral resolution enabling it to detect a wide range of trace gas variations in the atmosphere, including CO₂ and CH₄ sensitive in the mid and upper tropospheric regions between 5 and 12 km of altitude. IASI is an across track scanning system with a swath width of 2200 km, providing global coverage twice a day. The field of view is sampled by 2×2 pixels whose ground resolution

is 12 km at nadir. Both CO₂ and CH₄ are engineered and delivered by the Centre National de Recherche Scientifique (CNRS)-Laboratoire de Météorologie Dynamique (LMD) (Crevoisier et al., 2009a, 2009b, 2014). The two L2 products are delivered by the ESA GHG-Climate Change Initiative (Buchwitz et al, 2015) and the C3S Climate Data Store (https://cds.climate.copernicus.eu).

165 Table 1. Specifications of the satellite data used in the CAMS GHG reanalysis

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Gas Instrument - Period assimilated Ve		Version (data provider)	Reference	Peaking sensitivity	
Satellite					
SCIAMACHY –	20030101 - 20120324	CO2_SCI_BESD (v02.01.02, IUP-UB)	Reuter et al.,	Near Surface	
Envisat			2011)		
IASI – Metop-A	20070701 - 20150531	CO2_IAS_NLIS (v8.0, CNRS-LMD)	Crevoisier et al.	Middle and Upper troposphere	
IACI Matau D	20120201 20191120	CO2 IAC NI IC (24 2 224 CNDC I MD)	(2009a)	* *	
IASI – Metop-B		,		Middle and upper	
		· ·		troposphere	
TANSO-FTS -	20090601-20131231	CO2_GOS_SRFP (V2.3.6, SRON)	Butz et al.,	Near Surface	
GOSAT	20140101-20181231	CO2_GOS_SRFP(V2.3.8, SRON)	(2011); Guerlet		
	20190101-20201231	CO2_GOS_BESD (CAMS_NRT, IUP-UB)	et al. (2013);		
			Heymann et al.		
			(2015)		
SCIAMACHY -	20030108-20100601	CH4_SCI_IMAP (v7.2, SRON)	Frankenberg et	Near Surface	
Envisat			al., (2011)		
IASI – Metop-A	20070701-20150630	CH4_IAS_NLIS (V8.3, CNRS-LMD)	Crevoisier et al.,	Middle and Upper	
			(2009b, 2014)	troposphere	
IASI – Metop-B	20130201- 20181130	CH4_IAS_NLIS (V8.1_nrt, CNRS-LDM)		Middle and upper	
	20181201-20201231	CH4_IAS_NLIS (v4.0_nrt, CNRS-LDM)		troposphere	
TANSO-FTS -	20090601-20131231	CH4_GOS_SRFP (V2.3.6, SRON)	Butz et al.,	Near Surface	
GOSAT	20140101-20181231	CH4_GOS_SRFP (V2.3.8, SRON)	(2010);		
	20190101-20201231	CH4_GOS_SRPR (CAMS_NRT, SRON)	Schepers et al.,		
			2012		
	Satellite SCIAMACHY – Envisat IASI – Metop-A IASI – Metop-B TANSO-FTS - GOSAT SCIAMACHY – Envisat IASI – Metop-A IASI – Metop-A	Satellite SCIAMACHY – Envisat 20030101 – 20120324 IASI – Metop-A 20070701 - 20150531 IASI – Metop-B 20130201 – 20181130 20181201-20201231 2018001-20131231 GOSAT 20190101-20181231 20190101-20201231 20190101-20201231 SCIAMACHY – Envisat 20070701-20150630 IASI – Metop-A 20130201- 20181130 20181201-20201231 20181201-20201231 TANSO-FTS - GOSAT 20090601-20131231 20140101-20181231 20140101-20181231	Satellite SCIAMACHY – Envisat 20030101 – 20120324 CO2_SCI_BESD (v02.01.02, IUP-UB) IASI – Metop-A 20070701 - 20150531 CO2_IAS_NLIS (v8.0, CNRS-LMD) IASI – Metop-B 20130201 – 20181130 CO2_IAS_NLIS (v4.2_nrt, CNRS-LMD) 20181201-20201231 CO2_IAS_NLIS (v4.0_nrt, CNRS-LMD) TANSO-FTS - GOSAT 20090601-20131231 CO2_GOS_SRFP (V2.3.6, SRON) 20190101-20201231 CO2_GOS_BESD (CAMS_NRT, IUP-UB) SCIAMACHY - Envisat 20030108-20100601 CH4_SCI_IMAP (v7.2, SRON) IASI - Metop-A 20070701-20150630 CH4_IAS_NLIS (V8.3, CNRS-LMD) IASI - Metop-B 20130201- 20181130 CH4_IAS_NLIS (v4.0_nrt, CNRS-LDM) 20181201-20201231 CH4_IAS_NLIS (v4.0_nrt, CNRS-LDM) TANSO-FTS - 20090601-20131231 CH4_GOS_SRFP (V2.3.6, SRON) GOSAT 20140101-20181231 CH4_GOS_SRFP (V2.3.8, SRON)	Satellite SCIAMACHY - 20030101 - 20120324 CO2_SCI_BESD (v02.01.02, IUP-UB) Reuter et al., 2011)	

2.3 Surface fluxes and prescribed sources/sinks

The emissions and surface fluxes provide the surface boundary conditions for the atmospheric concentrations of CO₂ and CH₄. They play a crucial role in determining the variability and growth rate of both greenhouse gases in the atmosphere. Errors in the budget of the total flux will result into systematic errors or biases in forecast of atmospheric CO₂ and CH₄. In the CAMS reanalysis the surface fluxes (including sources and sinks) are not optimised by the assimilation system. This lack of surface

flux optimization can lead to biases in the analysis when the observing system coverage is sparse in space and time or when the observation error is large, and the analysis is strongly influenced by the model forecast.

175 Table 2 lists the datasets used to produce the CAMS reanalysis and Fig 3 shows the seasonal cycle and trend of the global mean values of each type of surface flux used in the simulations. They include:

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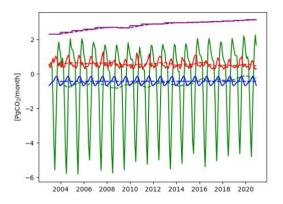
- Fire emissions derived using the CAMS Global Fire Assimilation System (GFAS) version 1.2 that assimilate fire
 radiative power observations from satellite-based sensors (Kaiser et al., 2012). GFAS produces daily estimates of
 wildfire and biomass burning emissions. The emissions are injected at the surface and distributed over the boundary
 layer by the model's convection and vertical diffusion scheme.
- Anthropogenic emissions from the Emission Database for Global Atmospheric Research (EDGAR) version 4.2FT2010 inventory (Janssens-Maenhout et al., 2011; Olivier and Janssens-Maenhout, 2012) excluding the short carbon cycle. The anthropogenic emissions are based on annual average values and include fossil fuel combustion and leakage, agricultural, landfill/waste emissions and aviation (based on the Atmospheric Chemistry and Climate Model Intercomparison Project (ACCMIP, Lamarque et al., 2013) nitric oxide (NO) emissions from aviation scaled to the annual CO₂ total emission from aviation from EDGAR). EDGAR produces global anthropogenic emissions for both CO₂ and CH₄ at a relatively high resolution of 0.1 degrees (compared to 80 km resolution of the CAMS re-analysis). The problem with EDGAR is that the latest version available at the time when the CAMS re-analysis started does not extend beyond 2010. Anthropogenic emissions of CO₂ are extrapolated from 2010 to 2014 with the time series of country totals from EDGARv4.3 (Janssens-Maenhout et al., 2016) and from 2015 to 2020, a persistent growth based on the last available year (2014) is applied. CH₄ anthropogenic emissions are fixed with the last year of available gridded data (2010) from 2011 to 2020. Note that CO₂ and CH₄ emissions are not adjusted for the COVID emission reduction in 2020 (Le Quéré et al., 2020).
- Biogenic CO₂ fluxes are based on the online CHTESSEL module (Boussetta et al., 2013) that relates CO₂ biogenic fluxes with radiation, precipitation, temperature, humidity, and soil moisture. CHTESSEL is used in conjunction with the biogenic flux adjustment system (BFAS) that improves the continental budget of CO₂ fluxes by combining information from fluxes estimates by a global flux inversion system (Chevallier et al., 2010), land-use information and the CHTESSEL online fluxes (Agustí-Panareda et al., 2016). The two-way interaction between the atmospheric forecast and the surface fluxes depicts how the forecast influences the surface fluxes and vice-versa, via the coupling of the biogenic fluxes to the atmospheric forecast (via radiation, temperature, humidity and soil moisture) and the influence of the resulting biogenic fluxes on the atmospheric CO₂ forecast.
- Wetland CH₄ monthly mean emissions come from a climatology (1990–2008) based on the LPJ-WHyMe model that is constrained by SCIAMACHY observations (Spanhi et al., 2011).

- A monthly modulation for CH₄ rice emissions is implemented based on the seasonal cycle of Matthews et al. (1991).
- The CH₄ chemical sink is represented by a monthly mean climatological loss rate from Bergamaschi et al. (2009) based on OH fields optimised with methyl chloroform (Bergamaschi et al., 2005; Houweling et al., 1998) and stratospheric radicals from the 2D photochemical Max-Planck-Institute (MPI) model (Brühl and Crutzen, 1993).
- Other sources and sinks include a CH₄ monthly soil sink (Ridgwell et al., 1991), CO₂ and CH₄ annual mean oceanic fluxes (Houweling et al., 1999; Lambert and Schmidt, 1993; Takahashi et al., 2009) and CH₄ monthly mean fluxes from termites (Sanderson, 1996) and wild animals (Houweling et al., 1999).

215 Table 2. Specifications of the emission and surface fluxes used in the CAMS GHG reanalysis

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Gas	Emission/Flux type	Data provider - Version
CO ₂	CO ₂ and CH ₄ fire emissions	GFAS Version 1.2 (Kaiser et al., 2012)
	CO ₂ ocean fluxes	Takahashi Climatology (Takahashi et al., 2009)
	CO ₂ emissions from aviation	Based on ACCMIP NO emissions from aviation scaled to annual total CO2 from EDGAR
		aviation emissions (Olivier and Janssens-Maenhout, 2012)
	CO ₂ ecosystem fluxes bias corrected with	Based on CHTESSEL (modelled online in IFS)
	BFAS	(Boussetta et al., 2013; Agustí-Panareda et al., 2016)
	CO ₂ anthropogenic emissions	EDGARv4.2FT2010 (2003-2010) (Olivier and Janssens-Maenhout, 2012)
CH ₄	CH ₄ total natural emissions	based on EDGARv4.2FT2010 (2003-2010) (Olivier and Janssens-Maenhout, 2012); LPJ-
		HYMN wetland climatology (Spahni et al., 2011);and other natural sources/sinks
		(Matthews et al., 1991; Ridgwell et al., 1999; Houweling et al., 1999; Lambert and
		Schmidt, 1993; Sanderson, 1996).
	CH ₄ chemical sink	Monthly mean climatology of CH4 loss rate from Bergamaschi et al. (2009)
	CH ₄ anthropogenic emissions	EDGARv4.2FT2010 (2003-2010) (Olivier and Janssens-Maenhout, 2012)



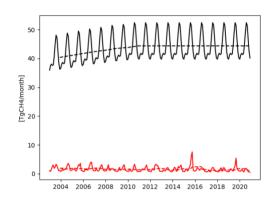


Figure 3. Monthly CO₂ and CH₄ surface fluxes. The CO₂ fluxes [PgCO₂/month] include modelled Net Ecosystem Exchange (NEE) fluxes (in green), anthropogenic emissions (in purple), ocean fluxes (in blue) and biomass burning emissions (in red). The total CH₄ fluxes [TgCH₄/month] excluding biomass burning emissions are shown by black line and CH₄ biomass burning emissions [TgCH₄/month] are depicted in red. The dash lines show the 1-year running mean for each of the fluxes.

2.4 Forecast model

The CAMS GHG reanalysis has been produced using the IFS model. The same model is used to produce operational numerical weather predictions (NWP) at ECMWF and the CAMS global forecast and analyses for reactive gas, aerosols and greenhouse gases at ECMWF (Fleming et al. 2015, Agustí-Panareda et al., 2017, Agustí-Panareda et al., 2022). The IFS model version used is IFS CY42R1, the same as in the CAMS reanalysis for reactive gases and aerosols (Inness et al., 2019). The forecasting model uses a reduced Gaussian grid with a resolution of TL255 corresponding to a horizontal resolution of approximately 80 km and 60 hybrid-sigma pressure vertical levels from the surface to 0.1hPa. The tracer advection is computed using a semiimplicit semi-Lagrangian scheme (Temperton et al., 2001; Diamantakis and Magnusson, 2016) that is not mass-conserving. This scheme leads to an error growth that can dominate the signal in the model simulations if it is not corrected. Thus, a mass fixer is required to ensure mass conservation at every time step (Diamantakis and Agustí-Panareda, 2017). The mass fixer is particularly important for long-lived greenhouse gases for which the interesting signals to monitor, e.g., trends or annual growth rates and large-scale spatial gradients, are weak compared to the large background values. The transport model also includes a turbulent mixing scheme (Sandu et al., 2013) and a convection scheme (Bechtold et al., 2014). For the CH₄ chemical sink in the troposphere and the stratosphere, climatological loss rates derived from the Max Planck Institute photochemical model are al., 2009). Full of **IFS** used (Bergamaschi documentation the be found can at https://www.ecmwf.int/en/forecasts/documentation-and-support/changes-ecmwf-model/ifs-documentation.

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2.5 Analysis procedure (data assimilation)

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The IFS system is using an incremental formulation of the 4-dimentional variational technique (4D-Var). The 4D-Var technique consists of minimizing a cost function that combines the model information and the observation information in order to obtain the best possible state of the atmosphere (analysis) accounting for the model and observation errors. The incremental 4D-Var cost function is quadratic and is formulated as follows:

$$J(\delta x) = \frac{1}{2} (\delta x - \delta x_b)^T B^{-1} (\delta x - \delta x_b) + \frac{1}{2} (G \delta x - d) R^{-1} (G \delta x - d)$$
(1)

where δx is the increment i.e., the difference between the model state x and the first guess x_g , δx_b is the difference between the background (the forecast started from the previous analysis) and the first guess, B the background error covariance matrix, C the observation operator or forward operator that translate the information from model space to observation space. The innovation vector is C0 is added to C1 to provide the analysis.

When the minimization of the cost function is complete, C2 is added to C3 to provide the analysis.

$$x_a = x_a + \delta x \tag{2}$$

The analysis is performed over 12-hour assimilation windows from 9:00 to 21:00 and from 21:00 to 9:00 UTC. The incremental 4D-Var assimilation involves the stepwise minimization of the linearised cost function (equation 1) by updating the first guess x_g and increasing the resolution. In the CAMS reanalysis setup, two minimizations are completed successively at TL95 (approximately 210 km) and TL159 (approximately 110 km) spectral truncations. Once the assimilation procedure is completed an analysis is generated that will serve to initialise the next forecast at the full TL255 resolution.

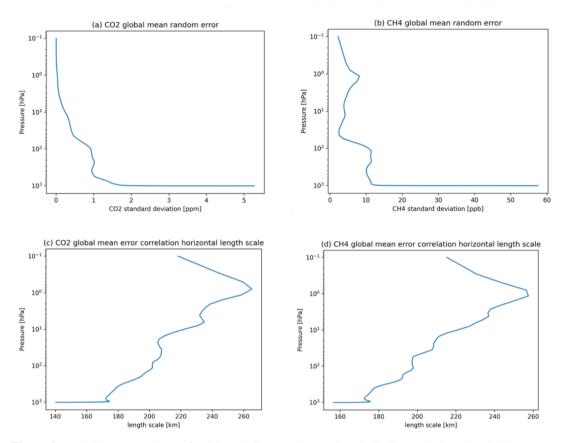


Figure 4. Model background error for CO₂ and CH₄ used in the CAMS GHG reanalysis: (a,b) global mean standard deviation and (c,d) global mean error correlation length scale across the vertical levels.

The background errors for CO₂ and CH₄ were produced from an ensemble of data assimilations (Massart et al., 2016), which allows the calculation of differences between pairs of background fields which have the characteristics of the background errors. The background errors for the greenhouse gas species are univariate, which means that there is no correlation between the greenhouse gas species and the dynamical fields. Hence each species is assimilated independently from the others. The background errors used for both the greenhouse gas species and the dynamical fields are also constant in time. In the ECMWF data assimilation system, the background error covariance matrix is given in a wavelet formulation (Fisher, 2004, 2006). This allows both spatial and spectral variations of the horizontal and vertical background error covariances globally. Figure 4 shows the global mean of the standard deviation and average horizontal correlation length scales for both CH₄ and CO₂. Following experimentation, the correlation length scales between the background errors were manually reduced in the atmospheric boundary layer (1km from the surface).

2.6 Monitoring the data assimilation system

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The time series of the departures (or differences) between the analysis (AN) and the assimilated satellite data (hereafter referred to as observations, OBS), as well as those between the underlying model simulation (or background, BG) and the observations, are used to monitor the performance of the analysis system and are shown in Figures 5 (for CO₂) and 6 (for CH₄). For each satellite retrieval product, both the BG departures (OBS-BG, green lines) and the AN departures (OBS-AN, red lines) are plotted (panel a: SCIAMACHY, panel b: IASI-A; panel c: IASI-B; panel d: GOSAT), together with the number of observations assimilated monthly (blue lines). Overall, both the random (i.e., standard deviation, dashed lines) and the systematic components of the departures (i.e., average values, solid lines) are shown to be reduced by the assimilation process, as highlighted by the AN departures (red lines) being closer to zero than the BG departures (green lines). Note that the difference between the BG and the AN departure is equal to the analysis increments associated with the related observations (i.e. AN-BG).

The number of observations assimilated is different for each satellite instrument and varies with time: IASI generates the largest number of data, with both instruments (IASI-A and IASI-B) providing between 150 000 and 200 000 XCO₂ or XCH₄ data per month; the observations taken by SCIAMACHY oscillate between 25 000 and 50 000 for CH₄ and between 5 000 and 10 000 for CO₂; the number of GOSAT XCO₂ data fluctuate around 2 500, whereas those from GOSAT XCH₄ are comprised between 5 000 and 10 000 per month. It is also clear from Figs 5(a,d) and 6(a,d) that fewer XCO₂ and XCH₄ data from SCIAMACHY, IASI and GOSAT-TANSO are assimilated during the winter months. A magenta vertical dashed line in Figs 5(c,d) and 6(c,d) indicates when the near-real time satellite products started to be assimilated in early 2019. This transition produced an abrupt change in the quality and availability of both IASI and GOSAT retrievals.

The modelled XCO₂ is systematically larger than the observations (leading to overall negative BG departures) because of the biases in the total fluxes (see section 2.3). Therefore, all instruments produced negative departures until 2013 (Fig. 5). From 2013 to 2019, the modelled values of XCO₂ became smaller than those measured by GOSAT (Fig. 5(d)), while the model continued to (slightly) overestimate the IASI XCO₂ observations in the mid to upper troposphere. This overestimation is consistent with a drift in the IASI CO₂ data towards a growing negative bias. After 2018, part of the drift is due to the fact that IASI (version v4.0) is saturating with increasing atmospheric CO₂. Note that this has been corrected with v9.1 (available on the C3S datastore). A sudden change in the IASI-B XCO₂ departures is visible in Fig. 5(c) around December 2018, in correspondence of the switch from the ESA-CCI reprocessed dataset to a near-real time LMD dataset used operationally in the CAMS GHG analysis. The transition to a new dataset was made necessary as the reanalysis production was running close to real-time and reprocessed observations were not available. After the transition to near-real time observations, the IASI XCO₂ increments are reduced to almost zero, as hinted by the overlap between the red (AN departure) and green line (BG departure)

in Fig. 5(c). At the same time, a drop in the number of assimilated IASI XCO₂ observations is observed (blue line, same panel and figure). Together with a drastic reduction in the magnitude of the increments, a large negative bias of approximately 5ppm in both the AN and BG departures emerges. This degradation in the quality of the IASI-B XCO₂ observations in the near-real time dataset is due to the change of the correction of the non-linearity of the detector of IASI-B that was made by CNES and EUMETSAT on August 17th 2018 and that introduced a bias of ~0.2 K on the channels used to perform the CO₂ retrieval. This change has been corrected in the versions of IASI-B MT-CO₂ that are available on the C3S datastore but were not used for this reanalysis. In January 2019, there was also a transition from the ESA-CCI GOSAT XCO₂ retrievals to the near-real time IUP-UB retrieval product (Heymann et al., 2015; Massart et al. 2016) as shown in Fig. 5(d). Consequently, the standard deviation of both the AN and the BG departures increases (cf. dashed lines, same panel and figure), suggesting that the near-real time data is noisier than the reprocessed dataset from ESA-CCI.

The mean XCH₄ departures (both AN and BG) of SCIAMACHY and IASI are relatively small (a few ppb) compared to GOSAT (up to 10 ppb), throughout the entire time period (see solid red and green lines in Fig. 6). The XCH4 SCIAMACHY data was not used from 9 April 2012 onwards (Fig. 6(a)). The standard deviation of both the AN and BG departures are smaller for GOSAT (around 10 ppb, dashed lines in Fig. 6(d)) than for SCIAMACHY (around 20 ppb, dashed lines in Fig. 6(a)), indicating that GOSAT provides less noisy observations. Similar to what was observed for CO₂, a discontinuity in the mean AN and BG departures of GOSAT XCH₄ emerges in January 2019, in correspondence of the transition from the ESA-CCI dataset and the NRT SRON retrievals (see dashed pink line in Fig. 6(d)). Both the AN and the BG departures change sign, indicating that while up to 2019 both the analysis and model were underestimating the GOSAT observations, they start to overestimate them since 2019. Since there was no modification to the model used for the reanalysis over this period, the cause of this negative bias emerging in both the AN and the BG departures since 2019 can only be attributed to the NRT GOSAT XCH₄ observations, and in particular to the fact that they are generated by using an extrapolated XCO₂ value in the proxy retrieval. In addition, the number of assimilated NRT GOSAT XCH₄ observations approximately doubles (blue line in Fig. 6(d)). Note that the switch to the near-real time retrievals for IASI-B XCH4 has a much more marginal impact on the system (Fig. 6(c)).

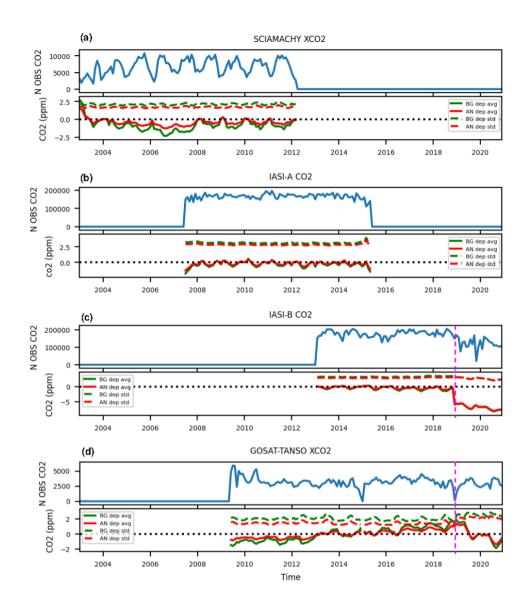


Figure 5. Time series of global monthly number of XCO_2 satellite data (blue) and monthly mean CO_2 analysis (AN) and model background (BG) departures of the various observations (OBS) assimilated in the reanalysis (AN-OBS and BG-OBS in red and green respectively, see legend). The solid lines show the monthly average of the departures, and the dash lines the monthly standard deviations. The magenta dash line indicates the switch to the near-real time satellite products. Note that the range of values in y-axis varies depending on the satellite product.

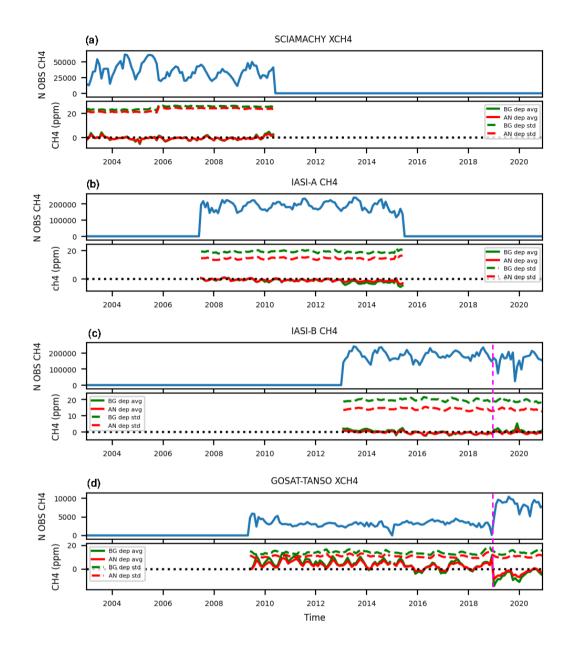


Figure 6. Time series of global monthly number of XCH₄ satellite data (blue) and monthly mean CH₄ analysis (AN) and model background (BG) departures of the various observations (OBS) assimilated in the reanalysis (AN-OBS and BG-OBS in red and green respectively, see legend) for different satellite products. The solid lines show the monthly average of the departures, and the dash lines the monthly standard deviations. The magenta dash line indicates the switch to the near-real time satellite products. Note that the range of values in y-axis varies depending on the satellite product.

3 Evaluation with independent observations

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Validation against a set of independent observations has been performed on the 18 years of the CAMS GHG reanalysis span. The independent data includes different types of observations (see Fig 7): in situ near-surface continuous observations of CO₂ and CH₄ mole fractions from the collaborative ObsPack datasets (Schuldt et al., 2020; Sarra et al., 2021; NOAA Carbon Cycle Group ObsPack Team, 2019; see Table A1); dry-air column-averaged mole fractions from the Total Carbon Observing Network (TCCON, Wunch et al. 2011, 2015); tropospheric and stratospheric partial columns for CH₄ from the Network for the Detection of Atmospheric Composition Change (NDACC, De Mazière et al., 2018) (see Table A2); AirCore vertical profiles of CO₂ and CH₄ mole fractions (Karion et al., 2010; Baier et al., 2021); and the NOAA global mean CO₂ and CH₄ mole fraction product based on the Greenhouse Gas Marine Boundary layer Reference (Conway et al., 1994, Dlugokencky et al., 1994, Massarie et al., 1995).

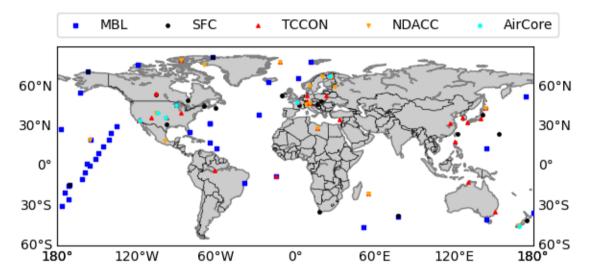


Figure 7 Map with observing sites used in the evaluation of the CAMS GHG reanalysis: MBL (blue squares) includes NOAA Marine Boundary Layer (MBL) reference sites used to compute the NOAA global mean CO2 and CH4 mole fraction product (see https://gml.noaa.gov/ccgg/mbl/mbl.html for further details); SFC (black circles) correspond to the in situ near-surface continuous observations of CO2 and CH4; TCCON and NDACC sites are depicted by red and orange trianges; and AirCore sites are shown by cyan circles.

3.1 Surface and column data

3.1.1 Carbon dioxide

Overall, the error is within ± 10 ppm and ± 4 ppm for most of the near-surface and total column stations respectively for the whole 18-year period (Figs 8 and 9). Near the surface (Fig 8), there is a large variability in the CO₂ error between continental stations influenced by local fluxes (e.g., CDL, FSD, AMT, HUN, see Table A.1) and oceanic stations sampling well-mixed air (ALT, BRW, MHD). Continental stations show large error variations with season (e.g., CDL, HUN), with an underestimation

- of CO₂ in the summer and an overestimation in the winter, indicating an underestimation of the amplitude of the CO₂ seasonal cycle largely driven by vegetation growth. Differences between stations will be determined by the footprint of observations having different contributions of fluxes from different biomes and from anthropogenic emissions. Accuracy of such fluxes can vary geographically.
- Overall, there is positive bias of a few ppm between 2003 and 2015 in the baseline surface stations (e.g. BRW, SMO, SPO) which is consistent with the XCO₂ error at the TCCON sites (Fig. 9). This positive bias decreases from 2007 to 2015 when IASI-A CO₂ data are assimilated, with values lower than 2 ppm, and becomes negative from 2015 to 2019 (from 0 to -2 ppm). From 2019 onwards, there is a positive trend in the bias, and it becomes positive (> +2 ppm) in 2020. There is consistency between the column and surface biases with a general positive bias at background stations before 2015 and a negative bias after 2015 (up to 2019) at the surface stations, although there is no data in 2020 from the surface stations.

The synoptic and large-scale variability of CO_2 is well represented by the reanalysis (lower panel in Fig 9). The root mean square error at TCCON stations is below 0.8 ppm for XCO_2 . The normalised standard deviation is around 1.0 (+/- 0.3) and the Pearson correlation coefficient is larger than 0.8.

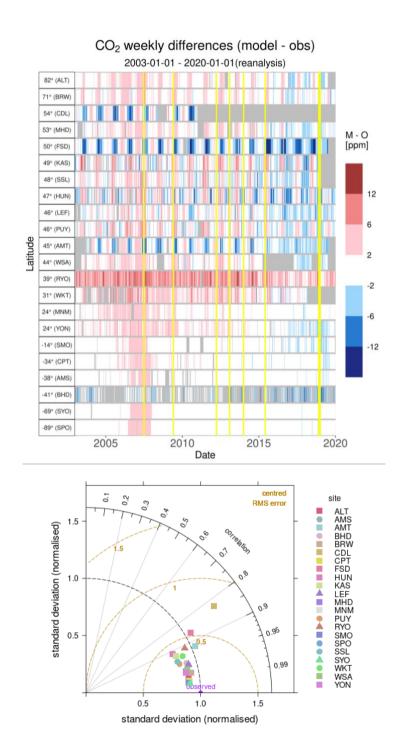
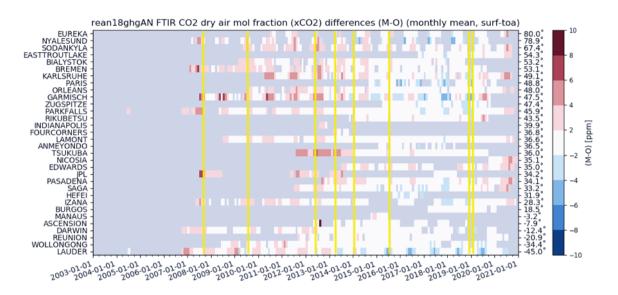
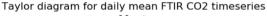
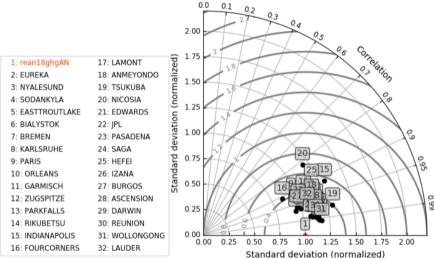


Figure 8. Top: Mosaic plot of CO₂ weekly biases (in ppm) of the CAMS GHG reanalysis compared to surface continuous observations of CO₂ mole fraction obtained from GLOBALVIEWplus CO₂ ObsPack v6.0 (Schuldt et al., 2020) and listed in Table A1. Each coloured vertical line represents a weekly mean. Vertical yellow lines depict the changes in the assimilated data documented in Figs 1, 5 and 6. Grey shading indicates no observations are available. Bottom: Taylor diagrams for the site dependent CO₂ comparison of the CAMS GHG reanalysis against same observations used in top panel. The standard deviation is normalised by dividing the









405 Figure 9. Top: Mosaic plot of the CAMS GHG reanalysis biases at all TCCON sites (see Table A2) for the column-averaged dry mole fraction of CO₂ [ppm] (XCO2) averaged daily around local noon (+/- 2.5 hours). Vertical yellow lines depict the changes in the assimilated data documented in Figs 1, 5 and 6. Grey shading indicates no observations are available. Bottom: Taylor diagrams for the station dependent XCO₂ comparison of the CAMS GHG reanalysis against TCCON FTIR data. The standard deviation is normalised by dividing the observed and modelled time series with the standard deviation of the model time series. The model has higher/lower variability compared to the observed data if the site is plotted with a distance smaller/larger than 1 from the origin.

3.1.2 Methane

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The CH₄ reanalysis fields are generally in good agreement with surface and tropospheric column observations with typical weekly and monthly errors within ±40 ppb and ±25 ppb respectively (Figs 10, 11 and 12). Stratospheric partial columns compared to NDACC data reveal a positive bias that is of the same order as the reported measurement uncertainty of 7% (Fig. 10, upper panel). The averaged relative differences in the troposphere across all NDACC sites are -0.4% for the reanalysis (Fig.11, lower panel), which is well within the measurement's uncertainty. The reanalysis overestimates the column-averaged CH₄ compared to TCCON observations (Fig. 12), for most mid- and high-latitude sites, with a relative difference of up to 2.5%, but shows a good agreement for the low latitude sites at Izaña, Darwin and Wollongong. At the surface the bias is overall positive up to 2007 (Fig. 10). With the introduction of IASI, the biases are reduced. However, with the switch to near-real time satellite data, the bias become negative at all sites reaching values lower than -20 ppb.

Differences between the surface and total column biases stem from the fact that the model suffers from large positive biases above the tropopause (between 100hPa and 10 hPa) of about 80-100 ppb during the months between September and November (Figs 5d and 6d of Verma et al., 2017) which affect the total column average. This stratospheric bias cannot be corrected systematically by CH₄ satellite data from SCIAMACHY, GOSAT and IASI.

For all observations (surface, partial and total columns) CH₄ shows a seasonality in the relative difference between observations and the reanalysis. The magnitude of the difference is site dependent. During local autumn/winter months the relative bias is increased (underestimation) at most surface sites and in the tropospheric columns. This underestimation is also seen in the TCCON time series. In the spring and summer there is an overestimation of CH₄ near the surface and in the total column. These biases are related to errors in the seasonal cycle of surface emissions, most likely from agriculture and wetlands, and the accuracy of the representation of the hydroxyl radical (OH) sink which overall have larger values in the climatology compared to CAMS IFS(CB05BASCOE) atmospheric chemistry model OH (Segers et al., 2020b, Williams et al., 2021). The XCH₄ root mean square error is around 1.4 ppb and the Pearson correlation coefficient is larger than 0.7 for XCH₄ except for

some outliers (Fig. 12, lower panel), indicating a good representation of the synoptic variability, as for XCO₂.

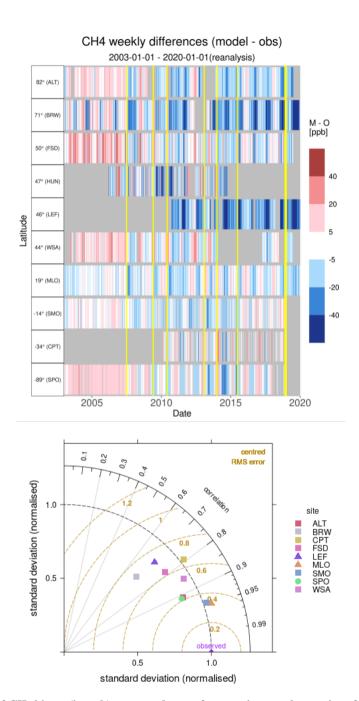
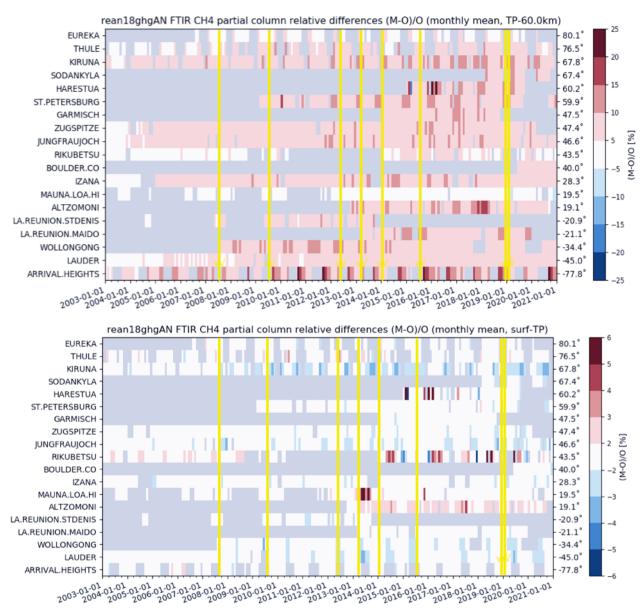
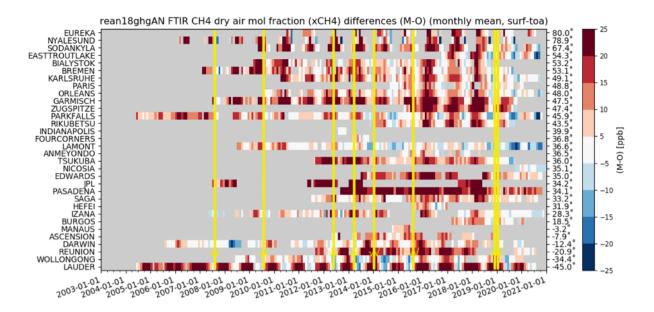


Figure 10. Top: Mosaic plot of CH₄ biases (in ppb) compared to surface continuous observations from GLOBALVIEWplus CH4

440 ObsPack v1.0 data product (Cooperative Global Atmospheric Data Integration Project, 2019) listed in Table A1. Each coloured vertical line represents a weekly mean. Vertical yellow lines depict the changes in the assimilated data documented in Figs 1, 5 and 6. Grey shading indicates no observations are available. Bottom: Taylor diagrams for the site dependent CH₄ comparison of the CAMS GHG reanalysis against same observations used in top panel. The standard deviation is normalised by dividing the observed and modelled time series with the standard deviation of the observations. The model has higher/lower variability compared to the observed data if the site is plotted with a distance larger/smaller than 1 from the origin.



450 Figure 11. Mosaic plot of seasonal relative CH₄ biases at all FTIR sites (see Table A2) for the stratospheric columns (top) and tropospheric columns (bottom) NDACC. Vertical yellow lines depict the changes in the assimilated data documented in Figs 1, 5 and 6. Grey shading indicates no observations are available.



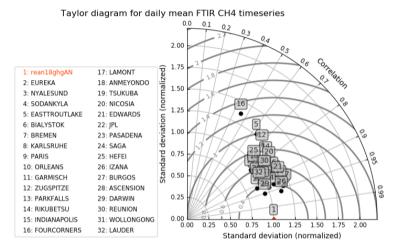


Figure 12 Top: Mosaic plot of monthly biases at all TCCON sites for the column-averaged mole fractions XCH4 [ppb] averaged daily around local noon (+/- 2.5 hours). Vertical yellow lines depict the changes in the assimilated data documented in Figs 1, 5 and 6. Grey shading indicates no observations are available. Bottom: Taylor diagrams for the station dependent XCH4 comparison of the CAMS GHG reanalysis against TCCON FTIR data. The standard deviation is normalised by dividing the observed and modelled time series with the standard deviation of the model time series. The model has higher/lower variability compared to the observed data if the site is plooted with a distance smaller/larger than 1 from the origin.

3.2 Vertical profiles

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The uncertainty of CAMS GHG reanalysis varies with height and the accuracy of the analysis vertical profiles depends mostly on the underlying model uncertainty, as the satellite data assimilated in reanalysis only provide integrated total or partial atmospheric column. The reanalysis has been evaluated using observations of CO₂ and CH₄ vertical profiles (Karion et al., 2010; Baier et al., 2021) from the NOAA AirCore dataset v20210813. It includes 133 vertical profiles from the surface to the lower stratosphere (up to around 40 hPa) from 2012 to 2020 at 7 sites listed in Table A.3.

475 Figure 13 shows that the largest mean error occurs (i) near the surface with a strong influence from surface fluxes; (ii) in Upper Troposphere/Lower Stratosphere (UTLS) region (between 500 hPa and 100 hPa) with a strong influence from longrange transport; and (iii) in the stratosphere (above 100 hPa) where uncertainties associated with chemical loss of CH₄ and the meteorology driving the tracer transport are largest, and the fact that satellite data used here are not able to constrain the stratospheric CO₂ and CH₄ in the reanalysis. Near the surface, there is a positive CO₂ bias associated with an overestimation 480 of the total flux in the model and a negative CH₄ bias which stems from both errors in the emissions and the chemical loss rate in the troposphere. The negative CO₂ bias in the UTLS agree with the tendency of the model to underestimate fine-scale highervalued CO₂ streamers associated with long-range transport. The large positive CH₄ bias in the stratosphere of around 200 ppb is consistent with the positive biases with respect to NDACC stratospheric column (Fig. 11, upper panel) and the documented model biases with respect to MIPAS and ACE-FTS by Verma et al. (2017). The errors associated with the stratospheric 485 chemical sink are thought to be the largest contributor to the stratospheric CH₄ bias as shown by tests using the IFS BASCOE-CB05 chemical loss rate (not shown here). In general, the reanalysis underestimates the CO₂ vertical gradient across the tropopause. This underestimation leads to a positive bias for CO₂ in the lower stratosphere of around 2 ppm. The analysis is not able to remove the large errors near the surface by only adjusting the atmospheric mole fractions, i.e., without adjusting the emissions in the data assimilation process, nor it is able to reduce the stratospheric errors in the model (Massart et al. 2017, 490 Verma et al. 2017). The vertical profiles have a large variability from day to day as shown in Figure 14 with a sequence of profiles at Traînou (France). The CAMS GHG reanalysis is able to capture these synoptic variations in the vertical profile, consistent with its skill to represent XCO₂ and XCH₄ synoptic variability (lower panels of Figs 9 and 12). For a full catalogue of all the individual AirCore vertical profiles used in Fig. 13 see Supplement.

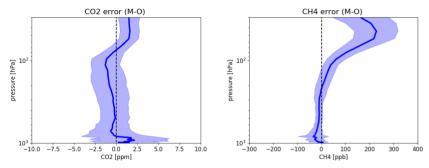


Figure 13: Vertical profiles of mean error (Model M- Observation O) of CAMS CO₂ (left) and CH₄ (right) reanalysis with respect to AirCore observations comprising 133 profiles at 7 sites (listed in Table A3) over the period from 2012 and 2020. The blue shading shows the +/- standard deviation of M-O with respect to the mean error. The vertical dash black line depicts the zero mean error.

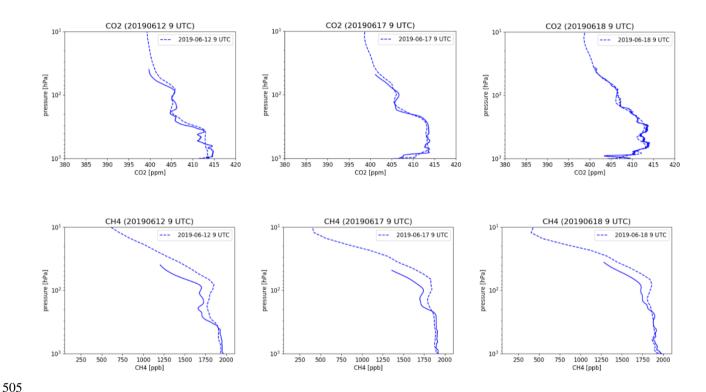


Figure 14: Vertical mole fraction profiles of CO₂ and CH₄ from the CAMS GHG reanalysis (dash line) and AirCore observations (solid line) at Traînou (France, see Table A3) over the period in June 2019.

3.3 Trends

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Although this reanalysis is using a consistent underlying model and re-processed observations of CO₂ and CH₄, the current system is not able to provide accurate enough atmospheric mole fraction that can be used to estimate trends and the atmospheric growth rate as computed by the changes in global mean CO₂ and CH₄ from one year to the next. The CO₂ and CH₄ global annual means based on Marine Boundary Layer (MBL) reference sites are compared to the NOAA Global Greenhouse Gas Reference Network (GGGRN) observations (https://gml.noaa.gov/ccgg/about.html, Andrews et al., 2014; Conway et al., 1994; Dlugokencky et al., 1994) in Fig. 15. Changes in the assimilated satellite data have a clear impact on the evolution of the global annual mean values of CO₂ and CH₄ in the CAMS GHG reanalysis. The reanalysis has a positive global bias in near-surface CO₂ and CH₄ of a few ppm and around 20 ppb respectively from 2003 to 2007. Note that this positive bias in the annual global mean does not imply that the bias will be positive everywhere, as shown by the negative surface CH₄ biases at the AirCore sites (Fig. 13) and the large temporal and geographical variability of the weekly bias illustrated in Figs 8 and 10. After the introduction of IASI in 2007 the global bias decreases and it is lowest during the period when the number of observations is largest in 2013 and 2014 (Figs 5 and 6). Finally, the change to the near-real time satellite retrievals in 2019 together with the incorrect trend in the emissions during the COVID slowdown period in 2020 (Le Quéré et al., 2020) lead to changes in the global bias from negative to positive for CO₂ and from positive to negative for CH₄. These changes in the global bias are consistent with the changes in the errors with respect to total-column and near-surface observations in Figs. 8 to 12. It is important to note that the changes in global bias associated with changes in the assimilated data are of the same order of growth rate of CO₂ (gml.noaa.gov/ccgg/trends) magnitude the observed atmospheric and CH_4 (gml.noaa.gov/ccgg/trends ch4). For this reason, this reanalysis product is not suitable for trend analysis.

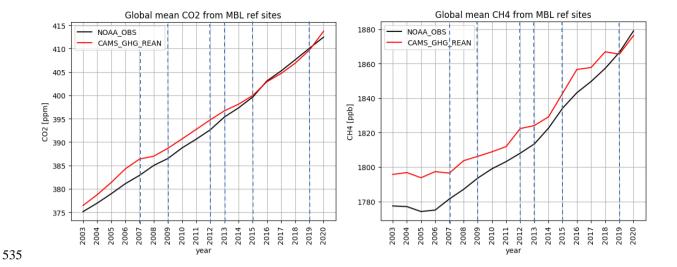


Figure 15: Global mean CO₂ [ppm] and CH₄ [ppb] from the CAMS GHG reanalysis (in red) and the NOAA global mean CO₂ and CH₄ (in black, https://gml.noaa.gov/ccgg/trends/global.html) based on the Greenhouse Gas Marine Boundary layer Reference (Conway et al., 1994, Dlugokencky et al., 1994, Massarie et al., 1995, Dlugokencky et al., 2021). The global mean of the CAMS GHG reanalysis has been computed based on the same NOAA Marine Boundary Layer (MBL) reference sites shown in Fig 7 (see https://gml.noaa.gov/ccgg/mbl/mbl.html for further details). The dash blue lines mark the years when there was a change in the observing system. The uncertainty associated with the computation of global mean using the MBL sites is estimated to be 0.1 ppm for CO₂ (Ed Dlugokencky and Pieter Tans, NOAA/GML, gml.noaa.gov/ccgg/trends/) and below 2 ppb for CH₄ (Ed Dlugokencky, NOAA/GML (gml.noaa.gov/ccgg/trends ch4/).

4. Limitations and caveats

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This section provides an overview of the shortcomings of the CAMS GHG reanalysis which users should consider when interpreting the data. The main issues documented in the previous sections are summarised below:

- 1. Emissions are prescribed and not adjusted by the data assimilation system in the CAMS reanalysis (Sect. 2.3). This leads to a growing model error for CO₂ and CH₄ that can be difficult to correct with a sparse observing system and 12-hour assimilation window. In addition, the prescribed emissions are not available in near-real time, which means they are either kept fixed since the last year available (e.g. 2010 for CH₄) or they are extrapolated with a climatological trend as done for CO₂ (see details in Sect. 2.3). Because of this, the CAMS GHG reanalysis is not suitable to investigate the impact of local emission changes, such as COVID impact studies, which require a large local emission adjustments to the prescribed inventories (e.g. Doumia et al., 2021) and atmospheric inversion systems to estimate the changes (e.g. McNorton et al, 2022).
- 2. Changes of satellite data used with different temporal, horizontal and vertical coverage cause changes in the quality of the reanalysis. For example, winter seasons have a lower number of observations because of light conditions and

the higher frequency of cloud presence. This affects the quality of the seasonal cycle and the inter-hemispheric gradient. Similarly, in regions where there is no observation coverage, such as the stratosphere, the reanalysis is based on the underlying model including its systematic errors (see discussion on stratospheric biases in Sect 3.2).

- 3. Changes in satellite retrievals affect the quality of the observations used in the CAMS GHG reanalysis. For example, the switch from the CCI re-processed satellite products to the near-real time products is associated with a marked change in the bias and random error (i.e. standard deviation) of the departures from XCO₂ and XCH₄ GOSAT observations, as well as in the bias of the departures from the XCO₂ IASI-B observations. This large increase in the bias of the assimilated CO₂ and CH₄ observations from 2019 onwards results into a large increase in the bias of the CAMS GHG reanalysis in 2019 and 2020 which has implications for the trend analysis (Sect. 3.3).
- 4. The fixed climatological chemical loss rate of CH₄ (Sect 2.3) has been shown to overestimate the atmospheric CH₄ chemical sink by Segers et al. (2020b). Preliminary tests coupling the IFS to the atmospheric loss rate derived from BASCOE-CB05 chemistry have indeed shown a large reduction in the CH₄ negative bias in mid-latitudes. Systematic errors in the CH₄ chemical sink used in this reanalysis may have contributed further to enhance the large negative CH₄ bias in the CAMS GHG reanalysis over the last period in 2020, when the increase in the CH₄ growth rate has been linked to a decrease in chemical loss rate (Stevenson et al., 2021).
 - 5. The large CH₄ and CO₂ biases in the stratosphere are currently under investigation. The CH₄ stratospheric bias is mainly associated with the use of a climatological loss rate (Sect 2.3), as preliminary tests using a different chemical loss rate based on IFS CB05BASCOE simulations show that the bias in CH₄ is greatly reduced.
- 6. Changes in systematic errors with time due to model error and changes in observation coverage and quality will affect trend analysis (see Sect. 3.3).

An up-to-date list of the known issues of the CAMS reanalysis can be found in the online CAMS documentation website (https://confluence.ecmwf.int/display/CKB/CAMS%3A+Reanalysis+data+documentation). Some of these issues will also be addressed in the future CAMS reanalysis (planned to start production in 2024), including the improvement of the prescribed emission trends, the consistent use of satellite retrieval products and the use of variable CH₄ chemical loss rate.

5. Summary and conclusions

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This technical report documents the first CAMS IFS reanalysis of CO₂ and CH₄ produced by ECMWF which complements the CAMS reanalysis of reactive gases and aerosols (Inness et al., 2019). The processing chain, assimilated satellite data and underlying model used are described and the resulting reanalysis is evaluated using independent in situ near-surface observations, total column retrievals and in situ atmospheric profile observations. The monthly systematic and random errors of CO₂ and CH₄ typically range within 1% from 2003 to 2020 with an overall good skill in the representation of synoptic spatial variability and seasonal cycle. The lowest systematic errors occur in the period with maximum number of observations

in 2013 and 2014. In 2019 there was a switch from C3S pre-processed satellite products to the near-real time CAMS satellite products because at the time of production the C3S products had not reached 2019. This caused a jump in the quality of the satellite data and the resulting CAMS GHG reanalysis. For this reason, a new re-run of the CAMS GHG reanalysis from 2019 onwards will be performed with consistent C3S satellite products in the near future.

The comparison of global mean values with observations shows variations in the bias that depend on changes in the assimilated satellite data of around 2 ppm and 10 ppb for CO₂ and CH₄ respectively, which have the same magnitude as the observed variations in their growth rate. For this reason, we do not recommend the use of this dataset to study changes in the atmospheric growth rate of CO₂ and CH₄. Similarly, large biases in stratospheric CO₂ and CH₄ should also considered when analysing stratospheric signals and trends in the CAMS GHG reanalyses. A list of caveats and limitations that users need to be aware of is provided in Sect. 4.

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The slow reduction of the lingering bias in the model background is associated with competing factors at play: (i) the error growth in the model background associated with the accumulation of systematic errors in emission and natural fluxes; (ii) the limited coverage of observations in time and space (both horizontal and vertical); (iii) the localised impact of observations associated with a short data assimilation window spanning 12 hours.

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In order to improve the CAMS reanalysis in future releases we recommend the following actions: (i) increase the number and coverage of satellite data assimilated from additional satellite missions such as the Copernicus Sentinel-5 Precursor (S5P), 2 3 (OCO-2, Orbiting Carbon Observatory and https://www.nasa.gov/mission_pages/oco2; OCO-3. https://www.jpl.nasa.gov/missions/orbiting-carbon-observatory-3-oco-3) and Greenhouse gases Observing SATellite-2 (GOSAT-2, https://global.jaxa.jp/projects/sat/gosat2) as well as the latest re-processed satellite products from C3S: (ii) improve the underlying anthropogenic emissions and natural fluxes by using the most recent flux data sources, with particular emphasis on the extrapolation of the prescribed flux data in near-real time; (iii) couple the chemical loss rate with the CAMS reanalysis of chemical species (Inness et al. 2019); and (iv) use the IFS inversion capability (McNorton et al., 2022) being developed within the CoCO2 project (coco2-project.eu) for future re-analyses and explore the possibility of applying a correction to the fluxes in order to match the observed global growth rate.

Table A.1 List of stations with in situ continuous observations of CO₂ and CH₄ from GLOBALVIEWplus CO₂ ObsPack v6.0 and CH₄ ObsPack v1.0 respectively used for the evaluation in Sect. 3.1.

Appendix A

Station, Country	Latitude/Longitu	Elevati	Data Reference
(site name)	de	on	
	[degrees]	[m asl]	
Alert, Canada (ALT)	82.45 62.51W	185	Worthy et al. (2003)
Barrow, AK, USA (BRW)	71.32N 156.61W	11	Peterson et al. (1986)
Candle Lake, Canada (CDL)	53.99N 105.12W	600	Worthy et al. (2003)
Mace Head, Ireland (MHD)	53.33N 9.90W	5	Ramonet et al. (2010)
Fraserdale, Canada (FSD)	49.88N 81.57W	210	Worthy et al. (2003)
Kasprowy Wierch, Poland (KAS)	49.23N 19.98E	1989	Rozanski et al. (2003)
Schauinsland, Germany (SSL)	47.92N 7.92E	1205	Schmidt et al. (2003)
Hegyhatsal, Hungary (HUN)	46.95N 16.65	248	Haszpra et al (2001)
Park Falls, WI, USA (LEF)	45.95N 90.27W	472	Andrews et al. (2014)
Puy de Dôme, France (PUY)	45.77N 2.97E	1465	Lopez et al. (2015); Colomb et al. (2020)
Argyle, ME, USA (AMT)	45.03N 68.68W	53	Andrews et al. (2014)
Sable Island, Canada (WSA)	43.93N 60.00W	5	Worthy et al. (2003)
Ryori, Japan (RYO)	39.03N 141.82E	260	Tsutsumi et al. (2005)
Moody, TX, USA (WKT)	31.31N 97.33W	251	Andrews et al. (2014)
Minamitorishima, Japan (MNM)	24.28N 153.98E	8	Tsutsumi et al. (2005)
Yonagunijima, Japan (YON)	24.47N 123.02E	30	Tsutsumi et al. (2005)
Tutuila, American Samoa (SMO)	14.25S 170.56W	42	Waterman et al. (1989)
Cape Point, South Africa (CPT)	34.35S 18.49E	230	Brunke et al. (2004)
Amsterdam Island, France (AMS)	37.80S 77.54E	55	Ramonet et al. (1996)
Baring Head Station, New Zealand (BHD)	41.41S 174.87E	85	Stephens et al. (2013)
Syowa Station, Antarctica, Japan (SYO)	69.01S 39.59E	14	Schuldt et al. (2020)
South Pole, Antarctica, USA (SPO)	89.98S 24.8W	2810	Conway et al. (1990)

Table A.2 List of total column stations used for the evaluation in Sect 3.1.

Station, country	Latitude/	Network	Data references
	Longitude		
	[degrees]		
Eureka, Canada	80.05N	TCCON+NDACC	Strong et al. (2019); Batchelor et al. (2009)
	86.42W		
Ny Ålesund,	78.9N	TCCON+NDACC	Notholt et al., (2019)
Norway	11.9E		
Thule, Greenland	76.53N	NDACC	Hannigan et al. (2009)
	68.74W		
Kiruna, Sweden	67.84N	NDACC	Bader et al. (2017)
	20.41E		
Sodankylä, Finland	67.37N	TCCON+NDACC	Kivi et al. (2014); Sha et al. (2021)
	26.63E		
Harestua, Norway	60.2N	NDACC	De Mazière et al. (2018)
	10.8E		
St Petersburg,	59.90N	NDACC	Makarova et al. (2015)
Russia	29.80E		
East Trout Lake,	54.35N	TCCON	Wunch et al. (2018)
Canada	104.99W		
Bialystok, Poland	53.23N	TCCON	Deutscher et al. (2015)
	23.02E		
Bremen, Germany	53.1N	TCCON	Notholt et al. (2014)
	8.85E		
Karlsruhe, Germany	49.1N	TCCON	Hase et al. (2015)
	8.44E		
Paris, France	48.85N	TCCON	Te et al. (2014)
	2.36E		
Orléans, France	47.97N	TCCON	Warneke et al. (2014)
	2.11E		
Garmisch, Germany	47.48N	TCCON+NDACC	Sussmann and Rettinger (2018a); Sussmann et al. (2012);
	11.06E		Hausmann al. (2016)

Zugspitze, Germany	47.42N	TCCON+NDACC	Sussmann and Rettinger (2018b)
	10.98E		
Jungfraujoch,	46.55N	NDACC	Zander et al. (2008)
Switzerland	7.98E		
Park Falls, WI,	45.94N	TCCON	Wennberg et al. (2017)
USA	90.27W		
Rikubetsu, Japan	43.46N	TCCON+NDACC	Morino et al. (2016); De Mazière et al. (2018)
	143.77E		
Boulder, CO, USA	39.99N	NDACC	Ortega et al. (2021)
	105.26W		
Indianapolis, IN,	39.86N	TCCON	Iraci et al. (2016)
USA	86W		
Four Corners, USA	36.8N	TCCON	Dubey et al. (2014)
	108.48W		
Lamont, OK, USA	36.5N	TCCON	Wennberg et al. (2016)
	97.49W		
Anmeyondo, South	36.54N	TCCON	Goo et al. (2014)
Korea	126.33E		
Tsukuba, Japan	36.05N	TCCON	Morino et al (2016)
	140.12E		
Nicosia, Cyprus	35.14N	TCCON	Petri et al. (2020)
	33.38E		
Edwards, CA, USA	34.96N	TCCON	Iraci et al. (2016)
	117.88W		
JPL, CA, USA	34.2N	TCCON	Wennberg et al. (2016)
	118.18W		
Pasadena Caltech,	34.14N	TCCON	Wennberg et al. (2015)
CA, USA	118.13W		
Saga, Japan	33.24N	TCCON	Kawakami et al. (2014)
	130.29E		
Heifei, China	31.9 N	TCCON	Liu et al. (2018)
	117.17E		

Izaña, Spain	28.3N	TCCON+NDACC	Blumenstock et al. (2014); García et al. (2021)
	16.48W		
Mauna Loa, HI,	19.54N	NDACC	Hannigan et al. (2009
United States	155.58W		
Altzomoni, Mexico	19.12N	NDACC	De Mazière et al. (2018);
	98.66W		
Burgos, Philippines	18.53N	TCCON	Morino et al. (2018)
	120.65E		
Manaus, Brazil	3.21S	TCCON	Dubey et al. (2014)
	60.6W		
Ascension Island,	7.92S	TCCON	Feist et al. (2014)
UK	14.33W		
Darwin, Australia	12.43S	TCCON	Griffith et al. (2014)
	130.89E		
Reunion St Denis,	20.9S	TCCON+NDACC	De Mazière et al. (2014)
France	55.49E		
Reunion Island,	21.1S	NDACC	Zhou et al. (2018)
Maido, France	55.4E		
Wollongong,	34.41S	TCCON+NDACC	Griffith et al. (2014); De Mazière et al. (2018);
Australia	150.88E		
Lauder, New	45.05S	TCCON+NDACC	Sherlock et al. (2014a, 2014b); Pollard et al. (2019)
Zealand	168.68E		Bader et al. (2017); Pollard et al. (2017)
Arrival Heights,	77.83S	NDACC	Bader et al. (2017)
Antarctica	166.67E		

Table A.3 List of AirCore sites (from NOAA AirCore data v20210813, Baier et al., 2021) used for the evaluation in Sect. 3.2.

Site, country	Latitude /Longitude [degrees]	
Boulder, CO, USA	40.03N 103.74W	
Lamont, OK, USA	36.85N 98.21W	
Lauder, New Zealand	45.50S 169.47E	
Sodankylä, Finland	67.41N 26.31E	
Park Falls, WI, USA	45.97N 90.32W	
Edwards, CA, USA	34.65N 117.29W	
Traînou, France	48.48N 1.16E	

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Code and data availability

The IFS forecasting and reanalysis system is not for public use as the ECMWF Member States are the proprietary owners. The resulting dataset is however freely available on the Copernicus Atmosphere Data Store. The CAMS GHG reanalysis can be accessed through the CAMS Atmosphere Data Store (ADS) at https://doi.org/10.24380/8fck-9w87. The format is available in both GRIB and NetCDF. The data record starts on 1 January 2003 00UTC and currently stops on 31 December 2020. Recent months will be added over time as soon as the reanalysis procedure and its validation are completed. The original data was available either as spectral coefficients with a triangular truncation of T255 or on a reduced Gaussian grid with a resolution of N128. But for the ease of the user, fields were interpolated from their native representation to a regular 0.75°x0.75° latitude longitude grid. For sub-daily data for the CAMS reanalysis is archived with a 3-hourly time step (0, 3, 6, 9, 12, 15, 18, 21 UTC). Pre-computed monthly averages are also directly available for all fields. The 3D fields are available on two different vertical resolution: 25 pressure levels (1000, 950, 925, 900, 850, 800, 700, 600, 500, 400, 300, 250, 200, 150, 100, 70, 50, 30, 10, 20, 7.5. 3. 2, hPa) 60 σ-hybrid model levels which described and are at https://www.ecmwf.int/en/forecasts/documentation-and-support/60-model-levels. The data records have 18 2D radiation fields, 2 vertically integrated atmospheric content of CO₂ and CH₄ (column-mean mole fractions, 14 2D surface fluxes variables, 32 2D meteorological fields and 16 3D fields including meteorological and greenhouse gases fields. A complete listing of the variables included in the **CAMS GHG** provided the reanalysis ADS (https://ads.atmosphere.copernicus.eu/cdsapp#!/home).

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(https://climate.esa.int/en/projects/ghgs/Data) and the Copernicus Climate Change Service (C3S) Climate Data Store (https://cds.climate.copernicus.eu). The data used to evaluate the CAMS GHG reanalysis were obtained from: the Total Carbon Column Observing Network (TCCON) Data Archive hosted by CaltechDATA at https://tccondata.org.; CH4 from the Network for the Detection of Atmospheric Composition Change (NDACC, www.ndacc.org; Sussmann et al., 2011; 2013; De Mazière et al., 2018) (see Table A2); the Observation Package (ObsPack) GLOBALVIEWplus Data Products https://gml.noaa.gov/ccgg/obspack; the NOAA AirCore dataset https://gml.noaa.gov/ccgg/obspack; and the NOAA Greenhouse Gas Marine Boundary Layer Reference https://gml.noaa.gov/ccgg/mbl/mbl.html.

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Author contributions

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Anna Agustí-Panareda, Jérôme Barré and Sébastien Massart prepared the manuscript with the rest of the co-authors, monitored the simulation, testing the impact of the surface fluxes and assimilated observations, as well as fixing bugs. Anna Agustí-Panareda was responsible of the forecasting model implementation and performed the evaluation of the vertical profiles and trends. Antje Inness led the CAMS reanalysis work and was responsible of running and monitoring the reanalysis simulations. Sébastien Massart developed the data assimilation of satellite products. Mark Parrington provided support on the biomass burning emissions and postprocessed the fire emission data in Fig. 3. Mel Ades worked on documenting the data assimilation aspects of the GHG reanalysis. Johannes Flemming, Zak Kipling and Luca Cantarello provided technical support for the diagnostics, bug fixes implemented during the testing stage and testing of the CAMS NRT satellite observations. Luke Jones provided technical support with the mass interpolation scheme and diagnostic tools. Miha Ratzinger was responsible for post-processing the dataset into the Copernicus Atmosphere Data Store. Roberto Ribas and Martin Suttie performed the satellite data acquisition and pre-processing. Richard Engelen and Vincent-Henri Peuch coordinated the efforts on the CAMS reanalyses. Bavo Langerock, Henk Eskes, Michel Ramonet and Jérôme Tarniewicz were responsible of validating the reanalysis dataset with independent observations including the production of surface and column CO₂ and CH₄ evaluation plots. Ilse Aben, Tobias Borsdorff, Michael Buchwitz, Cyril Crevoisier, Otto Hasekamp, Nicolas Meilhac, Stefan Noel,

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References

710

- Agustí-Panareda, A., Diamantakis, M., Bayona, V., Klappenbach, F. & Butz, A.: Improving the inter-hemispheric gradient of total column atmospheric CO₂ and CH₄ in simulations with the ECMWF semi-Lagrangian atmospheric global model. *Geoscientific Model Development* **10**, https://doi.org/10.5194/gmd-10-1-2017, 1–18, 2017.
- Agustí-Panareda, A., Diamantakis, M., Massart, S., Chevallier, F., Muñoz-Sabater, J., Barré, J., Curcoll, R., Engelen, R., Langerock, B., Law, R. M., Loh, Z., Morguí, J. A., Parrington, M., Peuch, V.-H., Ramonet, M., Roehl, C., Vermeulen, A. T., Warneke, T., and Wunch, D.: Modelling CO2 weather why horizontal resolution matters, Atmos. Chem. Phys., 19, 7347–7376, https://doi.org/10.5194/acp-19-7347-2019, 2019.
- Agustí-Panareda, A., McNorton, J., Balsamo, G., Baier, B.C., Bousserez, N., Boussetta, S., Brunner, D., Chevallier, F., Choulga, M., Diamantakis, M., Engelen, R., Flemming, J., Granier, C. Guevara, M., Denier van der Gon, H., Elguindi, N., Haussaire, J.-M., Jung, M., Janssens-Maenhout, G., Kivi, R., Massart, S., Papale, D., Parrington, M., Razinger, M., Sweeney, C., Vermeulen, A., Walther, S. Global nature run data with realistic high-resolution carbon weather for the year of the Paris Agreement. Sci Data 9, 160 (2022). https://doi.org/10.1038/s41597-022-01228-2, 2022.

725

- Alos, A. L. *et al.* The Copernicus Climate Data Store: ECMWF's approach to providing online access to climate data and tools. in *AGU Fall Meeting Abstracts* vol. 2019 PA31A-05, 2019.
- Andrews, A. E., Kofler, J. D., Trudeau, M. E., Williams, J. C., Neff, D. H., Masarie, K. A., Chao, D. Y., Kitzis, D. R., Novelli,
 P. C., Zhao, C. L., Dlugokencky, E. J., Lang, P. M., Crotwell, M. J., Fischer, M. L., Parker, M. J., Lee, J. T., Baumann, D. D.,
 Desai, A. R., Stanier, C. O., De Wekker, S. F. J., Wolfe, D. E., Munger, J. W., and Tans, P. P.: CO2, CO, and CH4 measurements from tall towers in the NOAA Earth System Research Laboratory's Global Greenhouse Gas Reference Network: instrumentation, uncertainty analysis, and recommendations for future high-accuracy greenhouse gas monitoring efforts, Atmos. Meas. Tech., 7, 647–687, https://doi.org/10.5194/amt-7-647-2014, 2014.

735

Bader, W., Bovy, B., Conway, S., Strong, K., Smale, D., Turner, A. J., Blumenstock, T., Boone, C., Collaud Coen, M., Coulon, A., Garcia, O., Griffith, D. W. T., Hase, F., Hausmann, P., Jones, N., Krummel, P., Murata, I., Morino, I., Nakajima, H.,

O'Doherty, S., Paton-Walsh, C., Robinson, J., Sandrin, R., Schneider, M., Servais, C., Sussmann, R. and Mahieu, E.: The recent increase of atmospheric methane from 10 years of ground-based NDACC FTIR observations since 2005, Atmos. Chem. 740 Phys., 17(3), 2255–2277, doi:10.5194/acp-17-2255-2017, 2017.

Baier, B., Sweeney, C., Newberger, T., Higgs, J., Wolter, S., & NOAA Global Monitoring Laboratory: NOAA AirCore atmospheric sampling system profiles (Version 20210813) [Data set]. NOAA GML, https://doi.org/10.15138/6AV0-MY81, 2021.

745

Batchelor, R. L., Strong, K., Lindenmaier, R., Mittermeier, R. L., Fast, H., Drummond, J. R. and Fogal, P. F.: A New Bruker IFS 125HR FTIR Spectrometer for the Polar Environment Atmospheric Research Laboratory at Eureka, Nunavut, Canada: Measurements and Comparison with the Existing Bomem DA8 Spectrometer, J. Atmos. Ocean. Technol., 26(7), 1328–1340, doi:10.1175/2009JTECHA1215.1, 2009.

750

Bechtold, P. et al. Representing Equilibrium and Nonequilibrium Convection in Large-Scale Models. Journal of the Atmospheric Sciences **71**, 734–753, 2014.

755

Bergamaschi, P., M. Krol, F. Dentener, A. Vermeulen, F. Meinhardt, R. Graul, M. Ramonet, W. Peters, and E. J. Dlugokencky, Inverse modelling of national and European CH4 emissions using the atmospheric zoom model TM5, Atmos, Chem. Phys., 5, 2431–2460, 2005.

Bergamaschi, P. et al.: Inverse modeling of global and regional CH4 emissions using SCIAMACHY satellite retrievals. Journal of Geophysical Research: Atmospheres 114, 2009.

760

Blumenstock, T., Hase, F., Schneider, M., Garcia, O. & Sepulveda, E. TCCON data from Izana (ES), Release GGG2014.R0 (Version GGG2014.R0) [Data set], https://doi.org/10.14291/TCCON.GGG2014.IZANA01.R0/1149295, 2014.

765

Boussetta, S. et al.: Natural land carbon dioxide exchanges in the ECMWF integrated forecasting system: Implementation and offline validation. Journal of Geophysical Research: Atmospheres 118, 5923–5946, 2013.

Brühl, C., and P. J. Crutzen, The MPIC 2D model, NASA Ref. Publ., 1292, 103–104, 1993.

Brunke, E-G., Labuschagne, C., Parker, B., Scheel, H.E. and Whittlestone, S. Baseline air mass selection at

Cape Point, South Africa: Application of 222Rn and other filter criteria to CO2. Atmospheric Environment 38, 33, 5693 – 5702, 2004.

- Buchwitz M, Reuter M, Schneising O, Boesch H, Guerlet S, Dils B, Aben I, Armante R, Bergamaschi P, Blumenstock T, Bovensmann H, Brunner D, Buchmann B, Burrows J, Butz A, Chédin A, Chevallier F, Crevoisier C, Deutscher N, Frankenberg C, Hase F, Hasekamp O, Heymann J, Kaminski T, Laeng A, Lichtenberg G, Demazière M, Noël S, Notholt J, Orphal J, Popp C, Parker R, Scholze M, Sussmann R, Stiller G, Warneke T, Zehner C, Bril A, Crisp D, Griffith D, Kuze A, O'dell C, Oshchepkov S, Sherlock V, Suto H, Wennberg P, Wunch D, Yokota T, Yoshida Y. The Greenhouse Gas Climate Change Initiative (GHG-CCI): Comparison and quality assessment of near-surface-sensitive satellite-derived CO2 and CH4 global data sets. REMOTE SENSING OF ENVIRONMENT 162; 2015. p. 344-362. JRC91437, 2015.
- Butz, A., Hasekamp, O. P., Frankenberg, C., Vidot, J., and Aben, I., CH4 retrievals from space-based solar backscatter measurements: Performance evaluation against simulated aerosol and cirrus loaded scenes, J. Geophys. Res., Volume 115, Issue D24, 27, DOI: 10.1029/2010JD014514, 2010.

- Butz, A., Guerlet, S., Hasekamp, O., et al., Toward accurate CO2 and CH4 observations from GOSAT, Geophys. Res. Lett., doi:10.1029/2011GL047888, 2011.
- Chevallier, F., Remaud, M., O'Dell, C. W., Baker, D., Peylin, P., and Cozic, A.: Objective evaluation of surface- and satellite-driven carbon dioxide atmospheric inversions, Atmos. Chem. Phys., 19, 14233–14251, https://doi.org/10.5194/acp-19-14233-790 2019.
 - Chevallier, F., et al.: CO₂ surface fluxes at grid point scale estimated from a global 21-year reanalysis of atmospheric measurements, J. Geo-phys. Res., doi:10.1029/2010JD013887, 2010.
- 795 Chevallier, F. Description of the CO2 inversion production chain 2020, Copernicus Atmosphere Monitoring Service. https://atmosphere.copernicus.eu/sites/default/files/2020-06/CAMS73 2018SC2 %20D5.2.12020 202004 %20CO2%20inversion%20production%20chain v1.pdf, 2000.
- Colomb, A., Delmotte, M., Pichon, J., Ramonet, M., Yver-Kwok, C., ICOS RI, 2020. ICOS ATC CO2 Release, Puy de Dôm e (10.0 m), 2016-08-25–2020-05-31, https://hdl.handle.net/11676/xMV5-Hxu7bA9q872kedNJIaF, 2020.
 - Conway, T. J. and K. W. Thoning. Short-term variations of atmospheric carbon dioxide at the South Pole, Antar ctic Journal of the United States, 236-238, 1990.

805 Conway, T.J., P.P. Tans, L.S. Waterman, K.W. Thoning, D.R. Kitzis, K.A. Masarie, and N. Zhang: Evidence for interannual variability of the carbon cycle from the NOAA/CMDL global air sampling network, J. Geophys. Res., 99, 22831-22855, 1994.

Crevoisier, C. *et al.*: First year of upper tropospheric integrated content of CO₂ from IASI hyperspectral infrared observations. *Atmospheric Chemistry and Physics* **9**, 4797–4810, 2009a.

810

Crevoisier, C. *et al.*: Tropospheric methane in the tropics – first year from IASI hyperspectral infrared observations. *Atmospheric Chemistry and Physics* **9**, 6337–6350, 2009b.

Crevoisier, C., Clerbaux, C., Guidard, V., Phulpin, T., Armante, R., Barret, B., Camy-Peyret, C., Chaboureau, J.-P., Coheur, P.-F., Crépeau, L., Dufour, G., Labonnote, L., Lavanant, L., Hadji-Lazaro, J., Herbin, H., Jacquinet-Husson, N., Payan, S., Péquignot, E., Pierangelo, C., Sellitto, P., and Stubenrauch, C.: Towards IASI-New Generation (IASI-NG): impact of improved spectral resolution and radiometric noise on the retrieval of thermodynamic, chemistry and climate variables, Atmos. Meas. Tech., 7, 4367–4385, https://doi.org/10.5194/amt-7-4367-2014, 2014.

820 Crisp and Co-authors: A constellation architecture for monitoring carbon dioxide and methane from space. CEOS Atmospheric Composition Virtual Constellation Greenhouse Gas Team Rep., 173 pp., http://ceos.org/document_management/Virtual_Constellations/ACC/Documents/CEOS_AC-VC_GHG_White_Paper_Version_1_20181009.pdf., 2018.

De Mazière, M. et al. TCCON data from Réunion Island (RE), Release GGG2014.R0 (Version GGG2014.R0) [Data set]. http://dx.doi.org/10.14291/tccon.ggg2014.reunion01.R0/1149288, 2014.

De Mazière, M. *et al.*: The Network for the Detection of Atmospheric Composition Change (NDACC): history, status and perspectives. *Atmospheric Chemistry and Physics* **18**, 4935–4964, 2018.

Deutscher, N. M. et al. TCCON data from Bialystok (PL), Release GGG2014.R1 (Version GGG2014.R1) [Data set]. https://doi.org/10.14291/TCCON.GGG2014.BIALYSTOK01.R1/1183984, 2015.

Diamantakis, M. & Magnusson, L.: Sensitivity of the ECMWF Model to Semi-Lagrangian Departure Point Iterations. *Monthly Weather Review* **144**, 3233–3250, 2016.

835

Doumbia, T., Granier, C., Elguindi, N., Bouarar, I., Darras, S., Brasseur, G. et al. Changes in global air pollutant emissions during the COVID-19 pandemic: a dataset for atmospheric modeling. Earth System Science Data, 13, 4191-4206. doi:10.5194/essd-13-4191-2021, 2021.

- Dlugokencky, E.J., L.P. Steele, P.M. Lang, and K.A. Masarie, The growth rate and distribution of atmospheric methane, J. Geophys. Res., 99, 17,021-17,043, 1994.
- Dlugokencky, E.J., J.W. Mund, A.M. Crotwell, M.J. Crotwell, and K.W. Thoning. Atmospheric Carbon Dioxide Dry Air Mole Fractions from the NOAA GML Carbon Cycle Cooperative Global Air Sampling Network, 1968-2020, Version: 2021-07-30, https://doi.org/10.15138/wkgj-f215, 2021.
 - Dubey, M. K. et al. TCCON data from Manaus (BR), Release GGG2014.R0 (Version GGG2014.R0) [Data set]. https://doi.org/10.14291/TCCON.GGG2014.MANAUS01.R0/1149274,1 2014.
- Dubey, M., R. Lindenmaier, B. Henderson, D. Green, N. Allen, C. Roehl, J.-F. Blavier, Z. Butterfield, S. Love, J. Hamelmann, D. Wunch. TCCON data from Four Corners, NM, USA, Release GGG2014R0. TCCON data archive, hosted by CaltechDATA, California Institute of Technology, Pasadena, CA, U.S.A. https://doi.org/10.14291/tccon.ggg2014.fourcorners01.R0/1149272, 2014.
- 855 Feist, D. G., Arnold, John, & Geibel. TCCON data from Ascension Island (SH), Release GGG2014.R0 (Version GGG2014.R0) [Data set]. https://doi.org/10.14291/TCCON.GGG2014.ASCENSION01.R0/1149285, 2014.
- Fisher, M.: Generalized frames on the sphere with application to background error covariance modelling, Seminar on recent developments in numerical methods for atmospheric and ocean modelling, 6–10 September 2004, Proceedings, ECMWF, pp. 87–101, available from ECMWF, Shinfield Park, Reading, Berk- shire, RG2 9AX, UK, 2004.
 - Fisher, M.: Wavelet Jb A new way to model the statistics of back- ground errors, ECMWF Newsletter, 106, 23–28, available from ECMWF, Shinfield Park, Reading, Berkshire, RG2 9AX, UK, 2006.
- Flemming, J., Huijnen, V., Arteta, J., Bechtold, P., Beljaars, A., Blechschmidt, A.-M., Diamantakis, M., Engelen, R. J., Gaudel, A., Inness, A., Jones, L., Josse, B., Katragkou, E., Marecal, V., Peuch, V.-H., Richter, A., Schultz, M. G., Stein, O., and Tsikerdekis, A.: Tropospheric chemistry in the Integrated Forecasting System of ECMWF, Geosci. Model Dev., 8, 975–1003, https://doi.org/10.5194/gmd-8-975-2015, 2015.
- Frankenberg, C., Aben, I., Bergamaschi, P., Dlugokencky, E. J., van Hees, R., Houweling, S., van der Meer, P., Snel, R., and Tol, P.: Global column-averaged methane mixing ratios from 2003 to 2009 as derived from SCIAMACHY: trends and variability, J. Geophys. Res.-Atmos., 116, D04302, doi:10.1029/2010JD014849, 2011.

Friedlingstein, P., Jones, M. W., O'Sullivan, M., Andrew, R. M., Bakker, D. C. E., Hauck, J., Le Quéré, C., Peters, G. P.,
Peters, W., Pongratz, J., Sitch, S., Canadell, J. G., Ciais, P., Jackson, R. B., Alin, S. R., Anthoni, P., Bates, N. R., Becker, M.,
Bellouin, N., Bopp, L., Chau, T. T. T., Chevallier, F., Chini, L. P., Cronin, M., Currie, K. I., Decharme, B., Djeutchouang, L.,
Dou, X., Evans, W., Feely, R. A., Feng, L., Gasser, T., Gilfillan, D., Gkritzalis, T., Grassi, G., Gregor, L., Gruber, N., Gürses,
Ö., Harris, I., Houghton, R. A., Hurtt, G. C., Iida, Y., Ilyina, T., Luijkx, I. T., Jain, A. K., Jones, S. D., Kato, E., Kennedy, D.,
Klein Goldewijk, K., Knauer, J., Korsbakken, J. I., Körtzinger, A., Landschützer, P., Lauvset, S. K., Lefèvre, N., Lienert, S.,
Liu, J., Marland, G., McGuire, P. C., Melton, J. R., Munro, D. R., Nabel, J. E. M. S., Nakaoka, S.-I., Niwa, Y., Ono, T., Pierrot,
D., Poulter, B., Rehder, G., Resplandy, L., Robertson, E., Rödenbeck, C., Rosan, T. M., Schwinger, J., Schwingshackl, C.,
Séférian, R., Sutton, A. J., Sweeney, C., Tanhua, T., Tans, P. P., Tian, H., Tilbrook, B., Tubiello, F., van der Werf, G., Vuichard,
N., Wada, C., Wanninkhof, R., Watson, A., Willis, D., Wiltshire, A. J., Yuan, W., Yue, C., Yue, X., Zaehle, S., and Zeng, J.:
Global Carbon Budget 2021, Earth Syst, Sci. Data Discuss. [preprint], https://doi.org/10.5194/essd-2021-386, in review, 2021.

García, O. E., Schneider, M., Sepúlveda, E., Hase, F., Blumenstock, T., Cuevas, E., Ramos, R., Gross, J., Barthlott, S., Röhling, A. N., Sanromá, E., González, Y., Gómez-Peláez, Á. J., Navarro-Comas, M., Puentedura, O., Yela, M., Redondas, A., Carreño, V., León-Luis, S. F., Reyes, E., García, R. D., Rivas, P. P., Romero-Campos, P. M., Torres, C., Prats, N., Hernández, M. and López, C.: Twenty years of ground-based NDACC FTIR spectrometry at Izaña Observatory-overview and long-term comparison to other techniques, Atmos. Chem. Phys., 21(20), 15519–15554, doi:10.5194/acp-21-15519-2021, 2021.

885

900

905

Goo, T.-Y., Oh, Y.-S. & Velazco, V. A. TCCON data from Anmeyondo (KR), Release GGG2014.R0 (Version GGG2014.R0) [Data set]. https://doi.org/10.14291/TCCON.GGG2014.ANMEYONDO01.R0/1149284, 2014.

895 Griffith, D. W. T. et al. TCCON data from Darwin (AU), Release GGG2014.R0 (Version GGG2014.R0) [Data set]. https://doi.org/10.14291/TCCON.GGG2014.DARWIN01.R0/1149290, 2014.

Griffith, D. W. T. et al. TCCON data from Wollongong (AU), Release GGG2014.R0 (Version GGG2014.R0) [Data set]. http://dx.doi.org/10.14291/tccon.ggg2014.wollongong01.R0/1149291, 2014.

Guerlet, S., Butz, A., Schepers, D., Basu, S., Hasekamp, O. P., Kuze, A., Yokota, T., Blavier, J.-F., Deutscher, N. M., Griffith, D. W., Hase, F., Kyro, E., Morino, I., Sherlock, V., Sussmann, R., Galli, A., and Aben, I.: Impact of aerosol and thin cirrus on retrieving and validating XCO2 from GOSAT shortwave infrared measurements, Journal of Geophysical Research: Atmospheres, 118, 4887–4905, https://doi.org/10.1002/jgrd.50332, 2013.

- Hannigan, J. W., M. T. Coffey, and A. Goldman. Semiautonomous FTS Observation System for Remote Sensing of Stratospheric and Tropospheric Gases. Journal of Atmospheric and Oceanic Technology 26, pp. 1814–1828. doi: 10.1175/2009JTECHA1230.1, 2009.
- 910 Hase, F., Blumenstock, T., Dohe, S., Gros, J. & Kiel, M. TCCON data from Karlsruhe (DE), Release GGG2014.R1 (Version GGG2014.R1) [Data set]. https://doi.org/10.14291/TCCON.GGG2014.KARLSRUHE01.R1/1182416, 2015.
- Haszpra, L., Barcza, Z., Bakwin, P. S., Berger, B. W., Davis, K. J., Weidinger, T. Measuring system for the l
 ong-term monitoring of biosphere/atmosphere exchange of carbon dioxide. J. of Geophysical Research 106D, 3057-3070,
 2001.
 - Hausmann, P., Sussmann, R., and Smale, D.: Contribution of oil and natural gas production to renewed increase in atmospheric methane (2007–2014): top–down estimate from ethane and methane column observations, Atmos. Chem. Phys., 16, 3227-3244, doi:10.5194/acp-16-3227-2016, 2016.
 - Hersbach, H. et al.: The ERA5 global reanalysis. Quarterly Journal of the Royal Meteorological Society 146, 1999–2049, 2020.

- Heymann, J., M. Reuter, M. Hilker, M. Buchwitz, O. Schneising, H. Bovensmann, J. P. Burrows, A. Kuze, H. Suto, N. M.
 Deutscher, M. K. Dubey, D. W. T. Griffith, F. Hase, S. Kawakami, R. Kivi, I. Morino, C. Petri, C. Roehl, M. Schneider, V. Sherlock, R. Sussmann, V. A. Velazco, T. Warneke, and D. Wunch, Consistent satellite XCO2 retrievals from SCIAMACHY and GOSAT using the BESD algorithm, Atmos. Meas. Tech., 8, 2961-2980, 2015.
- Houweling, S., F. Dentener, and J. Lelieveld, The impact of nonmethane hydrocarbon compounds on tropospheric photochemistry, J. Geophys. Res., 103(D9), 10,673–10,696, doi:10.1029/97JD03582, 1998.
 - Houweling, S., Kaminski, T., Dentener, F., Lelieveld, J. & Heimann, M. Inverse modeling of methane sources and sinks using the adjoint of a global transport model. *Journal of Geophysical Research: Atmospheres* **104**, 26137–26160, 1999.
- 935 ICOS RI: ICOS Atmospheric Greenhouse Gas Mole Fractions of CO2 for the period Sep 2015- May 2020, final quality c ontrolled Level 2 data for 23 stations (62 vertical levels), pre-release 2020-1, doi:10.18160/GZ5S-4GPR, 2020.
 - Inness, A., Ades, M., Agustí-Panareda, A., Barré, J., Benedictow, A., Blechschmidt, A.-M., Dominguez, J. J., Engelen, R., Eskes, H., Flemming, J., Huijnen, V., Jones, L., Kipling, Z., Massart, S., Parrington, M., Peuch, V.-H., Razinger, M., Remy,

- 940 S., Schulz, M., and Suttie, M.: The CAMS reanalysis of atmospheric composition, Atmos. Chem. Phys., 19, 3515–3556, https://doi.org/10.5194/acp-19-3515-2019, 2019.
- IPCC, Climate Change 2014: Synthesis Report. Contribution of Working Groups I, II and III to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change [Core Writing Team, R.K. Pachauri and L.A. Meyer (eds.)]. IPCC, Geneva, Switzerland, 151 pp., 2014.
- IPCC, Policymakers. 2021: Summary for In: Climate Change The Physical Science Basis. Contribution of Working Group I to the Sixth Assessment Report of the Intergovernmental Panel on Climate Change [Masson-Delmotte, V., P. Zhai, A. Pirani, S. L. Connors, C. Péan, S. Berger, N. Caud, Y. Chen, L. 950 Goldfarb, M. I. Gomis, M. Huang, K. Leitzell, E. Lonnoy, J.B.R. Matthews, T. K. Maycock, T. Waterfield, O. Yelekci, R. Yu and B. Zhou (eds.)]. Cambridge University Press. In Press., 2021.
- Iraci, L., J. Podolske, P. Hillyard, C. Roehl, P. O. Wennberg, J.-F. Blavier, J. Landeros, N. Allen, D. Wunch, J. Zavaleta, E. Quigley, G. Osterman, E. Barrow, J. Barney. TCCON data from Indianapolis, Indiana, USA, Release GGG2014R1. TCCON data archive, hosted by CaltechDATA, California Institute of Technology, Pasadena, CA, U.S.A. https://doi.org/10.14291/tccon.ggg2014.indianapolis01.R1/1330094, 2016.
 - Iraci, L. T. et al. TCCON data from Edwards (US), Release GGG2014.R1 (Version GGG2014.R1) [Data set]. https://doi.org/10.14291/TCCON.GGG2014.EDWARDS01.R1/1255068, 2016.

- Janssens-Maenhout, Greet; Crippa, Monica; Guizzardi, Diego; Muntean, Marilena; Schaaf, Edwin. Emissions Database for Global Atmospheric Research, version v4.2 (time-series). European Commission, Joint Research Centre (JRC) [Dataset] PID: http://data.europa.eu/89h/jrc-edgar-emissiontimeseriesv42, 2011.
- Janssens-Maenhout, G., Crippa, M.; Guizzardi, D.; Muntean, M.; Schaaf, E. JRC-EDGARv431_AP_timeseries. European Commission, Joint Research Centre (JRC) [Dataset] PID: http://data.europa.eu/89h/jrc-edgar-edgar-v431_timeseries, 2016.
 - Kaiser, J. W. *et al.*: Biomass burning emissions estimated with a global fire assimilation system based on observed fire radiative power. *Biogeosciences* **9**, 527–554, 2012.
 - Karion, A., Sweeney, C., Tans, P., Newberger, T. AirCore: An Innovative Atmospheric Sampling System, J. Atmos. Ocean. Technol., 27, 1839-1843, https://doi.org/10.1175/2010JTECHA1448.1, 2010.

- Kawakami, S. et al. TCCON data from Saga (JP), Release GGG2014.R0 (Version GGG2014.R0) [Data set]. 975 https://doi.org/10.14291/TCCON.GGG2014.SAGA01.R0/1149283, 2014.
 - Kivi, R., Heikkinen, P. & Kyro, E. TCCON data from Sodankylä (FI), Release GGG2014.R0 (Version GGG2014.R0) [Data set]. https://doi.org/10.14291/TCCON.GGG2014.SODANKYLA01.R0/1149280, 2014.
- Lamarque, J.-F., Shindell, D. T., Josse, B., Young, P. J., Cionni, I., Eyring, V., Bergmann, D., Cameron-Smith, P., Collins, W. J., Doherty, R., Dalsoren, S., Faluvegi, G., Folberth, G., Ghan, S. J., Horowitz, L. W., Lee, Y. H., MacKenzie, I. A., Nagashima, T., Naik, V., Plummer, D., Righi, M., Rumbold, S. T., Schulz, M., Skeie, R. B., Stevenson, D. S., Strode, S., Sudo, K., Szopa, S., Voulgarakis, A., and Zeng, G.: The Atmospheric Chemistry and Climate Model Intercomparison Project (ACCMIP): overview and description of models, simulations and climate diagnostics, 6, 179–206, https://doi.org/10.5194/gmd-6-179-2013, 2013, 2013.
 - Lambert, G. & Schmidt, S.: Re-evaluation of the oceanic flux of methane: Uncertainties and long-term variations. *Chemosphere* **26**, 579–589, 1993.
- 990 Le Quéré, C., Jackson, R.B., Jones, M.W. et al.: Temporary reduction in daily global CO2 emissions during the COVID-19 forced confinement. Nat. Clim. Chang. 10, 647–653, https://doi.org/10.1038/s41558-020-0797-x, 2020.
- Liu C., Wang, W., Sun, Y. TCCON data from Hefei, China, Release GGG2014R0. TCCON data archive, hosted by CaltechDATA, California Institute of Technology, Pasadena, CA, U.S.A. http://dx.doi.org/10.14291/tccon.ggg2014.hefei01.R0, 2018.
 - Lopez, M., M. Schmidt, M. Ramonet, J. L. Bonne, A. Colomb, V. Kazan, P. Laj and J. M. Pichon. Three years of semicontinuous greenhouse gas measurements at the Puy de Dôme station (central France). Atmospheric Measurement Techniques(8): 3941-3958, 2015.
 - Makarova, M. V., Kirner, O., Timofeev, Y. M., Poberovskii, A. V., Imkhasin, K. K., Osipov, S. I. and Makarov, B. K.: Analysis of methane total column variations in the atmosphere near St. Petersburg using ground-based measurements and simulations, Izv. Atmos. Ocean. Phys., 51(2), 177–185, doi:10.1134/S0001433815010089, 2015.
- 1005 Masarie, K. A., Peters, W., Jacobson, A. R. & Tans, P. P.: ObsPack: a framework for the preparation, delivery, and attribution of atmospheric greenhouse gas measurements. *Earth System Science Data* **6**, 375–384, 2014.

- Masarie, K.A. and Tans, P.P.: Extension and Integration of Atmospheric Carbon Dioxide Data into a Globally Consistent Measurement Record, J. Geophys. Res., .100, 11593-11610, 1995.
- Massart, S., and Bonavita, M.: Ensemble of Data Assimilations applied to the CAMS' greenhouse gases analysis. ECMWF Technical Memorandom 780, available at: https://www.ecmwf.int/en/elibrary/16438-ensemble-data-assimilations-applied-cams-greenhouse-gases-analysis (last access: xxx), 2016.

- 1015 Massart, S. *et al.*: Assimilation of atmospheric methane products into the MACC-II system: from SCIAMACHY to TANSO and IASI. *Atmospheric Chemistry and Physics* **14**, 6139–6158, 2014.
 - Massart, S. *et al.*: Ability of the 4-D-Var analysis of the GOSAT BESD XCO₂ retrievals to characterize atmospheric CO₂ at large and synoptic scales. *Atmospheric Chemistry and Physics* **16**, 1653–167, 2016.
 - Massart, S., Agusti-Panareda, A. and Flemming, J.: Evidence of a stratospheric methane bias in the IFS against MIPAS data, ECMWF Technical Memorandum, 814, 21 pages, https://www.ecmwf.int/sites/default/files/elibrary/2017/17698-evidence-stratospheric-methane-bias-ifs-against-mipas-data.pdf, 2017.
- Matthews, E., Fung, I. & Lerner, J.: Methane emission from rice cultivation: Geographic and seasonal distribution of cultivated areas and emissions. *Global Biogeochemical Cycles* **5**, 3–24, 1991.
- McNorton, J., Bousserez, N., Agustí-Panareda, A., Balsamo, G., Engelen, R., Huijnen, V., Inness, A., Kipling, Z., Parrington, M., and Ribas, R.: Quantification of methane emissions from hotspots and during COVID-19 using a global atmospheric inversion, Atmos. Chem. Phys. Discuss. [preprint], https://doi.org/10.5194/acp-2021-1056, in review, 2022.
 - Morino, I., Yokozeki, N., Matsuzaki, T., & Horikawa. TCCON data from Rikubetsu (JP), Release GGG2014.R1 (Version GGG2014.R1) [Data set]. https://doi.org/10.14291/TCCON.GGG2014.RIKUBETSU01.R1/1242265, 2016.
- Morino, I., Matsuzaki, T. & Horikawa, M. TCCON data from Tsukuba (JP), 125HR, Release GGG2014.R1 (Version GGG2014.R1) [Data set]. https://doi.org/10.14291/TCCON.GGG2014.TSUKUBA02.R1/1241486, 2016.
- Morino, I., V. A. Velazco, A. Hori, O. Uchino, D. W. T. Griffith. TCCON data from Burgos, Philippines, Release GGG2014R0. TCCON data archive, hosted by CaltechDATA, California Institute of Technology, Pasadena, CA, U.S.A. http://dx.doi.org/10.14291/tccon.ggg2014.burgos01.R0/1368175, 2018.

- NOAA Carbon Cycle Group ObsPack Team. (2019). Multi-laboratory compilation of atmospheric methane data for the period 1957-2017; obspack_ch4_1_GLOBALVIEWplus_v1.0_2019_01_08; [Data set]. NOAA Earth System Research Laboratory, Global Monitoring Division. https://doi.org/10.25925/20190108
- Notholt, J. et al. TCCON data from Ny Ålesund, Spitsbergen (NO), Release GGG2014.R1(Version R1) [Data set]. https://doi.org/10.14291/TCCON.GGG2014.NYALESUND01.R1, 2019.

- Notholt, J. et al. TCCON data from Bremen (DE), Release GGG2014.R0 (Version GGG2014.R0) [Data set]. https://doi.org/10.14291/TCCON.GGG2014.BREMEN01.R0/1149275, 2014.
 - Olivier, J. and Janssens-Maenhout G.: CO₂ Emissions from Fuel Combustion 2012 Edition, IEA CO₂ report 2012, Part III, Greenhouse-Gas Emissions, OECD Publishing, Paris, ISBN 978-92-64-17475-7, 2012.
- Ortega, I., J.W. Hannigan, R.R. Buchholz, G. Pfister. Long-term variability and source signature of gases emitted from oil & natural gas and cattle feedlot operations in the Colorado front range. Atmospheric Environment, Volume 263, 118663, ISSN 1352-2310, https://doi.org/10.1016/j.atmosenv.2021.118663, 2021.
- Peterson, J.T., W.D. Komhyr, L.S. Waterman, R.H. Gammon, K.W. Thoning, and T.J. Conway, Atmospheric CO2 variations at Barrow, Alaska, 1973 1982, J. Atmos. Chem., 4, 491 510, 1986.
 - Patra, P. K., et al.: TransCom model simulations of hourly atmo-spheric CO2: Analysis of synoptic-scale variations for the period 2002–2003, Global Biogeochem. Cycles, 22, GB4013, doi:10.1029/2007GB003081, 2008
- Patra, P. K., Houweling, S., Krol, M., Bousquet, P., Belikov, D., Bergmann, D., Bian, H., Cameron-Smith, P., Chipperfield, M. P., Corbin, K., Fortems-Cheiney, A., Fraser, A., Gloor, E., Hess, P., Ito, A., Kawa, S. R., Law, R. M., Loh, Z., Maksyutov, S., Meng, L., Palmer, P. I., Prinn, R. G., Rigby, M., Saito, R., and Wilson, C.: TransCom model simulations of CH4 and related species: linking transport, surface flux and chemical loss with CH4 variability in the troposphere and lower stratosphere, Atmos. Chem. Phys., 11, 12813–12837, https://doi.org/10.5194/acp-11-12813-2011, 2011.
 - Petri, C., C. Rousogenous, T. Warneke, M. Vrekoussis, S. Sciare, J. Notholt. TCCON data from Nicosia, Cyprus, Release GGG2014R0. TCCON data archive, hosted by CaltechDATA, California Institute of Technology, Pasadena, CA, U.S.A. http://doi.org/10.14291/tccon.ggg2014.nicosia01.R0, 2020.
- 1075 Pollard, D. F., Sherlock, V., Robinson, J., Deutscher, N. M., Connor, B. and Shiona, H.: The Total Carbon Column Observing Network site description for Lauder, New Zealand, Earth Syst. Sci. Data, 9(2), 977–992, doi:10.5194/essd-9-977-2017, 2017.

- Ramonet, M. and P. Monfray. CO2 baseline concept in 3-D atmospheric transport models. Tellus B 48(4): 502-520, 1996.
- 1080 Ramonet, M., P. Ciais, T. Aalto, C. Aulagnier, F. Chevallier, D. Cipriano, T.J. Conway, L. Haszpra, V. Kazan, F. Mei nhardt, J.-D. Paris, M. Schmidt, P. Simmonds, I. Xueref-Remy, and J. Necki, A recent build-up of atmospheric CO2 over Europe. Part 1: observed signals and possible explanations, Tellus B, DOI: 10.1111/j.1600-0889.2009.00442, 2010.
- Ramonet, M., Langerock, B., Warneke, T., Eskes, H.J.: Validation report of the CAMS greenhouse gas global re-analysis, years 2003-2020, Copernicus Atmosphere Monitoring Service (CAMS) report CAMS84_2018SC3_D5.1.2-2020_v0.1.pdf, April 2021, doi: 10.24380/438c-4597, 2021.
- Reuter, M., Bovensmann, H., Buchwitz, M., Burrows, J. P., Connor, B. J., Deutscher, N. M., Griffith, D. W. T., Heymann, J., Keppel-Aleks, G., Messerschmidt, J., Notholt, J., Petri, C., Robinson, J., Schneising, O., Sherlock, V., Velazco, V., Warneke, W., Wennberg, P. O., and Wunch, D.: Retrieval of atmospheric CO2 with enhanced accuracy and precision from SCIAMACHY: Validation with FTS measurements and comparison with model results, J. Geophys. Res., 116, D04301, https://doi.org/10.1029/2010JD015047, 2011.
- Ridgwell, A. J., Marshall, S. J. & Gregson, K.: Consumption of atmospheric methane by soils: A process-based model. *Global Biogeochemical Cycles* **13**, 59–70, 1999.
 - Rodgers, C. D.: Inverse Methods for Atmospheric Sounding: Theory and Practice. World Scientific, 2000.
- Sanderson, M. G.: Biomass of termites and their emissions of methane and carbon dioxide: A global database. *Global Biogeochemical Cycles* **10**, 543–557, 1996.
 - Sarra, A. G. D., Karion, A., Arlyn Andrews, Colomb, A., Scheeren, B., Viner, B., Myhre, C. L., Miller, C. E., Plass-Duelmer, C., Plass-Duelmer, C., Sloop, C. D., Sweeney, C., Kubistin, D., Jaffe, D., Dlugokencky, E., Vitkova, G., Manca, G., Huilin Chen, Lehner, I., ... Kazan, V. (2021). Multi-laboratory compilation of atmospheric carbon dioxide data for the years 20202021; obspack_co2_1_NRT_v6.1_2021-02-02 [Data set]. NOAA Earth System Research Laboratory, Global Monitoring
- 1105 2021; obspack_co2_1_NRT_v6.1_2021-02-02 [Data set]. NOAA Earth System Research Laboratory, Global Monitoring Laboratory. https://doi.org/10.25925/20210108, 2021.
 - Sandu, I., Beljaars, A., Bechtold, P., Mauritsen, T. & Balsamo, G.: Why is it so difficult to represent stably stratified conditions in numerical weather prediction (NWP) models? *Journal of Advances in Modeling Earth Systems* 5, 117–133, 2013.

- Schepers, D. *et al.*: Methane retrievals from Greenhouse Gases Observing Satellite (GOSAT) shortwave infrared measurements: Performance comparison of proxy and physics retrieval algorithms. *Journal of Geophysical Research: Atmospheres* 117, 2012.
- Schmidt, M., Graul, R., Sartorius, H., and Levin, L.: The Schauinsland CO2 record: 30 years of continental observations and their implications for the variability of the European CO2 budget, J. Geophys. Res.-Atmos., 108, 4619-4626, 2003.
- Schuldt, K. N., Mund, J., Luijkx, I. T., Jacobson, A. R., Cox, A., Vermeulen, A., Manning, A., Beyersdorf, A., Manning, A., Karion, A., Hensen, A., Arlyn Andrews, Frumau, A., Colomb, A., Scheeren, B., Law, B., Baier, B., Munger, B., Paplawsky, B., ... Loh, Z. Multi-laboratory compilation of atmospheric carbon dioxide data for the period 1957-2019; obspack_co2_1_GLOBALVIEWplus_v6.0_2020-09-11 [Data set]. NOAA Earth System Research Laboratory, Global Monitoring Division. https://doi.org/10.25925/20200903, 2020.
- Segers, Tokaya, J. & Houweling, S. Description of the CH4 Inversion Production Chain. 1125 https://atmosphere.copernicus.eu/sites/default/files/2021-01/CAMS73 2018SC3 D73.5.2.2-2020 202012 production chain Ver1.pdf, 2020a.
- Segers, A., Tokaya, J., Houweling, S.: Validation of the CH4 surface flux inversion reanalysis 1990-2019, Copernicus Atmosphere Monitoring Service EQC report, Dec 2020, https://atmosphere.copernicus.eu/sites/default/files/2021-1130 02/CAMS73 2018SC2 D73.2.4.1-2020 202012 validation CH4 1990-2019 v2.pdf, 2020b.
 - Stevenson, D., Derwent, R., Wild, O., and Collins, W.: COVID-19 lockdown NOx emission reductions can explain most of the coincident increase in global atmospheric methane, Atmos. Chem. Phys. Discuss. [preprint], https://doi.org/10.5194/acp-2021-604, in review, 2021.
- Sha, M. K., Langerock, B., Blavier, J.-F. L., Blumenstock, T., Borsdorff, T., Buschmann, M., Dehn, A., De Mazière, M., Deutscher, N. M., Feist, D. G., García, O. E., Griffith, D. W. T., Grutter, M., Hannigan, J. W., Hase, F., Heikkinen, P., Hermans, C., Iraci, L. T., Jeseck, P., Jones, N., Kivi, R., Kumps, N., Landgraf, J., Lorente, A., Mahieu, E., Makarova, M. V., Mellqvist, J., Metzger, J.-M., Morino, I., Nagahama, T., Notholt, J., Ohyama, H., Ortega, I., Palm, M., Petri, C., Pollard, D. F., Rettinger,
 M., Robinson, J., Roche, S., Roehl, C. M., Röhling, A. N., Rousogenous, C., Schneider, M., Shiomi, K., Smale, D., Stremme, W., Strong, K., Sussmann, R., Té, Y., Uchino, O., Velazco, V. A., Vigouroux, C., Vrekoussis, M., Wang, P., Warneke, T., Wizenberg, T., Wunch, D., Yamanouchi, S., Yang, Y. and Zhou, M.: Validation of methane and carbon monoxide from Sentinel-5 Precursor using TCCON and NDACC-IRWG stations, Atmos. Meas. Tech., 14(9), 6249–6304, doi:10.5194/amt-14-6249-2021, 2021.

- Sherlock, V. et al. TCCON data from Lauder (NZ), 125HR, Release GGG2014.R0 (Version GGG2014.R0) [Data set]. https://doi.org/10.14291/TCCON.GGG2014.LAUDER02.R0/11492981, 2014a.
- Sherlock, V., Connor, B., Robinson, J., Shiona, H., Smale, D., and Pollard, D.: TCCON data from Lauder, New Zealand, 120HR, Release GGG2014R0. CaltechDATA, 2014b.
 - Pollard, D., Robinson, J., and Shiona, H.: TCCON data from Lauder, New Zealand, 125HR, Release GGG2014R0. CaltechDATA, 2019.
- 1155 Spahni, R. et al.: Constraining global methane emissions and uptake by ecosystems. Biogeosciences 8, 1643–1665, 2011.
 - Stephens, B. B., G. W. Brailsford, A. J. Gomez, K. Riedel, S. E. Mikaloff Fletcher, S. Nichol, and M. Manning. Analy sis of a 39-year continuous atmospheric CO2 record from Baring Head, New Zealand. Biogeosciences 10, 4: 2683-2697, doi:10.5194/bg-10-2683-2013, 2013.
 - Strong, K., Roche, S., Franklin, J. E., Mendonca, J., Lutsch, E., Weaver, D., Fogal, P. F., Drummond, J. R., Batchelor, R., & Lindenmaier, R. (2019). TCCON data from Eureka (CA), Release GGG2014.R3 (Version R3) [Data set]. CaltechDATA. https://doi.org/10.14291/TCCON.GGG2014.EUREKA01.R3, 2019.
- Sussmann, R., Forster, F., Rettinger, M., and Bousquet, P.: Renewed methane increase for five years (2007-2011) observed by solar FTIR spectrometry, Atmos. Chem. Phys., 12, 4885-4891, doi:10.5194/acp-12-4885-2012, 2012.
 Sussmann, R., Ostler, A., Forster, F., Rettinger, M., Deutscher, N. M., Griffith, D. W. T., Hannigan, J. W., Jones, N., and Patra, P. K.: First intercalibration of column-averaged methane from the Total Carbon Column Observing Network and the Network for the Detection of Atmospheric Composition Change, Atmos. Meas. Tech., 6, 397-418, doi:10.5194/amt-6-397-2013, 2013.
 - Sussmann, R. & Rettinger, M. TCCON data from Garmisch (DE), Release GGG2014.R2 (Version R2) [Data set]. https://doi.org/10.14291/TCCON.GGG2014.GARMISCH01.R2, 2018a.
- Sussmann, R., M. Rettinger. TCCON data from Zugspitze, Germany, Release GGG2014R1. TCCON data archive, hosted by CaltechDATA, California Institute of Technology, Pasadena, CA, U.S.A. http://dx.doi.org/10.14291/tccon.ggg2014.zugspitze01.R1, 2018b.

Takahashi, T. et al.: Climatological mean and decadal change in surface ocean pCO2, and net sea-air CO2 flux over the global oceans. Deep Sea Research Part II: Topical Studies in Oceanography **56**, 554–577, 2009.

1180

Te, Y., Jeseck, P. & Janssen, C. TCCON data from Paris (FR), Release GGG2014.R0 (Version GGG2014.R0) [Data set]. https://doi.org/10.14291/TCCON.GGG2014.PARIS01.R0/1149279, 2014.

Temperton, C., Hortal, M. & Simmons, A. A.: two-time-level semi-Lagrangian global spectral model. *Quarterly Journal of the Royal Meteorological Society* **127**, 111–127, 2001.

Tsutsumi, Y., H. Matsueda, S. Nishioka, Consistency of the CO2 primary standards in JMA, 12th WMO/IAEA meeting of ex perts on carbon dioxide concentration and related tracers measurement techniques (Toronto, Canada, 15-18 September 2003), Global Atmosphere Watch Report No. 161 (WMO/TD-No.1275), 23-31, 2005.

1190

The Global Observing System For Climate: Implementation Needs, report number 200, World Meteorological Organization, 2016.

Verma,S., Marshall,J., Parrington,M., Agustí-Panareda,A., Massart, S., Chipperfield, M. P., Wilson, C., Gerbig, C.:
Extending methane profiles from aircraft into the stratosphere for satellite total column validation using the ECMWF C-IFS and TOMCAT/SLIMCAT 3-D model, Atmos. Chem. Phys., 17, 6663—6678, 2017.

Warneke, T. et al. TCCON data from Orléans (FR), Release GGG2014.R0 (Version GGG2014.R0) [Data set]. https://doi.org/10.14291/TCCON.GGG2014.ORLEANS01.R0/1149276, 2014.

1200

Waterman, L.S., D. W. Nelson, W.D. Komhyr, T.B. Harris, and K.W. Thoning, Atmospheric carbon dioxide measurements at Cape Matatula, American Samoa, 1976 1984., J. Geophys. Res., 94, 14817 14829, 1989.

Wennberg, P. O. et al. TCCON data from Park Falls (US), Release GGG2014.R1 (Version GGG2014.R1) [Data set]. https://doi.org/10.14291/TCCON.GGG2014.PARKFALLS01.R1, 2017.

Wennberg, P. O. et al. TCCON data from Lamont (US), Release GGG2014.R1 (Version GGG2014.R1) [Data set]. https://doi.org/10.14291/TCCON.GGG2014.LAMONT01.R1/1255070, 2016.

1210 Wennberg, P. O. et al. TCCON data from Caltech (US), Release GGG2014.R1 (Version GGG2014.R1) [Data set]. https://doi.org/10.14291/TCCON.GGG2014.PASADENA01.R1/1182415, 2015.

- Wennberg, P. O., D. Wunch, Y. Yavin, G. C. Toon, J.-F. Blavier, N. Allen, G. Keppel-Aleks. TCCON data from Jet Propulsion Laboratory, Pasadena, California, USA, Release GGG2014R0. TCCON data archive, hosted by CaltechDATA, California Institute of Technology, Pasadena, CA, U.S.A. http://doi.org/10.14291/tccon.ggg2014.jpl01.R0/1149163, 2016.
 - Williams, J. E., Huijnen, V., Bouarar, I., Meziane, M., Schreurs, T., Pelletier, S., Marécal, V., Josse, B., and Flemming, J.: Regional evaluation of the performance of the global CAMS chemical modeling system over the United States (IFS cycle 47r1), Geosci. Model Dev. Discuss. [preprint], https://doi.org/10.5194/gmd-2021-318, in review, 2021.
- Worthy, D.E., K.Higuchi, and D. Chan. North American influence on atmospheric carbon dioxide data collected at Sable Island, Canada. Tellus 55B, 105-114, 2003.

1230

- Wunch, D., G. C. Toon, J.-F. L. Blavier, R. A. Washenfelder, J. Notholt, B. J. Connor, D. W. T. Griffith, V. Sherlock, and P.
 O. Wennberg. The Total Carbon Column Observing Network, Philos. Trans. R. Soc. A Math. Phys. Eng. Sci., 369(1943), 2087–2112, doi:10.1098/rsta.2010.0240, 2011.
 - Wunch, D., G. C. Toon, V. Sherlock, N. M. Deutscher, C. Liu, D. G. Feist, and P. O. Wennberg. The Total Carbon Column Observing Network's GGG2014 Data Version, Pasadena, California, 2015.
 - Wunch, D., J. Mendonca, O. Colebatch, N. Allen, J.-F. L. Blavier, S. Roche, J. Hedelius, G. Neufeld, S. Springett, D. Worthy, R. Kessler, K. Strong. TCCON data from East Trout Lake, Canada, Release GGG2014R1. TCCON data archive, hosted by CaltechDATA, California Institute of Technology, Pasadena, CA, U.S.A. https://doi.org/10.14291/tccon.ggg2014.easttroutlake01.R1, 2018.
 - Zander, R., Mahieu, E., Demoulin, P., Duchatelet, P., Roland, G., Servais, C., Mazière, M. De, Reimann, S. and Rinsland, C. P.: Our changing atmosphere: Evidence based on long-term infrared solar observations at the Jungfraujoch since 1950, Sci. Total Environ., 391(2–3), 184–195, doi:10.1016/j.scitotenv.2007.10.018, 2008.
- 1240 Zhou, M., Langerock, B., Vigouroux, C., Sha, M. K., Ramonet, M., Delmotte, M., Mahieu, E., Bader, W., Hermans, C., Kumps, N., Metzger, J.-M., Duflot, V., Wang, Z., Palm, M. and De Mazière, M.: Atmospheric CO and CH4 time series and seasonal variations on Reunion Island from ground-based in situ and FTIR (NDACC and TCCON) measurements, Atmos. Chem. Phys., 18(19), 13881–13901, doi:10.5194/acp-18-13881-2018, 2018.