Disentangling methane and carbon dioxide sources and transport across the Russian Arctic from aircraft measurements

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Abstract. A more accurate characterization of the sources and sinks of methane (CH_4) and carbon dioxide (CO_2) in the vulnerable Arctic environment is required to better predict climate change. A large-scale aircraft campaign took place in September 2020 focusing on the Siberian Arctic coast. CH_4 and CO_2 were measured in situ during the campaign and form the core of this study. Measured ozone (O_3) and carbon monoxide (CO) are used here as tracers. Median CH_4 mixing ratios are fairly higher than the monthly mean hemispheric reference (Mauna Loa, Hawaii, US) with 1890-1969 ppb vs 1887 ppb respectively, while CO_2 mixing ratios from all flights are lower (408.09-411.50 ppm vs 411.52 ppm). We also report on three case studies. Our analysis suggests that during the campaign the European part of Russia's Arctic and Western Siberia were subject to long-range transport of polluted air masses, while the East was mainly under the influence of local emissions of greenhouse gases. The relative contributions of the main anthropogenic and natural sources of CH_4 are simulated using the Lagrangian model FLEXPART in order to identify dominant sources in the boundary layer and in the free troposphere. In western terrestrial flights, air masses composition is influenced by emissions from wetlands and anthropogenic activities (waste management, fossil fuel industry and to a lesser extent the agricultural sector), while in the East, emissions are dominated by freshwaters, wetlands, and the oceans, with a likely contribution from anthropogenic sources related to fossil fuels. Our results highlight the importance of the contributions from freshwater and ~~oceans~~ocean emissions. Considering the large uncertainties associated to them, our study suggests that the emissions from these aquatic sources should receive more attention in Siberia.

1 Introduction

The increasing greenhouse gas burden in the atmosphere led to a global surface temperature rise ~~of 1.09 °C in~~. In the last decade (2011-2020~~) compared to~~), global surface temperature was 1.09 °C higher than the last decades of the 19th century (1850-1900) (Masson-Delmotte et al., 2021). The global mean mixing ratio of carbon dioxide (CO_2) reached 413.2±0.20 ppm in 2020 (WMO, 2021). Anthropogenic CO_2 sources are dominated by fossil fuel combustion and cement production (9.6±0.5

35   GtC yr^{-1}) and land-use change (1.6±0.7 GtC yr^{-1}). While these emissions are steadily increasing, the ~~mains~~main sinks of CO_2, terrestrial vegetation (3.4±0.9 GtC yr^{-1}) and oceans (2.5±0.6 GtC yr^{-1}) have taken up a rather stable proportion (about 56% per year) of emissions from human activities albeit with regional differences (Friedlingstein et al., 2020; Masson-Delmotte et al., 2021).

Methane (CH_4), the second most abundant anthropogenic greenhouse gas, reached 1889±2 ppb in the atmosphere in 2020
40   (WMO, 2021), but has a global warming potential about 32 times higher than CO_2 on a 100-year horizon (Etminan et al., 2016). Since 2007, the CH_4 mixing ratio has been constantly increasing, up to 8-9 ppb yr^{-1} corresponding to an increase of the atmospheric burden of about 25 Tg CH_4 yr^{-1} (Platt et al., 2018). A total of 576 Tg CH_4 yr^{-1} has been injected into the atmosphere during the 2008-2017 decade combining anthropogenic and natural sources (Saunois et al., 2020). Anthropogenic emissions represent 60% of global total methane emissions with agriculture and waste management (191 to 223 Tg CH_4 yr^{-1}) and fossil
45   ~~fuels~~fuel exploitation (113 to 154 Tg CH_4 yr^{-1}) (Saunois et al., 2020). On a global scale, wetlands are the largest natural methane emission source (153 to 227 Tg CH_4 yr^{-1}) (Kirschke et al., 2013; Lan et al., 2021). Emissions from freshwaters have estimates ranging widely between 60 and 180 Tg CH_4 yr^{-1} (Saunois et al., 2016) and are mainly driven by diffusion, ebullition and release from bubble storage (Matthews et al., 2020). Fluxes from aquatic sources may be underestimated (Rosentreter et al., 2021). Ocean emissions by diffusion and ebullition are estimated to range between 2 and 40 Tg CH_4 yr^{-1}. However, these
50   estimated CH_4 emissions could potentially be higher due to large gas hydrate reservoirs on the seabed of Arctic Ocean (Platt et al., 2018). Widely debated emissions attributed to marine sources including methane hydrates dissociation have already been observed in the region (Shakhova et al., 2010; Berchet et al., 2016; Platt et al., 2018; Berchet et al., 2020; Thornton et al., 2020; Steinbach et al~~.,~~, 2021).

Siberia and the Russian Arctic are significant contributors to the CH_4 budget. Half of Siberian emissions originate from fossil
55   fuel exploitation (about 17 Tg CH_4 yr^{-1}), one third from natural wetlands (about 13 Tg CH_4 yr^{-1}) and a minor contribution from agriculture and waste (about 5 Tg CH_4 yr^{-1}) for a total average of 38 Tg CH_4 yr^{-1} (Saunois et al., 2016). The multiplicity of the sources and the seasonality of emissions (Berchet et al., 2015; Belikov et al., 2019) influence CH_4 variability over Siberia. Many uncertainties in the CH_4 emissions in Siberia therefore remain (Berchet et al., 2016; Elder et al., 2020; Matthews et al., 2020; Wik et al., 2016; Thornton et al., 2016). Uncertainties are driven by the multitude of adjacent sources present in the
60   region: natural wetlands, freshwaters (lakes, ponds and streams), oceans and anthropogenic activities (leaks from increasing oil and gas extraction and transport, as well as regional agriculture and waste management) and biomass burning. In addition to the currently dominant sources, the thawing of continental and submarine permafrost could release massive amount of carbon in the atmosphere since Siberia is considered to be one of the world's largest terrestrial carbon reservoirs (Belikov et al. 2019). Soils in the permafrost region retain twice as much carbon as the atmosphere does (Turetsky et al., 2019).

65   To reduce uncertainties at the scale of Siberia, atmospheric measurements have been performed at tower observatories (Belikov et al., 2019; Sasakawa et al., 2010; Sasakawa et al., 2017; Fujita et al., 2020), during oceanographic campaigns (Thornton et al., 2016; Berchet et al., 2020; Steinbach et al., 2021) or even by train (Skorokhod et al., 2016). The YAK-AEROSIB project

organizes intensive campaigns since 2006 across all of Siberia (Paris et al., 2008; Paris et al., ~~2010~~2010a). These annual campaigns provide measurements of atmospheric concentrations of CO_2, CH_4, O_3, carbon monoxide (CO) and aerosols with

70 an objective to better understand regional sources, dynamical processes and long-range transport in Siberia (Paris et al., 2008; Berchet et al., 2013). In situ measurements have the potential to document carbon sources and sinks in Siberia, and therefore to ultimately establish a more accurate ~~estimations~~estimation of future greenhouse gas trajectories. The previous campaigns have highlighted several key mechanisms in this poorly-studied region by representing vertical profiles of several greenhouse gases to determine the origin and impact of polluted air masses (Paris et al., 2008), identifying the source of upper tropospheric

75 O_3 depletion (Berchet et al., 2013) or also bringing new insights on specific events like the extensive wildfires that occur in Siberia during summer (Paris et al., 2009; Antokhin et al., 2018). In the present work, we analyze data from the last YAK-AEROSIB campaign that took place during September 2020 with a specific focus on Northern Russia and the coastal Arctic Ocean. This campaign included 13 flights across Russia with dedicated flights over the northern seas of Barents, Kara and Laptev, and the very East of the region with Bering Strait, the East Siberian Sea and the Chukchi Sea.

80 The present study focuses on the distribution of CH_4 and CO_2. This paper aims more specifically to identify and quantify the respective contributions from regional sources of CH_4 during the first half of September 2020 in the Russian Arctic. We document the measurement and data collected during the campaign. Ozone and carbon monoxide are used as tracers for the interpretation of CO_2 and CH_4 mixing ratio variability. Tropospheric ozone is a harmful pollutant produced from precursors including CO and CH_4 (Saunois et al., 2016), while CO is a pollutant and a minor greenhouse gas produced by combustion

85 processes.

In Sect. 2, we first describe the study area and the in-situ instrumentation for continuous measurements of CO_2, CH_4, O_3, and CO. We also explain our approach and the inputs to determine the contributions of each CH_4 source to the measurements made during the campaign, which is mainly achieved by combining the Lagrangian FLEXible TRAjectory model (FLEXPART) of transport and diffusion, with methane flux inventories. In Sect. 3 we present the 47 vertical profiles taken during the campaign

90 with an emphasis on CO_2 and CH_4 mixing ratios. We then ~~describe~~focus on four individual vertical profiles to characterize atmospheric transport of three ~~case studies based on specific signatures in species variability.~~different regions. We also discuss how ~~this~~the data ~~coincides~~coincide with simulations of CH_4 enhancements linked to anthropogenic and natural sources to give insights on the main CH_4 contributions at the time of the flight. We conclude by discussing the respective importance of anthropogenic activities and aquatic sources on CH_4 variability over Western Russia and Northeastern Siberia.

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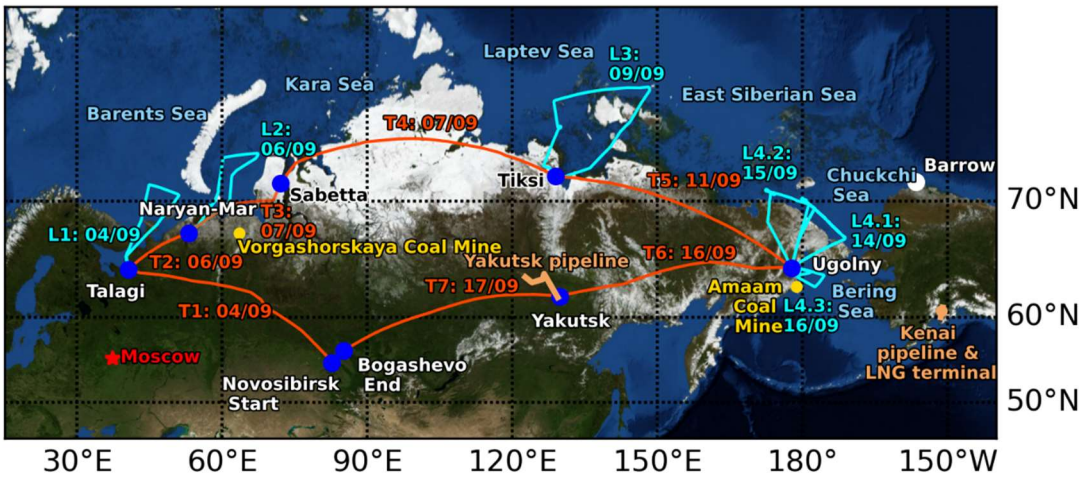
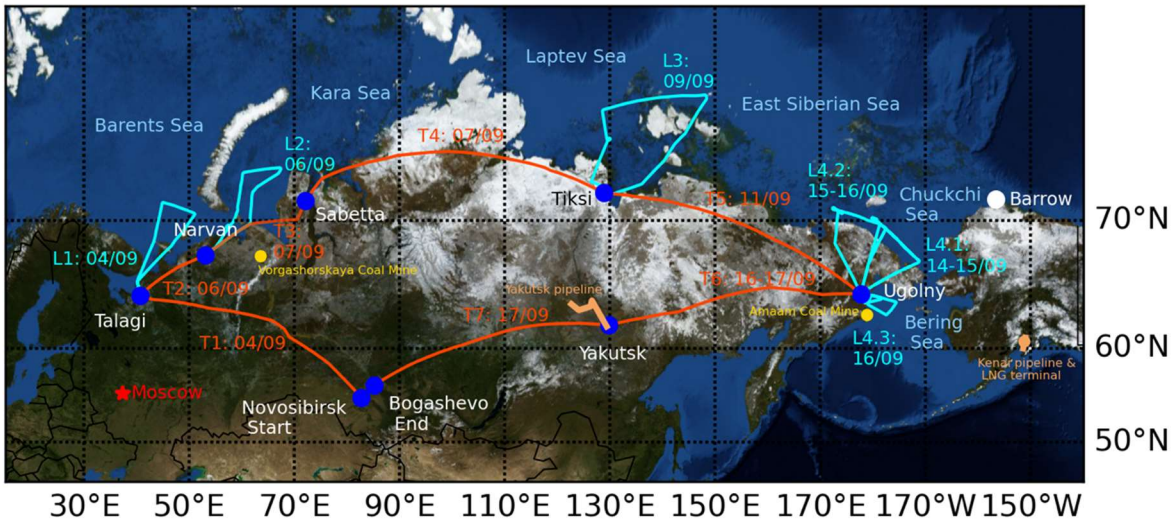
2 Methods

2.1 Description and synoptic situation of the campaign

The flight route of the September 2020 campaign is shown in Fig. 1 and the campaign is described in Table 1. An overview of the campaign can be found in Belan et al. (2022). The aircraft used for the campaign was the "Optik" Tupolev-134A-3M (SKh) operated by the V.E. Zuev Institute of Atmospheric Optics (Tomsk, Russia) (~~Anokhin~~Antokhin et al., ~~2011~~2018; Belan et al., 2022). The maximum range of the plane is 3000 km with an average observed airspeed of 172 m s^{-1} and a vertical speed of 6.5 m s^{-1} for both ascents and descents. The plane configuration is illustrated in Fig. 2 and more details can be found in ~~Antokhin~~Belan et al. (~~2011~~2022).

The campaign took place between September 4th and 17th, 2020. The period of the campaign was chosen to be during summer to have measurements when the ocean is open. While September is not expected to be a period of maximum emissions, it presents minimum sea ice (Fig. A1) and maximum thaw. Thirteen different flights covered Russia from 55° N to 76° N between 40° E and 172° W for a cumulative ground distance of 25000 km with altitudes up to 11 km. The flights are grouped in two different categories: 1) loops above lands and oceans to explore specific environments with flight legs at typical altitudes of 200, 500 and 5000 m (these flights are referred to with a prefix "L",), and 2) transit flights between airports (prefix "T"). The campaign started in Novosibirsk and ended in Bogashevo (Tomsk airport). The first flights (T1, L1, T2, L2 and T3) crossed the cities of Arkhangelsk and Naryan-Mar and encompassed different biomes such as taiga (boreal forest), extensive wetlands and steppe regions (Belikov et al., 2019). Then the plane travelled along the Northern shores of the Russian Federation (T4, L3 and T5), above section of the Arctic Ocean in the seas of Kara, Laptev and the East Siberian Sea, crossing the small towns of Sabetta and Tiksi. The Arctic Ocean is at its lowest annual sea-ice extent in September and September 2020 was the second record low in Arctic sea-ice extent after 2012 (Fetterer et al., 2017). The seas that were covered by the campaign were essentially free of ice during the campaign (Fig. ~~B1~~A1). The East Siberian Arctic Shelf (ESAS) contains up to 40% of Arctic marine permafrost (Ruppel, 2015). Several loops (L4.1, L4.2, and L4.3) were carried out in the Russian Far East characterized by Arctic deserts, tundra, and forest tundra for the northern regions while the south is dominated by forests (Petäjä et al., 2021). All these loops used the airport of Ugolny (Anadyr). The final transit flights (T6 and T7) crossed regions covered by forests of coniferous trees and by agricultural lands to the South (Bartalev et al., 2003), and landed at the airports of Yakutsk and Tomsk respectively. Large oil and gas (O&G) infrastructures (Yakutsk pipeline, Kenai terminal in Alaska) and coal mines (Ammam, Vorgashorskaya) from the Global Monitor Energy (2022) database are displayed on the map.

125 **Figure 1: September 2020 campaign flight plan. The 13 flights are indicated either as "loop" (prefix L, cyan line) or "transit" (prefix T, orange line). Oil and gas (O&G) infrastructures and coal mines are shown in light orange and yellow respectively (data from "Global Coal Mine Tracker" and "Global Fossil Infrastructure Tracker", Global Energy Monitor 2022, under CC BY-NC-SA 4.0 License). Map background: "Blue Marble Next Generation", September 2004 (Stöckli et al., n.d.).**

130 **Table 1: Campaign description**

Flight	Date UTC	Time UTC	Local Time	Airports	Latitude	Longitude
T1	04 Sept. 20	06:55-11:01	13:55-15:01	Novosibirsk North-Talagi	55-65° N	83-41° E
L1	04 Sept. 20	12:58-17:06	15:58-20:06	Talagi	65° N	41° E
T2	06 Sept. 20	08:07-09:31	11:07-12:31	Talagi-Naryan Mar	65-68° N	41-53° E

L2	06 Sept. 20	10:55-14:36	13:55-17:36	Naryan Mar	68° N	53° E
T3	07 Sept. 20	05:01-06:27	08:01-11:27	Naryan Mar-Sabetta	68-71° N	53-72° E
T4	07 Sept. 20	08:14-10:48	11:14-19:48	Sabetta-Tiksi	71-72° N	72-129° E
L3	09 Sept. 20	03:09-06:57	12:09-15:57	Tiksi	72° N	129° E
T5	11 Sept. 20	01:48-05:00	10:4817:00	Tiksi-Ugolny	72-65° N	129-178° E
L4.1	14-15 Sept. 20	23:28-02:57	11:28-14:57	Ugolny	65° N	178° E
L4.2	15-16 Sept. 20	22:32-01:42	10:32-13:42	Ugolny	65° N	178° E
L4.3	16 Sept. 20	03:38-05:51	15:38-17:51	Ugolny	65° N	178° E
T6	16-17 Sept. 20	23:16-02:39	11:16-11:39	Ugolny-Yakutsk	65-62° N	178-130° E
T7	17 Sept. 20	03:57-07:49	14:57-14:49	Yakutsk	62° N	130° E[CN1]

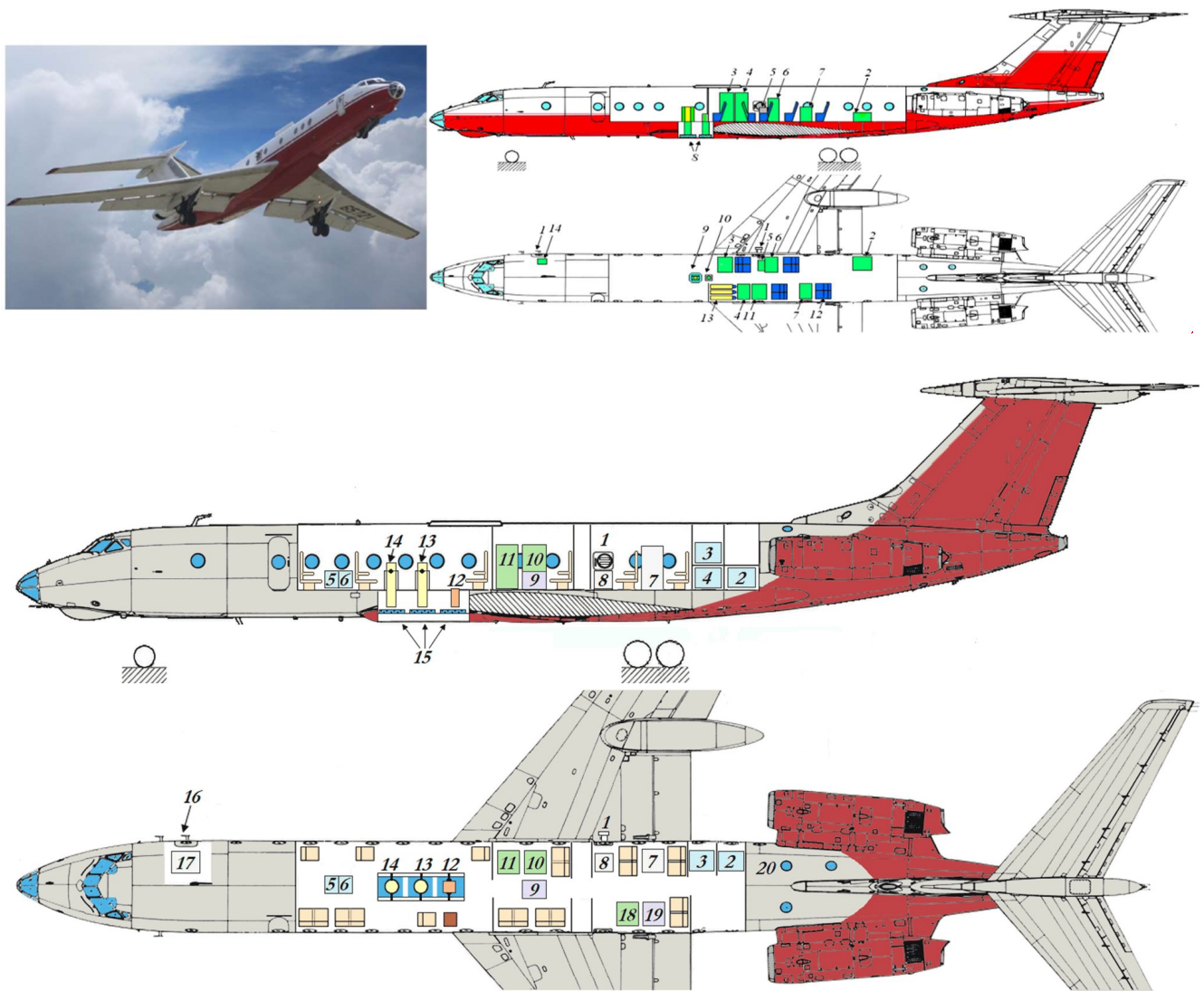



Figure 2: Arrangement of the instrument suite on board ~~of~~ the OPTIK TU-134 aircraft laboratory: 1 – ambient air inlets and RH&T probe; 2 – aircraft electrical power distribution unit (28 VDC); 3, 4, 5, 6 – inverters (28 VDC/220 VAC) and UPSs (Delta RT-2K); 7 – aerosol instrument rack: ~~aethalometer~~Aethalometer (MDA-02) and photoelectric aerosol nephelometer (FAN-M); 8 – aerosol instrument rack: diffusional particle sizer (DPS), optical particle counter (Grimm 1.109), filter, and bioaerosol sampling suite; 9 – navigation system (CompaNav-5.2 IAO); 10 – gas analysis rack: CO2/_CH4/_H2O (Picarro G2301-m): 11 – gas analysis rack: O3 (TEI ~~Model~~model 49C), CO (TEI ~~Model~~model 48C); 12 – spectroradiometer (Spectral Evolution PSR-1100F); 13, 14 – aerosol lidars; 15 – camera hatches; 16 – ambient air inlet; 17 – sampling unit for organic aerosol analysis; 18 – gas analysis rack: ~~NOX~~NOx (Thermo Scientific ~~Model~~model 42i-TL); 19 – main data acquisition system (NI PXI-1042); 20 – GLONASS/GPS antennas. ~~Copyright to Mikhail Yu. Arshinov~~Figure and caption by Belan et al. (2022).

Figure 3 shows the geopotential height at 500 hPa as provided in the dataset ERA5 from the European Centre for Medium-Range Weather Forecasts; (ECMWF; Hersbach, 2018) on the 6th and the 15th of September. At the beginning of the campaign (Fig. 3a), there ~~was~~were a low-pressure system over central Siberia (80° E, 60° N) and a high-pressure ridge coming from Western Europe and going in a north-easterly direction to the seas of Barents and Kara. The period was characterized by an "Omega" whirling around troughs over Eastern Europe and eastern Asia that can be seen distinctly at the end of the campaign (Fig. 3b). The combination of these two events lifted an airmass most likely affected by western European emissions into the free troposphere over Siberia during the campaign. This may have influenced greenhouse gases concentrations as measured during the campaign. Over Eastern Siberia (180° E, 55° N), there was a low-pressure system and winds blown from western Alaska to Bering Strait (Fig. 3b).

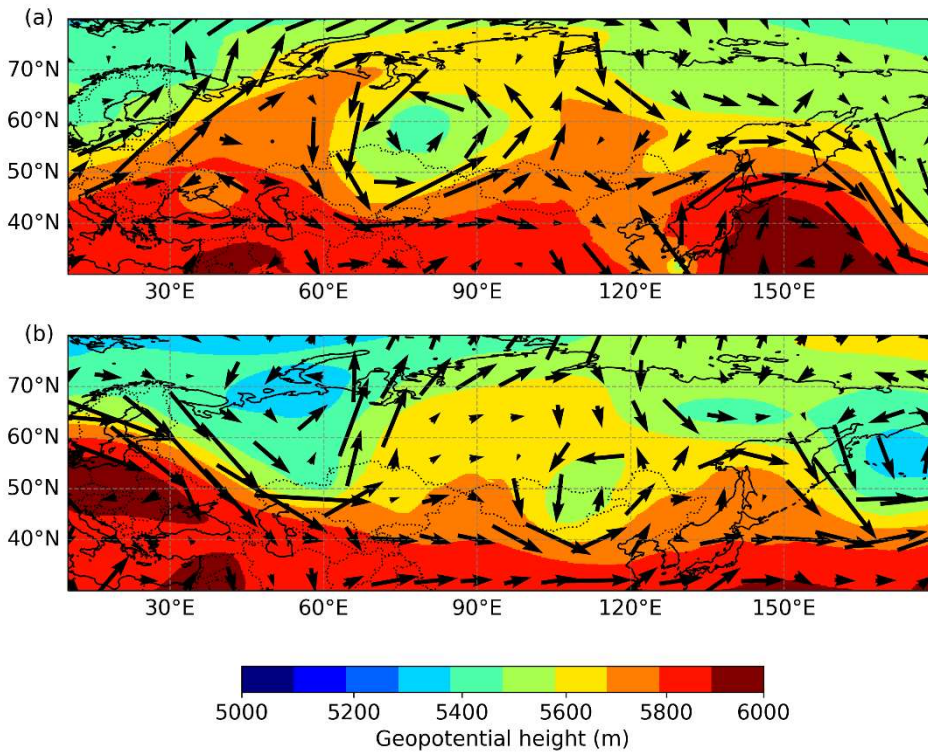



Figure 3: (a) Geopotential height and wind (speed and direction) at 500 hPa from ECMWF reanalyses for 06 September at 13:00 UTC. (b) Same as (a) for 15 September at 01:00 UTC

A series of 47 vertical profiles was collected, each lasting between 20 and 30 min. Loop flights were composed of three ascending and descending profiles. Transit flights had only vertical profiles during take-off and landing, and took place at high altitude (about 10 km) in between. The plane stayed at horizontal plateaus when top altitudes were reached between 9000 m and 11000 m, as well as close to the ground at about 200 m and some on descents at 5000 m and at 500 m. ~~60% of the~~

measurements were taken above the Arctic Circle (> 66° N). About 80% of the data was acquired above 2000 m which is generally above the planet boundary layer (BL) at these latitudes and time of the year. Finally, 60% of the measurements were taken above North of the Arctic Circle (> 66° N).

2.2 Instrumentation of the campaign

165 For in situ analysis air is sampled from inlets. The length of the tubes between the inlets and instruments was 34.8 m. ConcentrationsMixing ratios of CO_2 and CH_4 were measured by a Picarro G2301-m GHG analyzer using Cavity Ring-Down Spectroscopy. It is a modification of the G2301 model specially conceived for aircraft measurements which is designed to minimize effects induced by aircraft vibration and roll, and suitable for rapidly changing altitude (up to 1000 m min^{-1}) and temperature (up to 15 °C hr^{-1}) conditions.). It has an acquisition rate of 1 Hz and a precision of <0.20 ppm for CO_2 and <1.5

170 ppb for CH_4. Water vapor mole fraction is also quantified to include internal water dilution correction and thereby express dry air mole fractionsmixing ratios of carbon dioxide and methane. Three calibration gases are transported in high-pressure cylinders with respective CO_2 mixing ratios of 370.91±0.005 ppm, 390.33±0.009 ppm and 430.30±0.006 and CH_4 mixing ratios of 1814.35±0.150 ppb, 1960.99±0.094 ppb, and 2205.01±0.106 ppb. Their values were determined before the campaign at Laboratoire des Sciences du Climat et de l'Environnement (LSCE) according to World Meteorological Organization (WMO)

175 standards (Paris et al., 2008). The drift is then corrected according to the difference between the measurements and the calibrated value of the reference gas. The whole campaign has been sampled at an interval of 1 s without data gaps, except for the transit T1 (the first of the series) during which the instrument software "froze" for the first half of the flight and it was restarted.was restarted after. The calibrated measurements can be compared to values of stations considered as reference sites for species measured during September 2020:

180   -   The calibrated measurements can be compared to values of reference of stations at Mauna Loa, Hawaii (19.54° N, 155.58° W, 3397 m a.s.l) (Oltmans and Levy 1994; Dlugokencky et al., 2021a; Dlugokencky et al., 2021b) and Mace Head, Ireland (53.20° N, 09.54° W, 25 m a.s.l) (Hazan et al., 2016) that can be considered as background sites for CO_2, CH_4, CO (only Mace Head) and O_3 (only Mauna Loa) concentrations during September 2020. Measurements from for CO_2, CH_4 and O_3. Barrow, Alaska (71.32° N, 156° W, 11 m a.s.l, see Fig. 1) were also used for comparison.

185   -   Mace Head marine sector, Ireland (53.20° N, 09.54° W, 25 m a.s.l) (Hazan et al., 2016) for CO_2, CH_4 and CO. Marine sector means that a selection on data based on standard deviation, wind speed and wind direction has been made to only keep air masses from westerly sectors (Biraud et al., 2000).

   -    Barrow, Alaska (71.32° N, 156° W, 11 m a.s.l, see Fig. 1) for CO_2 and O_3.

Ozone was measured by a commercial fast response ozone analyzer (Thermo Environmental Instruments Model 49C USA),

190 with modifications for internal calibration and aircraft operation safety. It is based on UV absorption in two parallel cells and has a precision of 2 ppb for an integration time of 4 s. In practice, values have been acquired with periods that vary between 4 and 10 s. Air is pressurized prior to the detection by a Teflon KNF Neuberger pump model N735 also used for CO. Before the

campaign, the instrument has been calibrated against a NIST related reference calibrator Model49PS and verification of the O_3 analyzer have been made before and after the campaign. The length of the Teflon tubes between the inlet and O_3 and CO analyzers was 2m.

Carbon monoxide is acquired by an instrument based on a commercial infrared absorption correlation gas analyzer (Thermo Electron Model 48C, USA). It has been improved with addition of a periodical in-flight accurate zero measurements, new IR detector, pressure increase and regulation in the absorption cell, increased flow rate to 4 l min^{-1}, water vapor trap and ozone filter. The response time of the instrument is 30 s and it has a precision of 5 ppb, with a lower detection limit at 10 ppb ("YAK-Aerosib Measurements", 2021). For this campaign, the CO data were not post-processed due to technical problems and were therefore of lower quality than in previous studies. We applied a median filter with a kernel size of 75 samples. When computing the statistics for each flight, CO standard deviations were ranging between 10 and 35 ppb for mean ~~mole fractions~~mixing ratios between 73 and 103 ppb. ~~We decided to not use~~In the present study, CO measurements ~~for a quantitative study but~~are ~~almost used~~discussed as ~~a~~ tracer for analysis of the other gases.

To be compared with simulations (see Sect. 3.4), the measured mixing ratios were 1-minutes averages with the background removed. The background is defined for each flight as follows: 1) only CH_4 values in air that had a corresponding O_3 mixing ratio < 70 ppb were kept, to remove influence from air of fresh stratospheric origin. 2) Then the 5th percentile of the remaining data was taken as the background value. Background mixing ratio were calculated flight by flight.

2.3 Back trajectories with the Lagrangian model FLEXPART

We used the Lagrangian particle transport and dispersion model FLEXPART (Stohl et al., 1998; Stohl and Thomson 1999; Stohl et al., 2005; Pisso et al., 2019) for long-range transport analysis and for determining the origin of the polluted air masses measured during the campaign. We adopted the backward ("receptor-oriented") approach of the model suitable when the number of sources is superior to the number of receptors and designed to quantify the remote contributions in a single plume (Seibert and Frank 2004). In our configuration FLEXPART ~~computes~~computed the position of 2000 particles backwards in time, following the atmospheric conditions with a stochastic contribution representing the diffusion (including small scale turbulence not included in averaged meteorological fields) (Fleming, Monks, and Manning 2012). These implementations ~~allow~~allowed a more realistic representation of transport in planet boundary layer. The lifespan of the virtual particles was set to 10 d as the transport precision decreases after 10 days due to accumulated transport errors and insufficient number of particles to represent it (Stohl et al., 1995). The meteorological inputs were ERA5 datasets from ECMWF with a spatial resolution of 1° and a temporal resolution of 3 h covering the north hemisphere, limiting the resolution of the output grid to 0.5° x 0.5°. A simulation was performed for each flight, beginning 10 days before the first acquisition and ending with the last acquisition. Receptor (i.e. aircraft) positions every 1 s during the campaign are aggregated in four dimensions boxes (0.1° x 0.1° x 100 m x 60 s). The potential emission sensitivity (PES) calculated by the model for each receptor position is defined as

the residence time of the backward particles in a grid cell below a threshold altitude (here 500 m and 2000 m). We convolved this output with CH$_4$ flux inventories to get maps of potential sources contributions that are integrated over the Northern Hemisphere to finally get the simulated mixing ratios of each source every minute.

~~To be compared with simulations, the measured mole fractions~~mixing ratios ~~were down-sampled into 1-minute bins and the values were averaged. In addition,~~minutes averages with the background ~~removed. The background is defined as the minimal mole fraction for each flight was removed~~follows: 1) only CH$_4$ ~~values in air that had a corresponding O$_3$ mixing ratio < 70 ppb were kept, in this way influences from each~~stratospheric air ~~were eliminated as much as possible. 2) Then the 5th percentile was taken as the background~~ value. ~~Background mixing ratio were calculated flight by flight. This background is used in Section 3.4 where it is subtracted from measurements~~ to have comparable CH$_4$ ~~calculate measured~~ enhancements.

2.4 Methane flux inventories

For ~~the present study~~our simulations, five categories of CH$_4$ sources were considered~~.~~:

- Emissions of CH$_4$ by anthropogenic activities were characterized with EDGAR v6.0 (Emission Database for Global Atmospheric Research from PBL Netherlands Environmental Assessment Agency) with a grid resolution of 0.1° x 0.1° (Crippa et al., 2019). EDGAR inventories are published as yearly or monthly files. We chose to use the month of September 2018 (the last available) to benefit from the seasonality of the emissions. We aggregated the 21 EDGAR sectors into three categories: agriculture, exploitation and use of fossil fuels and waste management (see grouping definition in Table ~~A1~~B1).

- Biomass burning and wildfires emissions were represented using GFED4.1s (Global Fire Emissions Database) based on van der Werf et al. (2017) at a resolution of 0.25° x 0.25° for the monthly mean of September 2020.

- ~~Wetlands~~Wetland fluxes were based on the process-based model ORCHIDEE as described in Ringeval et al. (2012) at a horizontal resolution of 0.5 x 0.5° for September 2017 only for the North Hemisphere provided to Global Carbon Project 2019 (Melton et al., 2013; Wania et al., 2013; Zhang et al., 2021).

- The representation of ~~freshwaters~~freshwater CH$_4$ fluxes, based on the work of Thonat et al. (2017), combined two inventories: the lake biogeochemical model (Blake4me) (Tan et al., 2015) for sources above 60° N and Global Lakes and Wetlands Database (GLWD3) distribution for the rest of the Northern Hemisphere (Lehner and Döll, 2004), with a resolution of 0.25° x 0.25° based on data from 1971 to 2013. [CN2][CN3]

- The ocean fluxes were based on the work of Weber et al. (2019), a synthesis of measurements made between 1980 and 2016 with an output grid in resolution 0.25° x 0.25° from the MarinE MethanE and NiTrous Oxide (MEMENTO) database (Kock and Bange, 2015). The total marine fluxes are provided as diffusive fluxes from sediments to the surface and ebullition of bubbles released from bottom sediments that reached the atmosphere. We combined the two fluxes in one emission map.

When it was possible, we preferred using inventories averaged on September instead of inventories averaged on several months to put an emphasis on the seasonality of emissions. All inventories were re-gridded to 0.5° x 0.5° in order to be aligned with the grid of the FLEXPART footprints. Figure ~~A1~~B1 illustrates the fluxes indexed by inventories from fossil fuel ~~exploitation~~emissions, agriculture, wetlands, freshwaters, and oceans on Northern Hemisphere.

3 Results

3.1 Distribution of CH_4 and CO_2 during the campaign

Hereafter we refer to "low altitude" and "high altitude" for data that are respectively acquired below and above an arbitrary altitude threshold of 2000 m. The majority (80 %) of the data have been measured at "high altitude". "Low altitude" measurements are analyzed separately to highlight sensitivity to local or regional sources. Reported uncertainties are 1σ uncertainties.

3.1.1 CH_4 mixing ratios

The average CH_4 mixing ratio of the campaign is 1946±45 ppb. This is higher than the monthly mean mixing ratio measured at Mauna Loa 1887 ppb used as Hemispheric reference. Figure ~~4b~~4a shows that the median value of each flight (except for T6) is above the Mauna Loa mean value. At low altitude, the average ~~mole fraction~~mixing ratio is higher with a value of 2011±33 ppb and so exceed ~~both~~ mixing ratios observed at Mauna Loa ~~and~~, Mace Head ~~(1979~~marine sector (1956 ppb) and Barrow (1991 ppb). This suggests the possible influence of significant regional methane sources. ~~This observation is emphasized by the presence of wide interquartile ranges (especially~~The highest mixing ratio ~~rre observed in Western Arctic flights (for the first flights T1, T2, L1 and L2), pointing to strong gradients in CH_4 mole fractions (up to +200 ppb) caused by significant methane emissions close to the ground.~~Many flights present a high CH_4 variability~~.~~. ~~F~~flights in the western Siberian Arctic have the highest 3rd quartile of the campaign (L1, L2 and T2) then followed by the flights in the eastern Arctic (L3, L4.2 and L4.3). ~~This strong CH_4 variability had already been observed during previous campaigns (Paris et al., 2008)~~Large mixing ratio are expected to be encountered in the boundary layer, linked to regional sources, ~~and will be better investigated in the present study~~these will be discussed ~~by using vertical profiles~~ in the Sect. 3.2 by using vertical profiles~~to determine whether the sources are local or not~~. Flights T1 and T6 registered the lowest CH_4 mixing ratios of the campaign (respectively 1812 ppb and 1777 ppb with an aircraft altitude above 10000 m), coming from stratospheric air masses which results in strong gradients for these two flights.

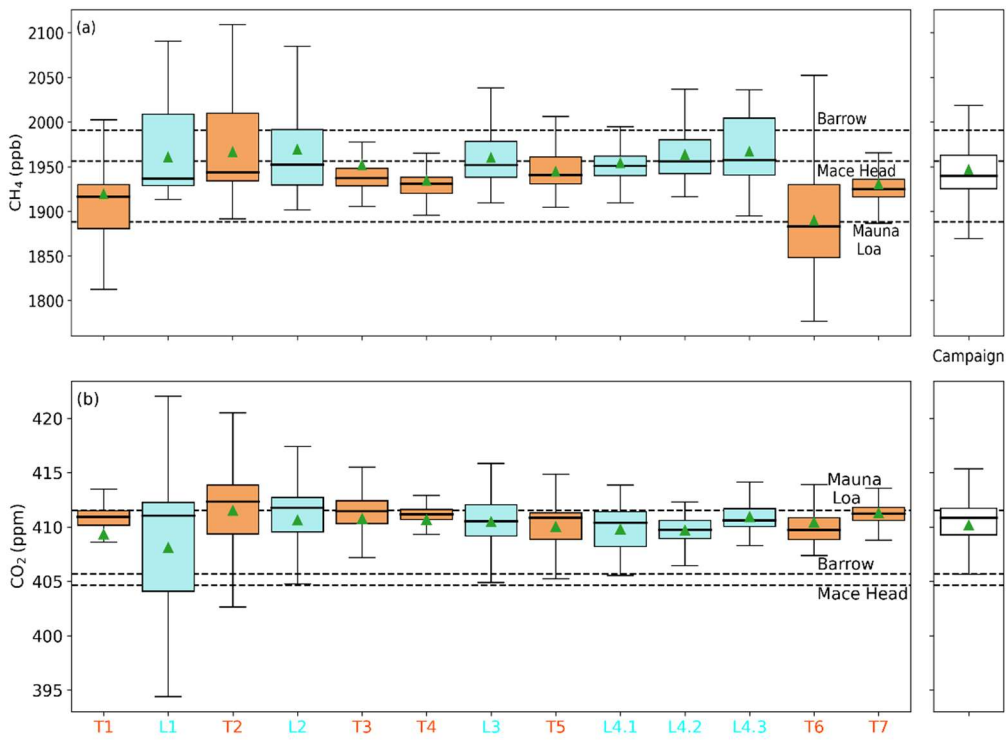
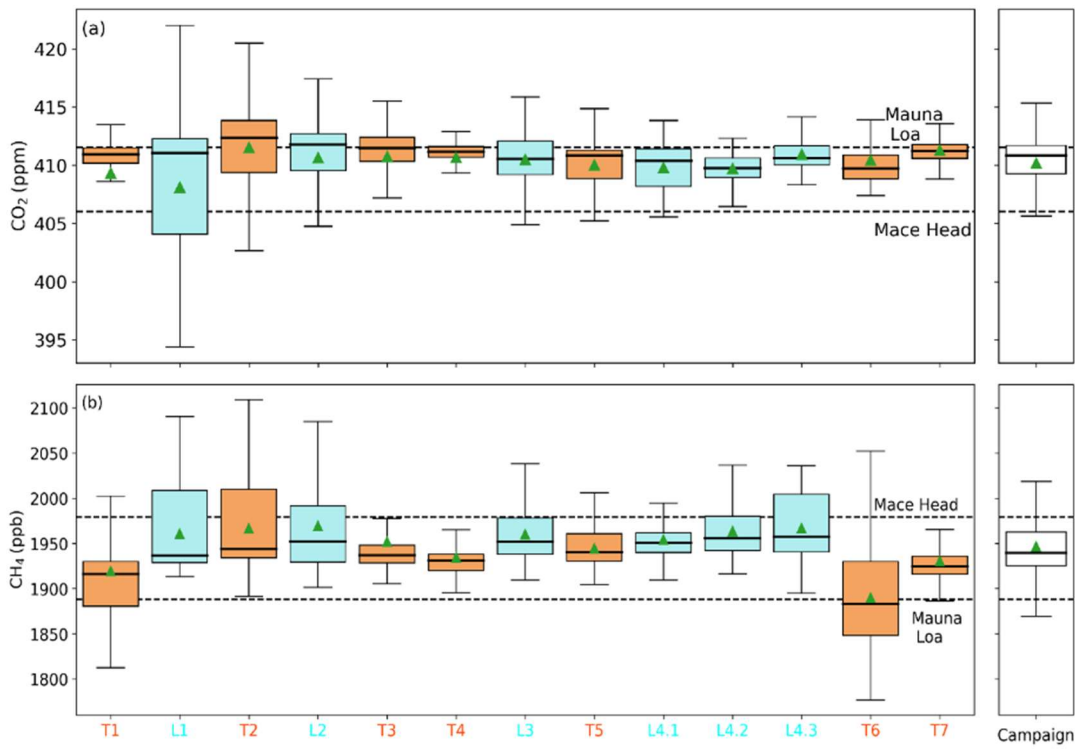



Figure 4: (a) CH₄CO₂ **mixing ratios variability by flight. Statistics for the entire campaign are shown in the right panel. The boxplot shows the interquartile range (25-75%) and the median value. The whiskers extend to the lowest and highest values, ignoring outliers beyond 1.5 times the interquartile range. Flight means are shown as green triangle. Dashed lines show respectively the mean monthly mixing ratios at Mauna Loa and Mace Head marine sector during September 2020. (b) Same as (a) for** CH₄CO₂**.**

Generally, loop flights exhibit a higher first quartile than transit flights for low altitude values. These specific flights have indeed longer low-altitude legs and overall more frequent passages at low altitude (three ascent and descent for loops versus one ascent and one descent only for transits). High CH₄ concentrations in narrow enhancements close to the seas might illustrate the contribution of regional marine and coastal sources. This will be further discussed in Sect. 3.3.

3.1.2 CO_2 mixing ratios

The average CO_2 mixing ratio of the campaign is 410.17±3.29 ppm, slightly below the seasonal average at Mauna Loa (411.52 ppm) but higher than the Mace Head value (406.00marine sector and Barrow values (respectively 404.62 ppm and 405.67 ppm). Figure 4a4b presents the mixing ratio statistics for each flight and the whole campaign. Almost all median values are below the Mauna Loa reference, suggesting that air masses observed during the campaign may have intersected with regional CO_2 sinks. Candidate areas potentially explaining the sink include all terrestrial biomes in Russia (Bartalev et al., 2003; Belikov et al., 2019; Petäjä et al., 2021). Flight T1 flying over the Western Siberian taiga exhibits median values at low and high altitude respectively of 397.73 ppm and 410.92 ppm (with standard deviation of 4.30 ppm), highlighting the influence of these sinks. This biospheric uptake by Siberian boreal ecosystems (> 50° N) has already been observed during previous YAK-Aerosib campaigns realized in late-August or early-September. Vertical profiles for western flights at this time of the year present low CO_2 concentrations close to the surface jointly with a positive strong gradient when the altitude is increasing (Paris et al., 2008; Paris et al., 2010a). Paris et al. (2010b) demonstrated based on the relation between measured CO_2 concentration and airmasses' residence time in the lowest 300 m that for September campaign, the longer the airmasses resided over local areas (boreal and sub-arctic Siberia, > 50° N), the stronger the CO_2 uptake by Siberian ecosystems.

Two flights (L1 and T2) exhibit a large variability compared to the rest. For both flights, there are considerable gradients in CO_2 mole fractionsmixing ratios (+15 ppm and +10 ppm for respectively L1 and T2) between high and low altitudes suggesting the presence of effective sinks (forests and oceans in this case) at low altitude and the presence of "polluted" air masses at high altitude from long-range transport.

The highest mixing ratio measured during the campaign (451.85 ppm) was registered at the end of flight T6 while the aircraft was landing at the airport of Yakutsk. This is most likely driven by local pollution[CN4]. At the time of landing the airport was fogged in (the aircraft was landed with difficulty due to the absence of enough fuel for flying to an alternate airfield).

3.1.3 Comparing CO_2 and CH_4 variabilities

When comparing CO_2 and CH_4 profiles, two different ~~situations~~types of flights appear: 1) both CO_2 and CH_4 present high variability such as in flights T2, L1, L2 or L3 illustrating active regional sources for both species and a strong uptake for CO_2. 2) CH_4 present high variability (with relatively low values) while CO_2 is steady such as in flights T1 and T6. For CH_4, it illustrates the influences of strong regional emissions for the highest values and of influx of stratospheric air for the lowest values. The CO_2 measured is subject to an attenuated vertical propagation of seasonal surface fluxes as demonstrated in Gerbig et al. (2003) leading to this ~~poor~~relatively smaller variability during mostly high-altitude transit flights. ~~Apart from the study on~~To put these limited observations on the greenhouse gas variability in perspective, ~~no other conclusions can be made from this figure as there is no common ground for the reference values between CH_4 and CO_2 statistics, only showing that Siberian territory is subject to complex transfer functions for each gas which needs a vertical representation for a finer analysis~~the following section discusses vertical distribution of mixing ratios.

3.2 Average vertical distribution of the gases

All data were binned every 1000 m in a range of ±500 m to get vertical profiles in Fig. 5. Data were also stratified according to their longitude, separating regions West and East of the 130° E meridian. As reported in Sect. 3.1, western flights exhibit large vertical gradient in mean gas mixing ratios (high minus low altitude means ranging between -64 to -109 ppb, +9.10 to +14.80 ppm and -14 to -29 ppb respectively for CH_4, CO_2, ~~CH_4~~ and O_3) compared to eastern flights.

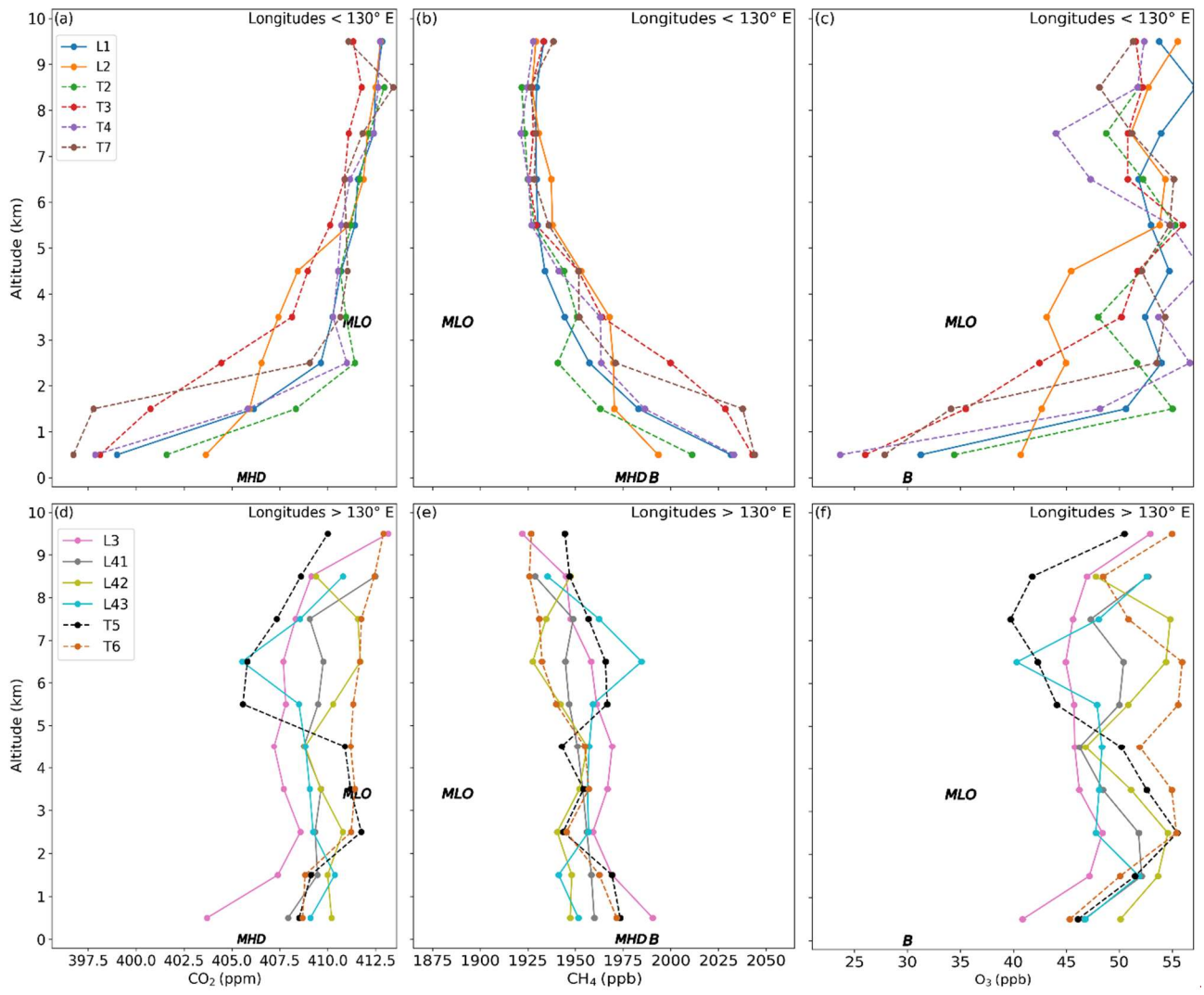
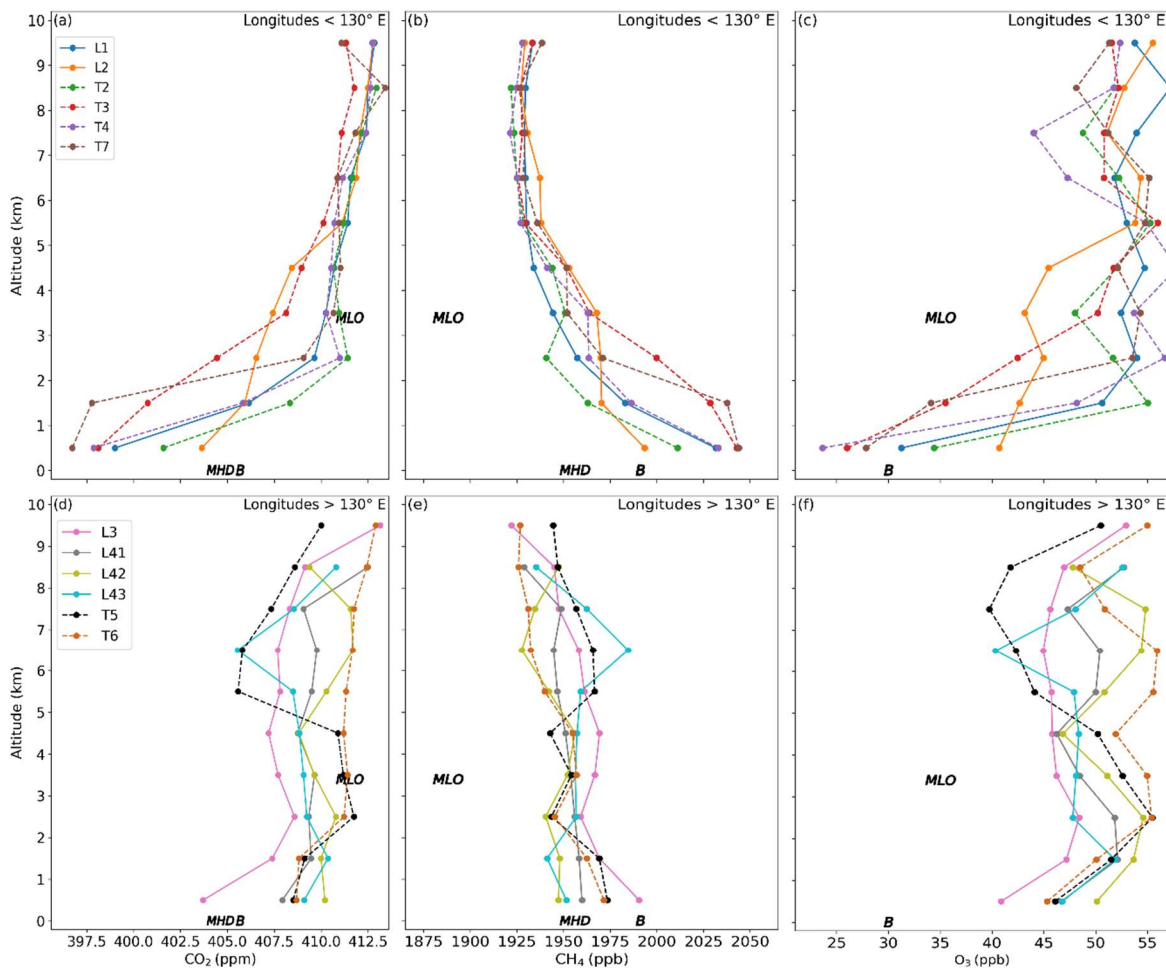



Figure 5: Flight mean vertical mixing ratio profiles for CO_2 (a, d), CH_4 (b, e), and O_3 (c, f). Top row shows flights West of 130°E while bottom row shows flights East of 130° E. Flight T1 is not shown due to missing data. Monthly mean values at Mauna Loa (MLO), Mace Head marine sector (MHD) and Barrow (B) in September 2020 are shown in each panel.

For CH_4 at low altitude, western flights (Fig. 5b) present higher mixing ratios than eastern flights (Fig. 5e) with the biggest values in T3 and T7, closely followed by L1 and T4. ~~These high mole fractions are recorded close to Moscow, western Russian cities and Western Europe with anthropogenic emissions as well as the Vasyugan swamp (wetlands) reported as a serious CH_4 source in Belikov et al. (2019). Influence of such sources is checked in Sect.~~Another feature of western flights is the difference of gradient between low and high altitudes: all western flights except L2 have a much steeper gradient between 0 and 3000 m (ranging from -74 ppb for L1 to -42 ppb for T3 with increasing altitude) than between 3000 and 10000 m (ranging from +11 ppb for L1 to +38 ppb for L2 with increasing altitude). Combining this information with the observation of the biggest CH_4 mixing ratio close to the ground indicates a strong local source that are the Ural wetlands as reported in Belikov et al. (2019).

At high altitude, no conclusion on the origin of the influences can be made at this moment. This will be investigated in Sect. 3.4.1 using the simulated CH_4 enhancements and the footprints. On the opposite, low altitude values for eastern flights (Fig. 5e) are encompassed in a range between ~~1940~~1947 ppb and ~~1960~~1990 ppb. These mixing ratios still represent an enhancement of +60 ppb to + ~~80~~100 ppb compared to the hemispheric ~~background~~reference (Mauna Loa monthly mean) of 1888 ppb. The

350 references of Mace Head marine sector and Barrow respectively at ~~1979~~1956 ppb and 1991 ppb are in the same range as the easternmost flights' low altitude values but below western flights low altitude values. ~~For Eastern~~The simulated enhancements presented in Sect. 3.4.1 are also necessary to identify the sources at the origin of the CH_4 measured here. For flights T5 and L4.3, a layer of elevated CH_4 mixing ratios is crossed in the mid troposphere between approximately 6 and 8 km altitude (Fig. 5e). In Flight L4.3, for example, the vertical CH_4 gradient represents an excess of ~20 ppb between 6500 m and 8500 m

355 compared to the underlying layer (between 4500 m and 6500 m). This increase of CH_4 at top altitudes for flight L4.3 correlates with a decrease of O_3 ~~mole fraction~~mixing ratio of -10 ppb between 7500 m and 8500 m and a decrease in CO_2[CN5][CN6]. The possible origin of this high-altitude O_3 depletion will be investigated in Sect. 3.3.1 using individual vertical profiles that exhibit the same distribution for other flights such as L3 (also T4 or L2 not shown in the study).

O_3 vertical profiles generally increase with altitude, especially in the western flights, and follow a trend opposite to that of CH_4

360 mixing ratios. Eastern flights present an enhancement of +10 ppb to +20 ppb as compared to respective Mauna Loa and Barrow mixing ratios ~~respectively at~~of 35 ppb and 30 ppb for the same altitude range, while western mixing ratios agree with the value from Barrow but above the one from Mauna Loa. O_3 and CH_4 measurements are above the corresponding ~~background~~reference values and therefore reflect regional variability of transport that need deeper investigation.

For CO_2, our measurements are comparable to the Mauna Loa monthly mean (411.52 ppm) in the free tropospheric altitude

365 range of the site (2000-3000 m). Mean CO_2 measured during the western flights in the lowest 1000 m (397-403 ppm) ~~are~~is consistently lower than ~~the~~ Mace Head marine sector and Barrow monthly ~~mean (406~~means (respectively 404.62 ppm~~).~~ and 405.67 ppm). As for CH_4, all western CO_2 vertical profiles (except for L2) have a steeper gradient between 0 and 3000 m (ranging from 6.3 ppm for T3 to 13.1 ppm for T4 with increasing altitude) than between 3000 and 10000 m (ranging from 0.4 ppm for T7 to 3.2 ppm for T3 with increasing altitude) conjointly supporting the ~~observation~~inference made in Sect. 3.1.2 on

370 the strong uptake by Siberian ecosystems. On the opposite ~~the,~~ eastern flights except L3 present mean CO_2 mixing ratios below 1000 m ~~is relatively similar to the~~higher than Mace Head ~~mean~~marine sector and Barrow means. The four highest mean CO_2 mixing ratios close to the ground are collected during eastern profiles (Fig. 5d). These are the three loops of Bering Strait and the transit T6 landing at Yakutsk. For flight T6, half of the low altitude values were taken in the vicinity of the city of Yakutsk and could be subject to local pollution. Concerning the city of Yakutsk, CH_4 ~~and CO as well present~~also presents high mixing

375 ratios at low altitudes ~~for T6~~with values of 1972 ppb and 2045 ppb for respective flights T6 and T7. Highest average mixing ratios ~~of 1972 ppb and 103 ppb,~~at low altitudes for CO are also measured during flights T6 and ~~for~~T7 with ~~respective mixing ratios~~similar values of ~~2045~~103 ppb and 107 ppb respectively. The loops L4.1, L4.2 and L4.3 in remote marine environments

will be investigated in Sect. 3.2.2. Flights L4.3 and T5 present a lower CO_2 layer at higher altitude (respectively 6500 m and 5500 m) already noted previously.

380   These observations show that the atmosphere over Siberia is affected by a complex pattern of influences from local emissions and long-range transport of polluted air masses. It highlights the importance of untangling the different influences on atmospheric composition (Petäjä et al., 2021) that will be further discussed in the next section.

3.3 Individual vertical profiles and regional atmospheric transport

3.3.1 North-Western Russia

385   Figure 6 displays four selected profiles (either ascent or descent) that are representative of vertical distribution of trace gases ~~mole fractions~~mixing ratios encountered during the campaign. Figure 6a is a profile of flight L1 characterized by taiga and ~~wetlands~~wetland environments. The flight L1 was highlighted in Sect. 3.1.1. for having the largest CO_2 interquartile range of the whole campaign. Fig. 6b is a profile of flight L3 close to the ESAS. Fig 6c shows a profile of flight T7 crossing the city of Yakutsk in regions covered by forests of coniferous trees and agricultural lands, and Fig. 6d represents a profile of flight L4.1

390   located near the Bering straits in biomes such as Arctic deserts, tundra, and ~~forests~~forest tundra.

The vertical profile shown in Fig. 6a presents a large CO_2 depletion of 14 ppm in the lowest 2000 m depth layer, highlighting the strong drawdown of CO_2 in BL air. Higher altitude air corresponds to free tropospheric air masses that mostly resided over western European countries and United States to finally arrive in Russia from high altitudes (see footprint in Fig. 7a). These regions are important CO_2 and CH_4 emitters through fossil fuel exploitation and combustion, agriculture and waste

395   management as documented in the EDGAR inventories (Crippa et al., 2019). This shows that the tropospheric air over the area of flight L1 is dominated by the outflow of European BL air interplaying with CO_2 uptake in the BL. Both CH_4 and O_3 profiles present an inversion corresponding to a "chemical" BL around 1000 m (with gradients respectively -90 ppb and +25 ppb). Below 3000 m, CH_4 is anti-correlated with O_3 while the two gases present common features above this altitude, with thin positively correlated layers at 4000 m and 5000 m.

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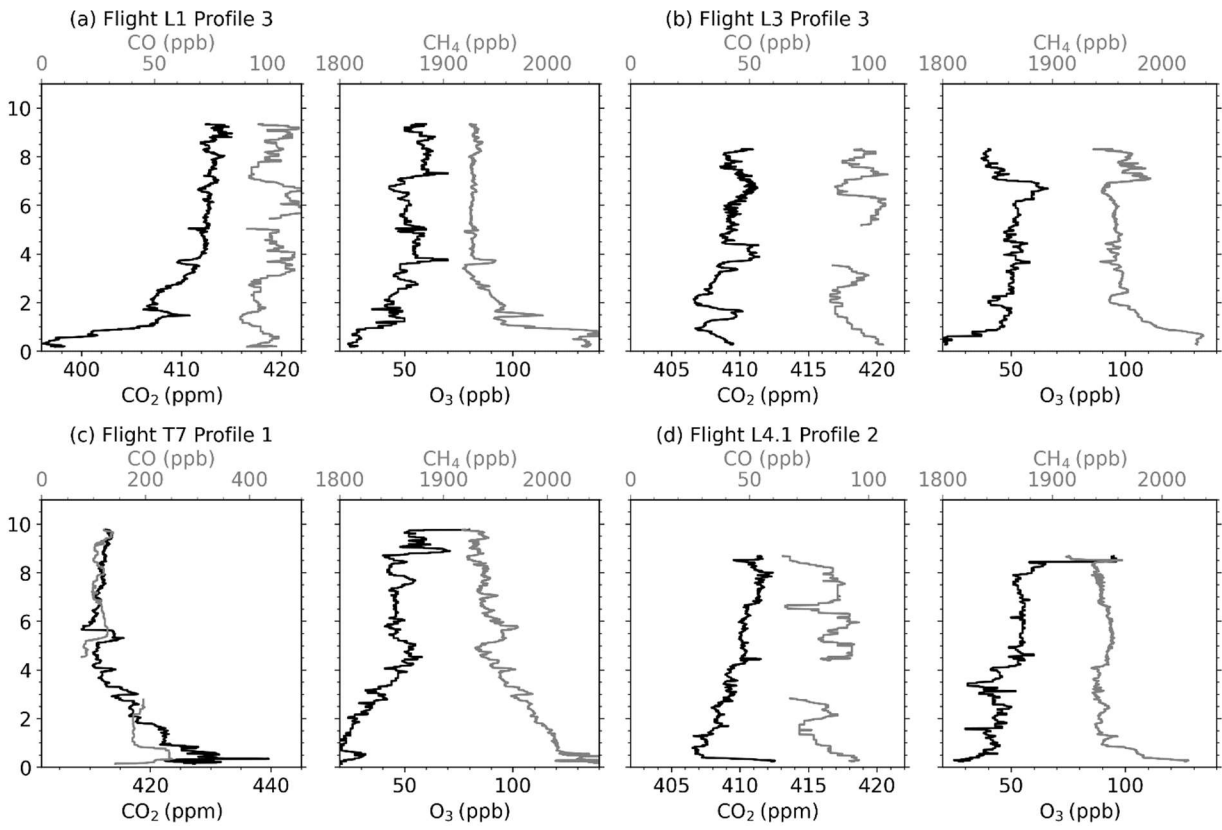
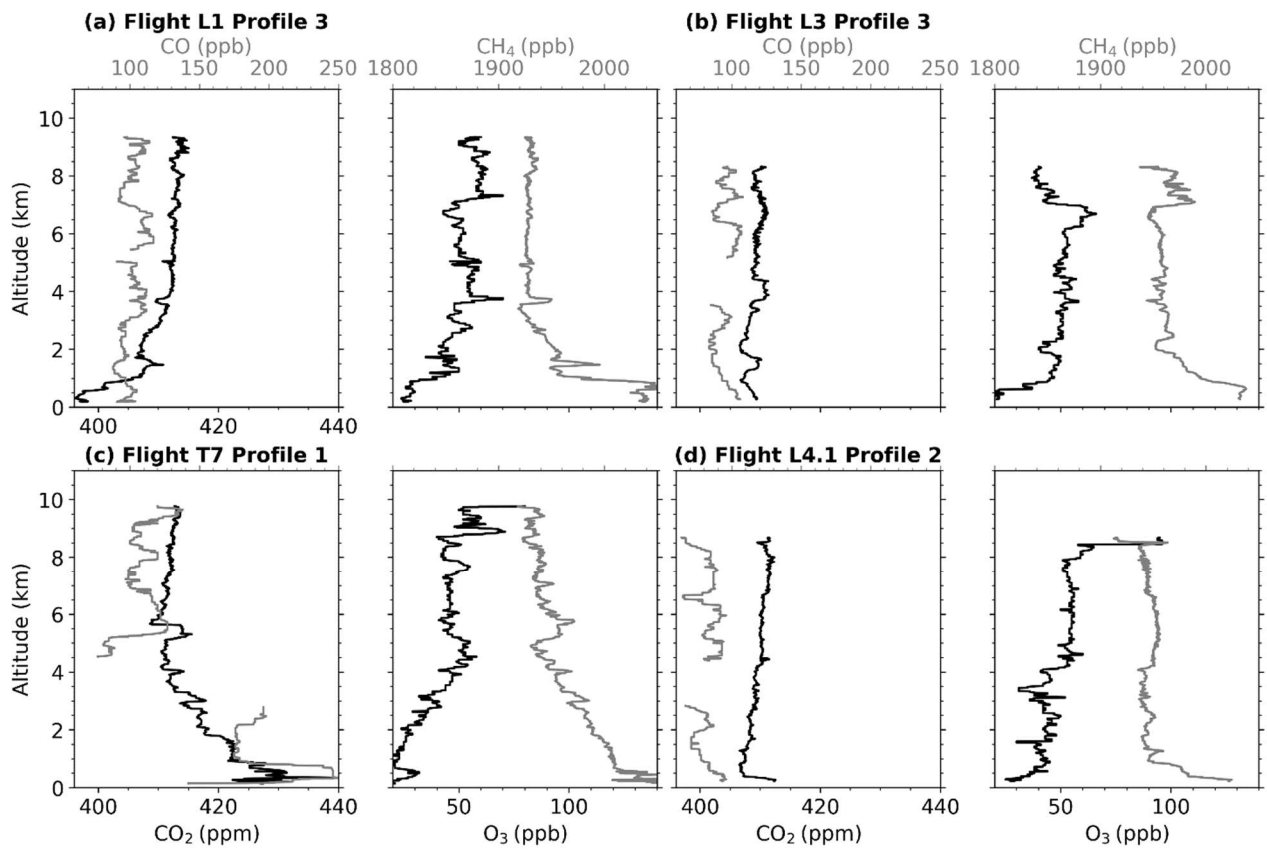


Figure 6: Four selected vertical profiles of the campaign. CO₂ and CO are respectively in black and gray on each left plot. O₃ and CH₄ are respectively in black and gray on each right plot. ~~Please note the different scales for CO₂ and CO.~~

405

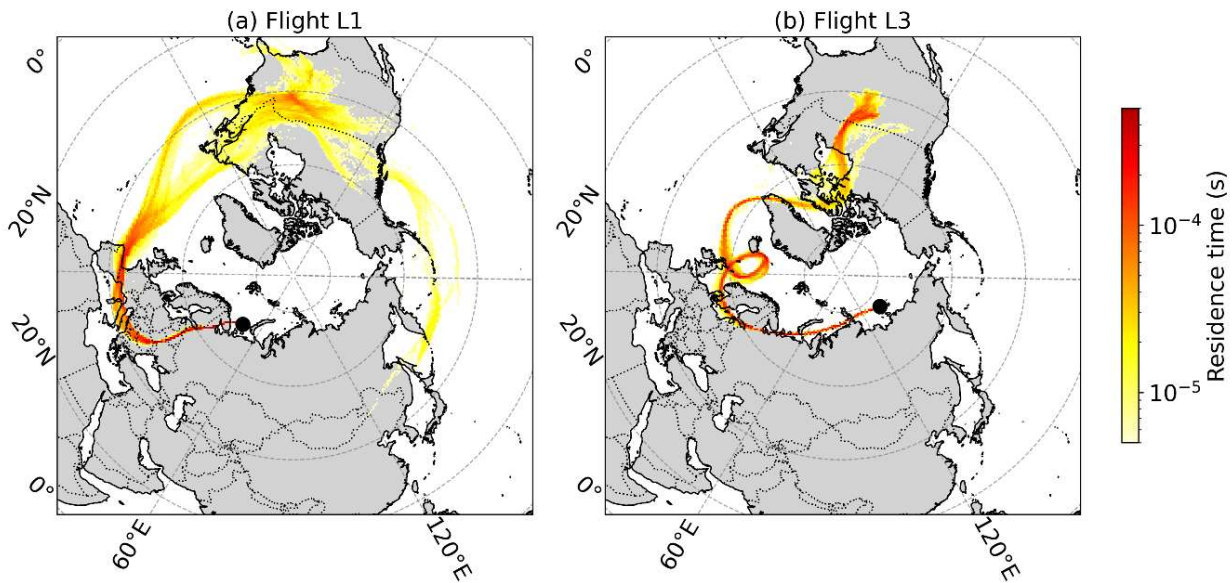



Figure 7: Total column footprints for simulated retro-transport of particles released at 14:53 UTC at an altitude of 9324 m during flight L1 (a) and at 05:01 UTC at an altitude of 5256 m during flight L3 (b). The black dot represents the receptor position.

410

Figure 6b shows the second ascent of L3, in the North of Siberia. Here, CO_2 ~~mole fraction~~mixing ratio shows less variability than flight L1 profile 3. A slight decrease of -3 ppm of CO_2 and -20 ppb of CO with altitude can be observed under 1000 m, possibly indicating the influence of emissions by local combustion, although this ascent does not start at an airport. We can also observe a stratification in the CO_2 and CO profiles with mixing ratio changes in stacked layers whose thickness ~~varies~~vary

415   between 500 m and 1500 m. This stratification has already been observed in previous campaigns (Paris et al. 2008) and is due to slow stirring in the troposphere under reduced vertical mixing. This profile is also characterized by significant sources of CH_4 in the BL as it is suggested by the strong enhancement of 100 ppb close to the ground (Fig. 6b). Among potential sources is methane hydrates degradation from the East Siberian Arctic Shelf (Berchet et al., 2016; Thornton et al., 2016; Berchet et al., 2020). The influence of potential sources will be discussed in Sect 3.4. In the same profile, O_3 is anti-correlated with CH_4

420   above 3000 m, except at the top altitude around 8500 m where O_3 ~~mole fractions~~mixing ratios unexpectedly decrease closer to the stratosphere. This O_3 depletion mentioned in Sect 3.2. (-24 ppb compared to immediately lower layers) occurs in the air mass whose history is shown in Fig 7b. The same backward transport simulation with a threshold altitude of 500 m (appendix Fig. C1) shows an air mass coming from the BL in Northern America ~~and~~then crossing the Atlantic Ocean and Northern Europe~~.~~ (only visible on the total column footprint). This layer with poor ozone ~~concentration~~mixing ratio is likely to be an

425   uplift of BL air with the origin of the depletion potentially coming from another location further upstream. The O_3 ~~mole fraction~~mixing ratio at top altitude varies from 37 to 40 ppb which is comparable to local ~~mole fraction~~mixing ratio at 2000 m (40 to 42 ppb) of the same flight, or flight L1 (41 to 42 ppb just under 2000 m) corresponding to regional BL air.

3.3.2 Yakutsk urban area

Another singular event formerly mentioned was a possible pollution enhancement in the area of Yakutsk. The corresponding
430 vertical profile in Fig. 6c shows high CO_2, CH_4 and CO mixing ratios (respectively 439.60 ppm, 2070 ppb and 251 ppb) as the aircraft took off from Yakutsk. At low altitude, both CO and CH_4 present high correlation with CO_2 (respectively $R^2 = 0.92$ and $R^2 = 0.84$) as shown in Fig. 8a and Fig. 8b while there is no correlation at higher altitude. Such suggests combustion emissions possibly related to fossil ~~fuels~~fuel exploitation and/or use. This region is indeed characterized by the presence of operating O&G infrastructures and coal mines (Fig. 1). We can also observe the lowest O_3 ~~mole fraction~~mixing ratio of the
435 campaign at 2 ppb at the ground level of Yakutsk (value too low to be seen on the figure). Having this O_3 near-ground minima over landing air-strips in polluted city plumes of Yakustk suggests a titration of O_3 by NO.

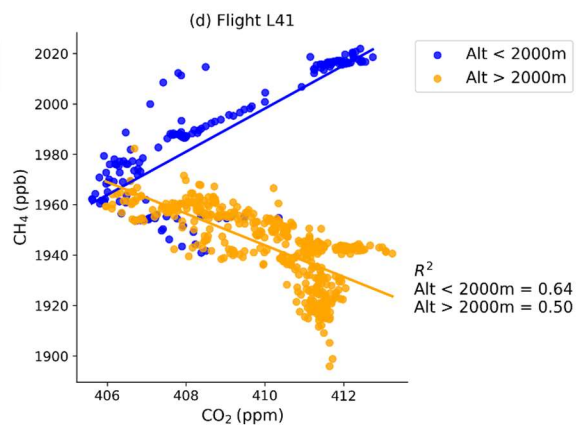
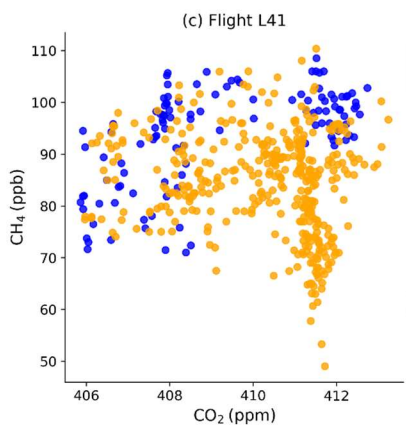
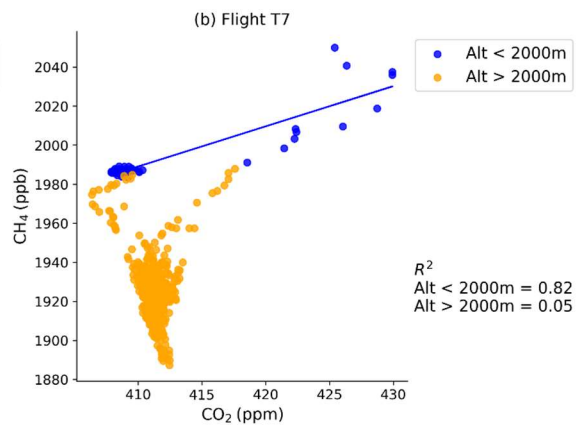
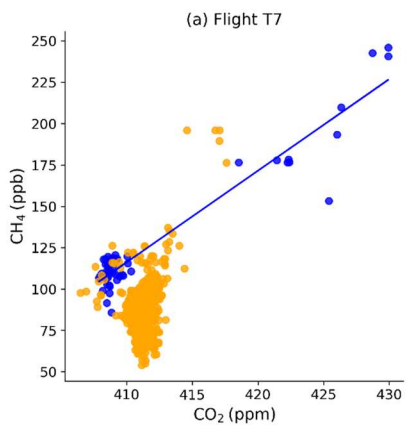
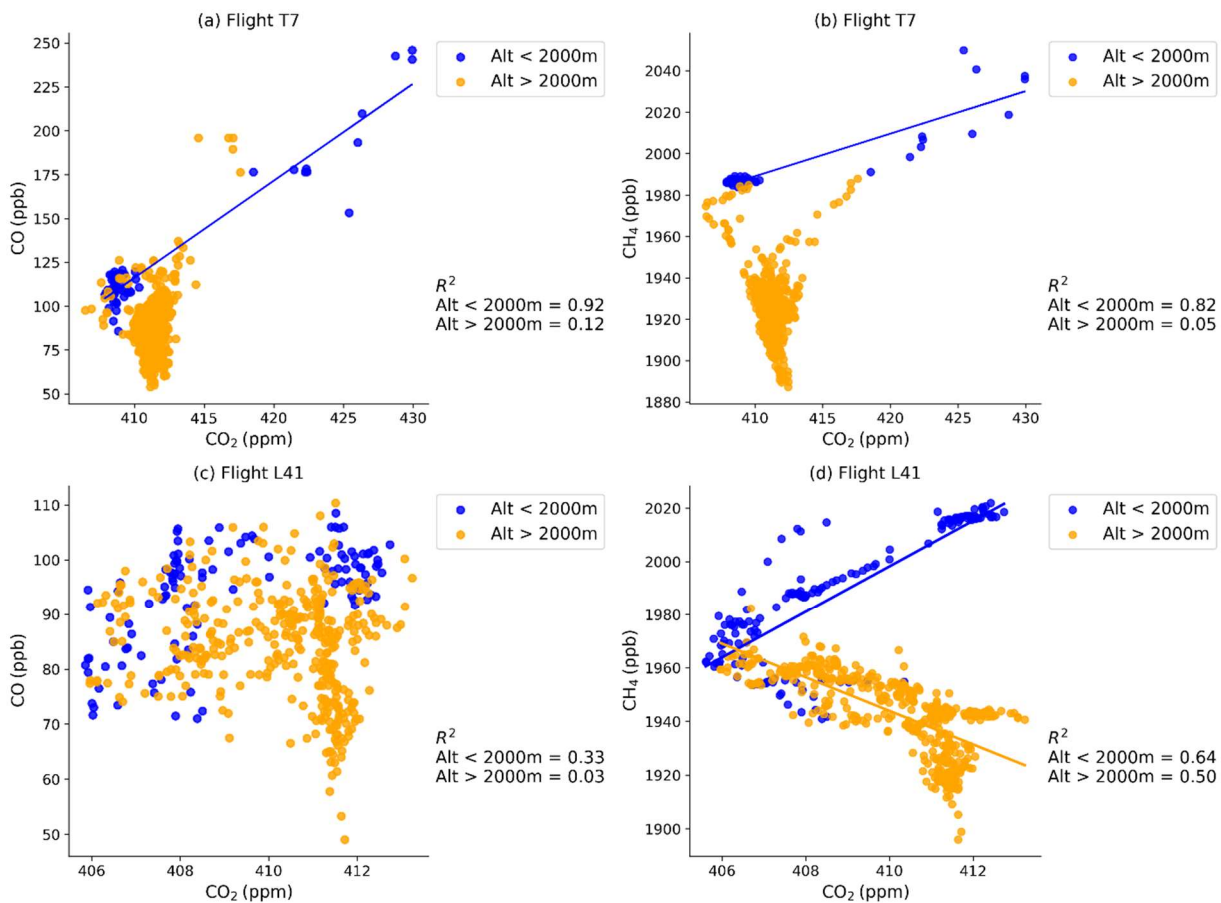



Figure 8: Scatter plots of CO₂ vs CO (a) and CO₂ vs CH₄ (b) for flight T7. Same for flight L4.1 in (c) and (d). Blue dots and respective regression lines are for data acquired under 2000 m. Orange dots and respective regression lines are for data acquired above 2000 m. Correlation coefficient R^2 are a displayed for each set.

3.3.3 CO₂ enhancement at low altitude in the Far East

~~Althgouh~~Although the average CO_2~~2~~ profiles of Flights L4.1, L4.2 and L4.3 show little variability over the Bering Strait ~~measured~~ we find significant CO₂ enhancements at very low altitude (<500m; Sect. 3.2). Here we focus our analysis on the loop L4.1 as the three flights have similar features. In the vertical profile of the first descent (second profile), remote from any airport, a high CO₂ mixing ratio of 412.5 ppm has been observed close to the ~~ground~~ surface (200m), against 407 ppm at 500 m altitude (Fig. 6d). The CH₄ mixing ratio also decreases from 2025 ppb to 1980 ppb between 200 m and 500 m altitude. On the other hand, CO presents only a small, opposite gradient of about -10 ppb across the same altitude range. Above 500m, the CO_2~~2~~ profile exhibits a more typical shape of marginally increasing vertical gradient.

Figure 8c shows a scatter plot of CO against CO_2 for the whole flight L4.1 (including all six profiles). The correlation coefficient under 2000 m is significant but slightly ~~low~~lower (R^2 = 0.33, ~~*p* = xxxx~~p = 10^{-11}) tha~~n~~t correlation coefficients between CO_2 and CH_4 shown on Fig. 8d (R^2 = 0.64, p = 10^{-37}~~*p* = xxxx~~)~~but still significant,~~ suggesting that the CO_2 enhancement may be ~~not be strongly~~only driven by a mixture of emissions ~~from~~ including local combustion processes. ~~Fig. 8d presents good correlation between CH_4 and CO_2, both at low and high altitude (respective R^2 = 0.64 and R^2 = 0.50).~~ The regression slope below 2000 m[JP7] is 8.5 ppb of CH_4 ~~/~~per ppm of CO_2. Non-combustion sources emitting both CO_2 and CH_4 leading to such ~~as~~ signature could include one or several fossil fuel exploitation areas co-emitting the two species. According to the Global Fossil Infrastructure Tracker map from Global Monitor Energy, no operating O&G ~~pipelines~~pipeline or terminal is reported in the East Siberian region, and the closest one being Kenai Alaska liquid natural gas terminal (Fig. 1). Still according to Global Energy Monitor, only Amaam North Coal Mine is active in East Siberian region (Fig. 1) but there are other coal mine activities reported in Russian Far East (Petäjä et al., 2021). Section 3.4.1~~xxx~~ below shows that influence on this part of the flight L4.1 are mostly of natural origin and linked to the potential emission sensitivity of~~to~~ Alaskan emissions.

Multiple regional and diffuse sources may be at play in explaining the enhancement observed. Another possibility is an emission from marine CH_4 sources. The absence of sea ice at the time of the campaign in the involved area (Fig. ~~B1~~A1) (Fetterer et al., 2017) is enabling air-sea exchange, notably for CH_4.

~~-~~The anti-correlation between CO_2 and CH_4 above 2000 m (Fig. 8d) could be a residual from active sources for CH_4 and active sinks for CO_2 during the summer. While the air in this region has been relatively well-mixed as shown by the flat vertical profiles of CO_2 and CH_4 for flights L4.1, L4.2 and L4.3 ~~on~~(-Fig. 5) and the reduced value ranges for CO_2 and CH_4 on the scatter plot of Fig. 8d (compared to the value ranges on Fig. 8b), ~~there are some imperfections~~mixing of airmasses affected simultaneously by CO_2~~2~~ sinks and CH_4~~4~~ sources in the past might be incomplete resulting in this residual anti-correlation. This will be better investigated in the Sect. 3.4.1 by using the simulated mixing ratios for flight L4.1.

3.4 Untangling the different methane sources with a Lagrangian model[CN8]

3.4.1 Contributions to measured CH_4 ~~depending on the geographical~~ by source type and by location

This section aims at identifying the main emissions influencing the CH_4 ~~concentrations~~mixing ratios through case studies of selected flights. Fig. 9 shows a scatter plot of measured and simulated CH_4 enhancements for each flight ~~(enhancement being defined~~. Enhancement here ~~as~~refers to the measured mixing ratio minus the ~~lowest mixing ratio~~background of the flight~~)~~. (background is defined in Sect. 2.3). Correlation coefficient R^2 range between 0.~~26~~28 (for flight ~~T6~~T5) and 0.~~87~~86 (for flight L1), with associated p-values that are inferior to 10^{-10} for every flight. For most flights, the agreements between simulations and measurements are satisfactory and enable comparison of simulated source contributions to observed CH_4 enhancements. However, ~~we will show also that there is~~the 1:1 dashed line highlights an underestimate of simulated ~~mole fractions compared to the measurements~~mixing ratios that will be discussed in this section and the next one. For the study, we select one flight in

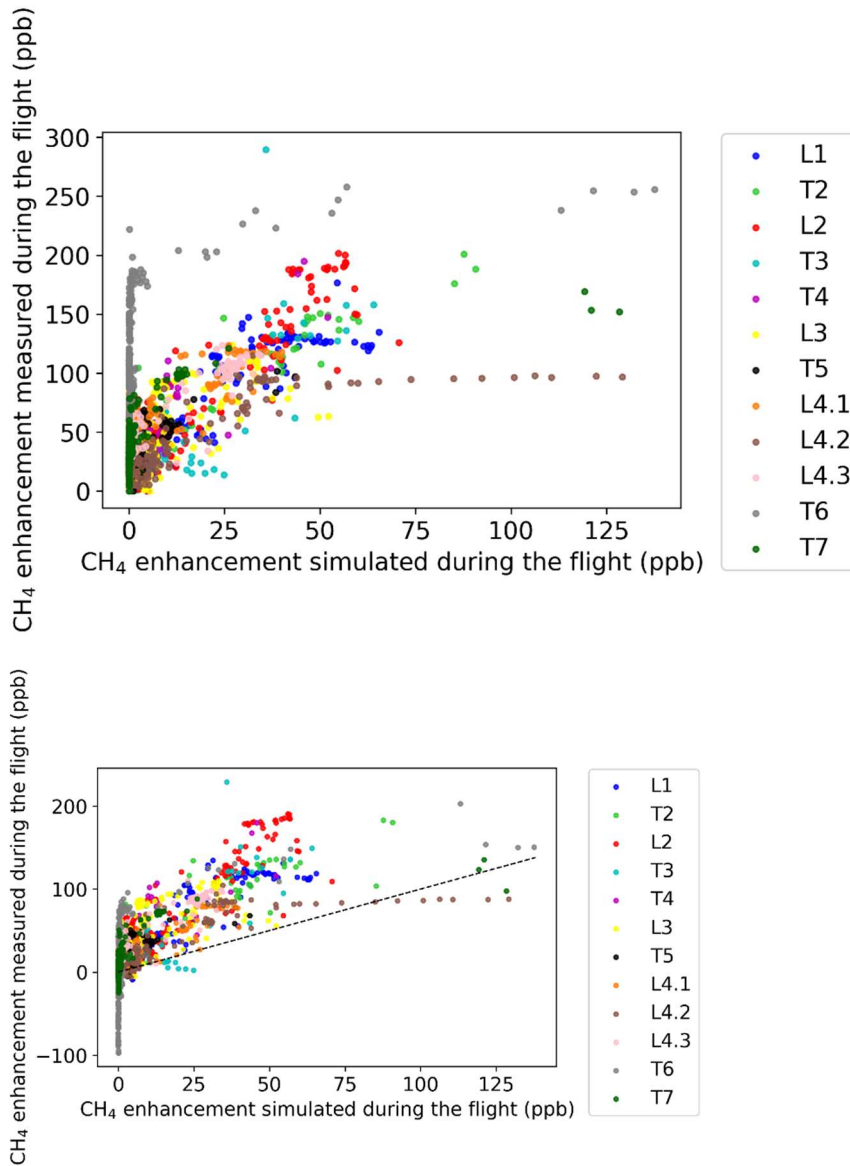
western Siberia: T2 that has a correlation coefficient $R^2 = 0.\text{85}\underline{82}$, and one flight in eastern Siberia: L4.1 that has a correlation coefficient $R^2 = 0.67$.

485

Figure 9: Scatter plots of simulated CH₄ enhancement vs measured CH₄ for each flight of the campaign except T1~~.~~ with the 1:1 black dashed line. Data were aggregated into 1 min bins to produce this figure.

Figure 10 shows the simulated CH₄ ~~mole fractions~~mixing ratios aggregating contributions from tagged sources during flight

490 T2. The simulated signal at the receptor position is compared to measured CH₄ enhancement. The CH₄ enhancement variability

is reasonably well reproduced by the model (Fig. 10a) but there is a consistent underestimation in total enhancement values. ~~Some simulated contributions are underestimated by a factor 3 to 4.~~ The possible reasons of this underestimate will be discussed later in Sect. 3.4.2. However, the ability of the model to reproduce the peaks and trough of CH$_4$ during the entire campaign allows some confidence in the comparative tagged tracer analysis intended here.

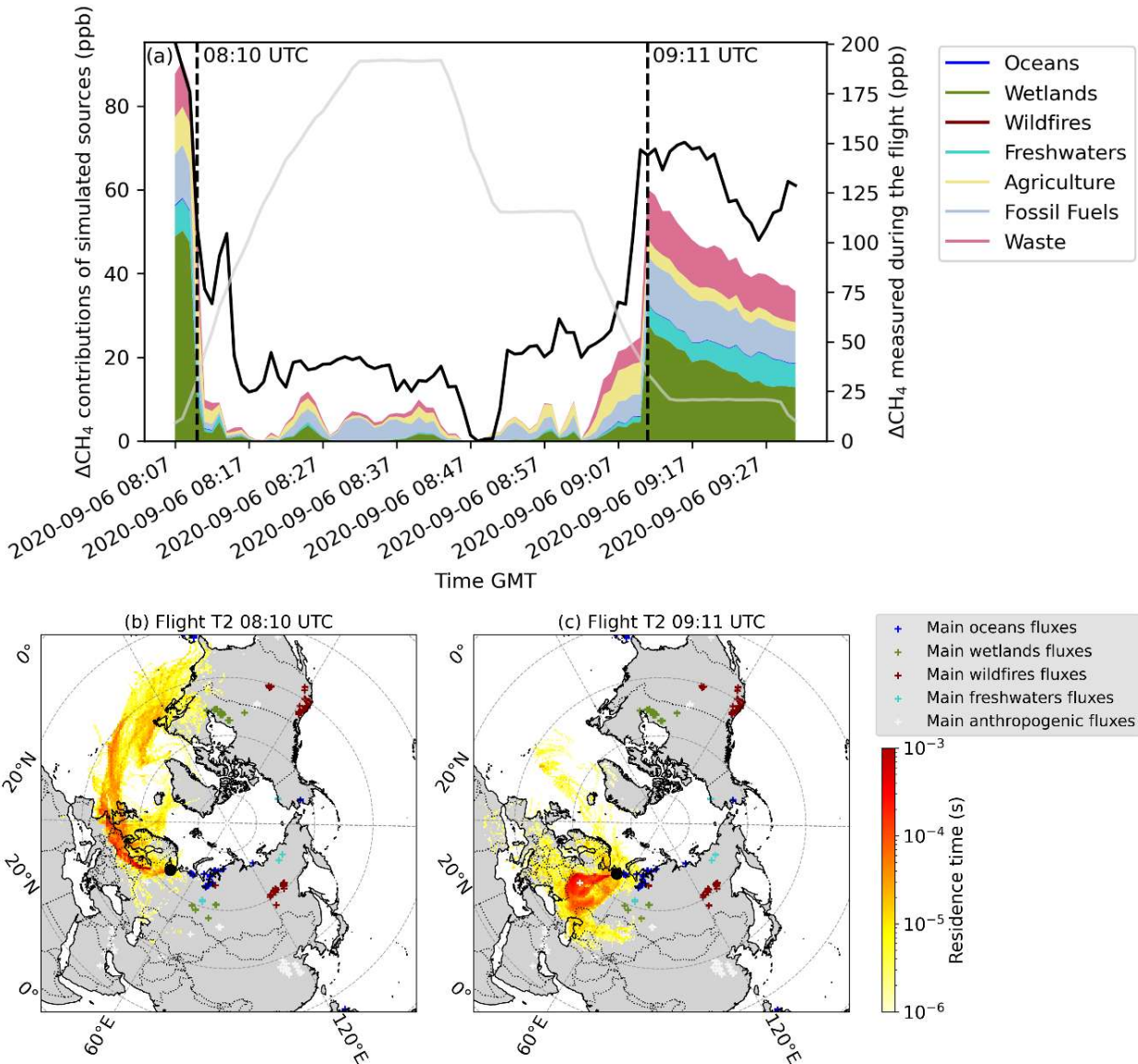
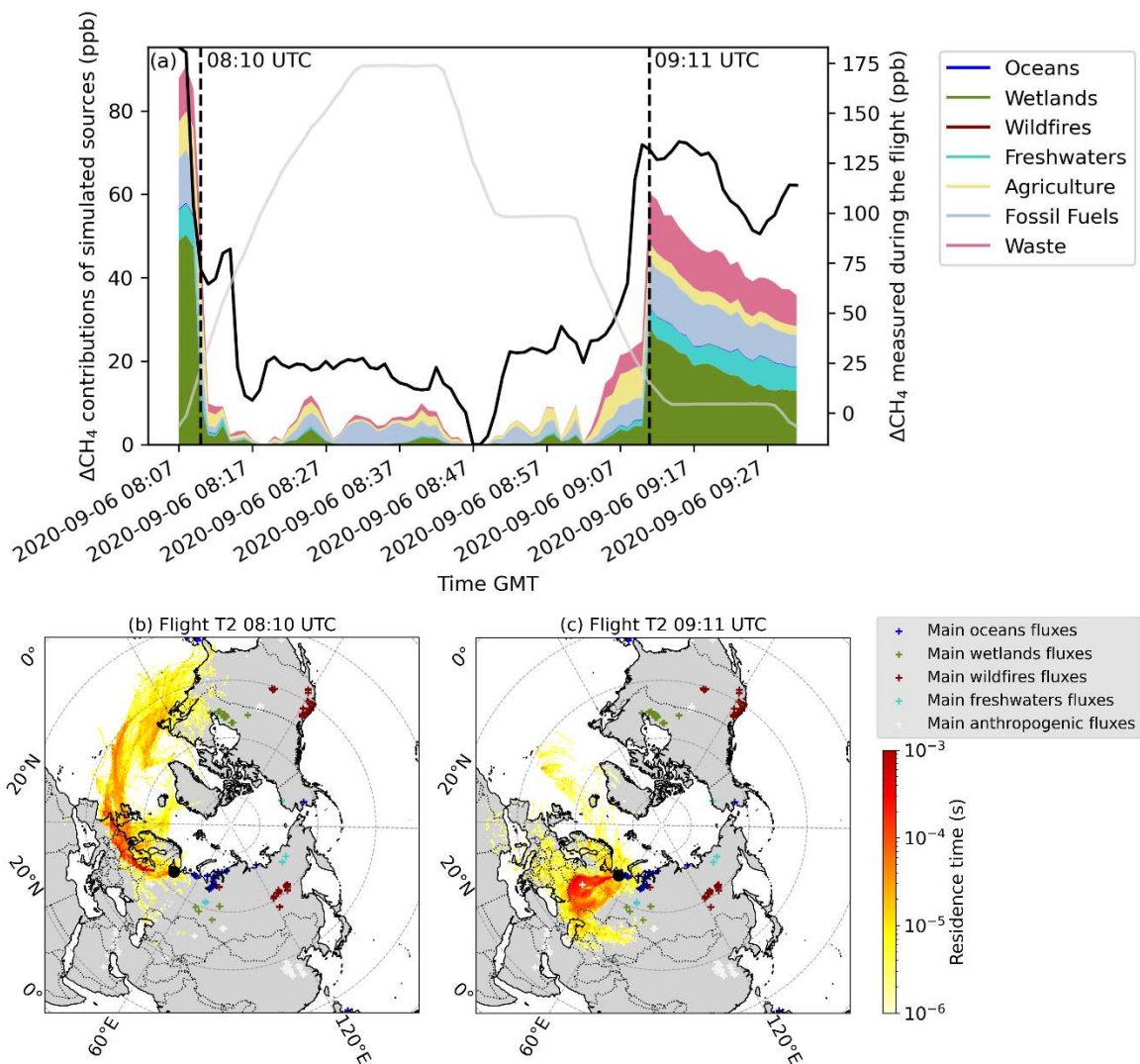



Figure 10: (a) Simulated CH₄ enhancement (coloured stacked plot) and measured CH₄ (black line) for flight T2. Note the different Y-axes. Altitude is shown in grey. (bottom at 12 m and top at 8963 m). The two vertical dotted lines indicate the measurements represented in the following footprints. (b) 10 d potential emission sensitivity (PES) for particles released at 08:10 UTC. (c) Same as (b) at 09:11 UTC. The colored "+" symbols represent the fluxes with biggest intensities derived from each inventory. The black dot represents the receptor position.

Figure 10b and c show PES for two selected positions representative for the flight. The footprint in Fig. 10b corresponds to the beginning of the ascent after leaving the airport (first dotted line at 08:10 UTC in panel a). At this position, an enhancement of 50 ppb was simulated. This enhancement is simulated to originate from a dominant anthropogenic contribution distributed between fossil fuel ~~exploitation~~emissions, agriculture and waste management for a total of 36 ppb (the following highest

contribution being wetlands with 10 ppb of contribution). The footprint Fig. 10b shows that the air mass has mostly resided over Western Russia, Central Europe, the Atlantic and, to a lesser extent USA (although this corresponds to an air mass age close to the limit of particle tracking of 10 days). This is consistent with the dominance of agricultural CH_4 fluxes in

510 anthropogenic fluxes in Europe and the USA (appendix Fig. A1bB1b) while agriculture is much less significant in the Russian emission inventories (Crippa et al., 2019). This documents long-range transport of free tropospheric air masses with relatively high CH_4 (comparable to values encountered in the BL) in western Siberia.

The footprint on Fig. 10c, close to the Arctic Ocean at lower altitude (second dotted line at 09:11 UTC in panel a), corresponds to mixed enhancements of 28 ppb by wetlands and 27 ppb by human activities. The PES is concentrated on western Russia,

515 with a major sensitivity to anthropogenic flux at the core representing the city of Moscow (white-filled "plus" sign on Fig. 10c). The simulated enhancements are driven by with the CH_4 fluxes from fossil fuelsfuel exploitation and use (appendix Fig. A1aB1a) in western Russia, and from wetlandswetland fluxes in Western Siberia (appendix Fig. A1cB1c), especially in the Vasyugan swamp.

At the end of the flight, when getting closer to Naryan Mar, Fig. 10a depicts an enhancement of 7 ppb associated with

520 freshwater [JP9][CN10]sources that is represented by the constant light blue area between 09:11 UTC and the last data at 09:31 UTC. A large number of lakes are present around Naryan Mar and its surroundings (Fig. C1D1), supporting the importance of freshwater contribution in the atmospheric CH_4 burden. Also, it is likely that a part of the simulated enhancements due to wetlands may be associated to freshwatersfreshwater CH_4 emissions because there may be some overlap in inventories regarding small lakes and wetlandswetland sources, especially in complex regions with presence of many wetlands, lakes or

525 rivers like in Naryan Mar outskirts.

Focusing now on Eastern Siberia, Fig. 11 illustrates the simulated and measured CH_4 enhancements, and two PES footprints associated with two specific positions of the flight. Overall, measured and simulated enhancements are lower than for the Western Siberian flight previously discussed. The first case (00:25 UTC during flight L4.1, first dotted line) is a very thin plume of elevated CH4 in the lower free troposphere. It has a simulated enhancement of 14 ppb dominated by contributions

530 due to fossil fuel exploitationemissions (7 ppb), agriculture (4 ppb) and waste management (3 ppb). The PES footprint in Fig. 11b shows that the air masses have partially resided over north-east China, very close to fossil fuels exploitationfuel emission sources (see the white "+" sign on Fig. 11b) and where important CH_4 fluxes from agriculture are present (Fig. A1B1). Several fossil-fuel extraction infrastructures are located in this region (Global Energy Monitor, 2022). A large part of the air masses has also resided over East China Sea and Bering Strait where there are CH_4 fluxes due to offshore fossil fuel according to

535 EDGAR inventories (appendix Fig. A1B1) explaining the domination of fossil fuel exploitationemission contribution in the selected peak. Overall at altitudes above 2000 m, most contributions are missed by the model showing that the air in the free troposphere has not been into contact with the surface for more than 10 days. This supports the anti-correlation observed in the Sect. 3.3.3 between CO_2 and CH_4 for flight 3.1 that is a residual of the active sources and sinks during the past summer.

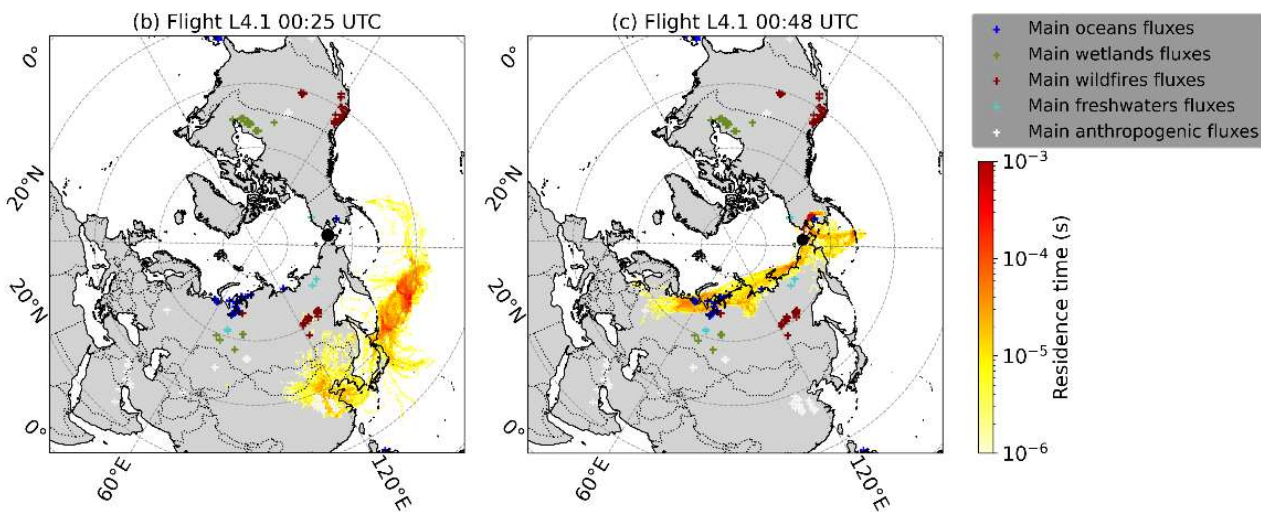
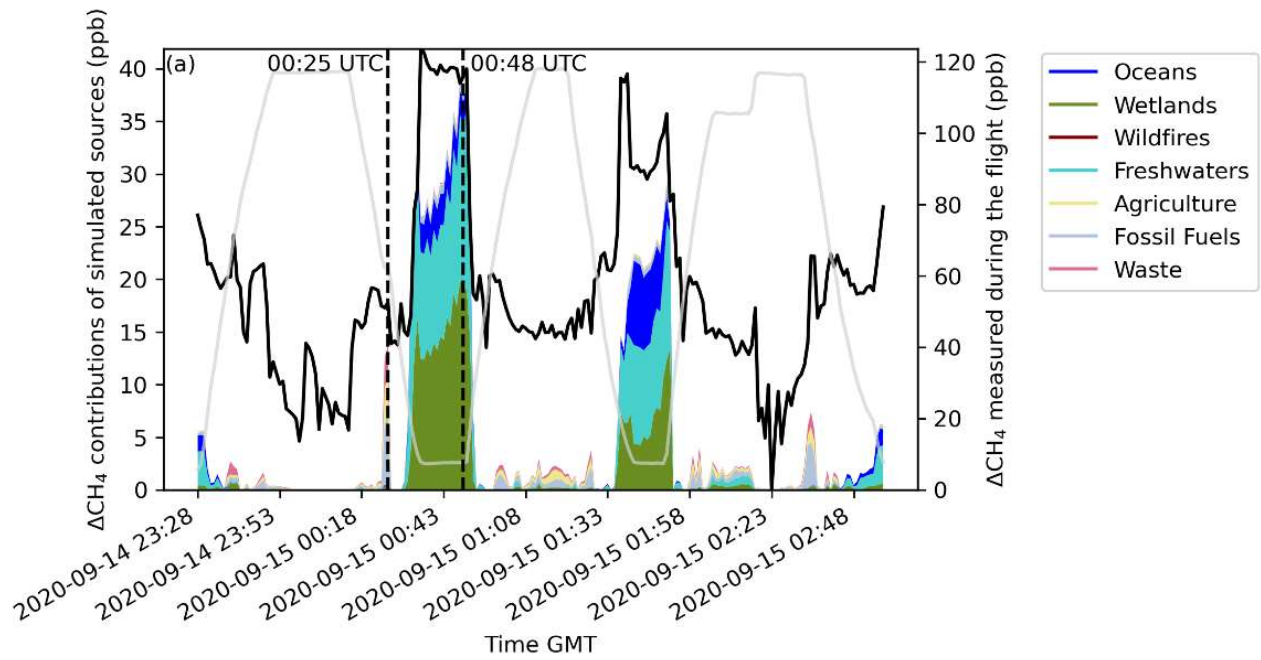
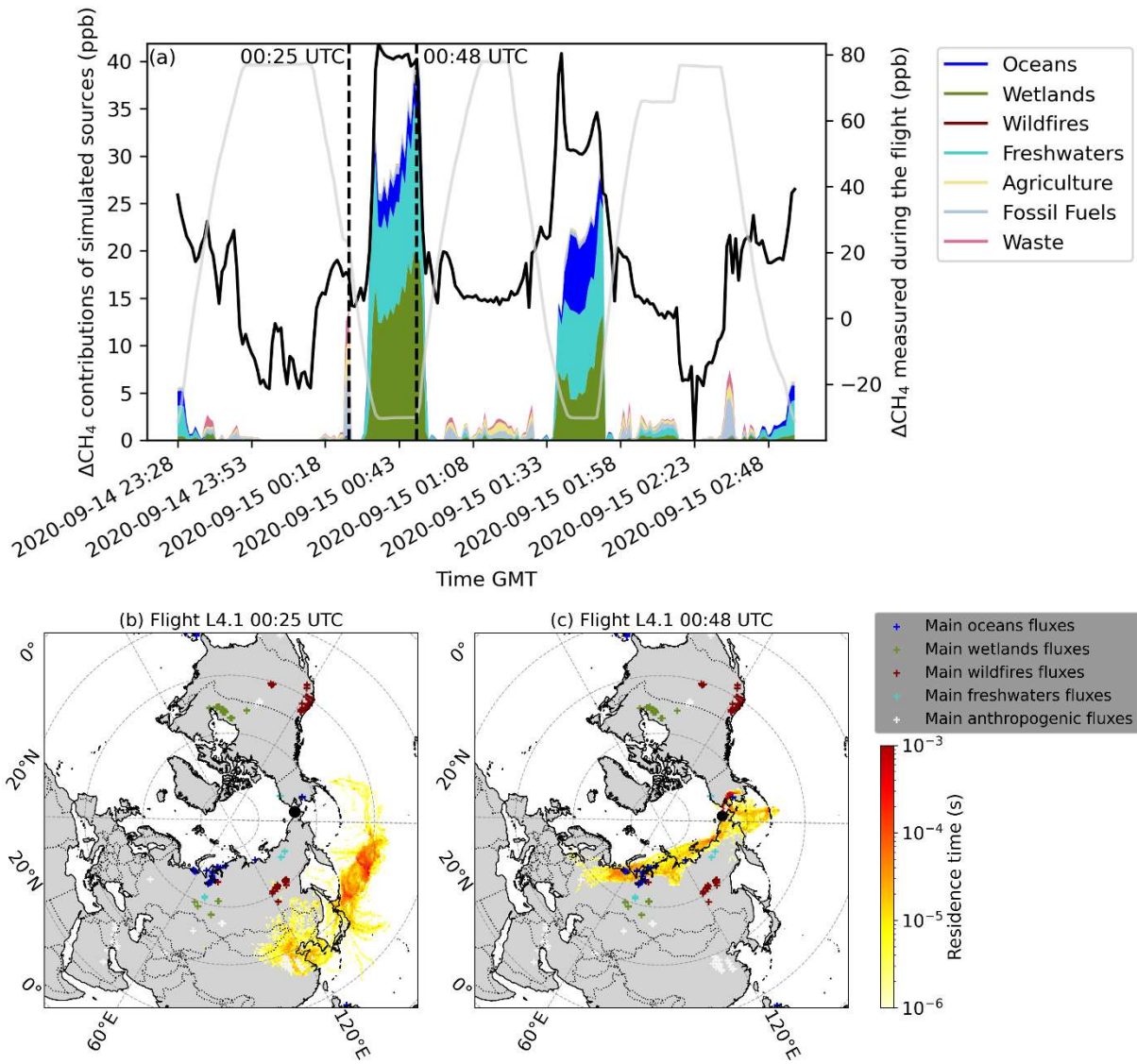



Figure 11: (a) Simulated CH₄ enhancement (coloured stacked plot) and measured CH₄ (black line) for flight L4.2. Note the different Y-axes. Altitude is shown in grey. (bottom at 38 m and top at 8707 m). The two vertical dotted lines indicate the measurements represented in the following footprints. (b) 10 d potential emission sensitivity (PES) for particles released at 00:25 UTC. (c) Same as (b) at 00:48 UTC. The colored "+" symbols represent the fluxes with biggest intensities derived from each inventory. The black dot

 represents the receptor position.

The second dotted line at 00:48 UTC in Fig. 11a is associated with simulated CH₄ enhancement of 38 ppb [JP11]close to the BL. The simulation indicates that air masses travelled mostly over the Bering Strait, East Siberian Sea and Laptev Sea (Fig.

11c). The enhancement is dominated (in relative terms) by the freshwater contribution with 15 ppb. Wetlands still play a large role with simulated mixing ratios that can represent 20% to 50% of the peaks at low altitude for this flight. Methane from the Arctic Ocean significantly appears in the simulated low-altitude enhancement but is lower than the other natural sources, with a maximum at 8 ppb in the same flight (Fig. 11a).

Over these two flights, we can observe that the peak‑‑to‑‑peak amplitude of simulated mixing ratios is ~~larger~~higher than the ~~peak-to-peak~~ amplitude of measured mixing ratios ~~(e.g. looking at the decrease just after the first peak of Fig. 10a[JP12]),~~ indicating that some contributions are likely missing in our model~~.~~ (e.g. the second simulated peak of Fig. 10a, between 08:07 UTC and 08:17 UTC is missing some contributions). Some simulations though have smaller occurrences of underestimations of observed peaks, such as flight T3. Wildfires have influenced previous campaigns during specific episodes, especially close to the sources, as reported in Paris et al. (2008) and in Antokhin et al. (2018). Here ~~we notice~~ the ~~absence of~~wildfires are simulated ~~wildfires contribution throughout~~as well, but are not visible since the contributions are negligible compared to the other ones over the whole campaign.

The contributions of the oceanic sources are not dominant in the simulation over any freshwater source in this study. Here, freshwaters appear as a dominant term in the CH$_4$ emitter influencing variability over Northern Siberia. Given the modelling uncertainties, it is challenging to attribute missing simulated CH4 to a minor source term. Hence, comparing our simulated and measured CH$_4$ time series do not provide sufficient confidence for assuming that the Weber et al. (2019) marine inventory is underestimated. The two PES footprints reveal a local CH$_4$ production in this part of Siberia, even in polluted air masses at high altitude whereas in the West they were resulting from long-range transport. In addition, there might be a sampling bias due to the campaign observation strategy. CH$_4$ measured from loops flights more often originates from natural sources compared to in transit flights, which sample BL air essentially in the vicinity of airports. The aircraft flies more frequently at low altitude and over remote areas during the loop flights.

Overall, our model-data comparison indicated that CH$_4$ variability in Western Siberia is dominated by a combination of Western Siberia wetland sources and human activities largely related to fossil fuel emissions while eastern Siberia is characterized by strong natural sources such as wetlands, lakes, ponds and, to a smaller extent, oceans.

3.4.2 Sources of uncertainties and the underestimation of measurements

Although peaks and ~~trough~~troughs are well simulated, total CH$_4$ ~~mole fractions~~mixing ratios across all flights were consistently underestimated. While FLEXPART is performing well at high altitudes in representing the signal variability, it has some limitations in simulating local pollution in the boundary layer. Stohl et al. (1995) reported that the representation of vertical transport leads to greater errors than lateral transport. This might be due either to the low resolution of meteorological data, poor emission inventory accuracy in Siberia, or a simplification of flux densities under the planetary boundary layer (Stohl et al., 1995). In addition, the spatial resolution of 1° of the meteorological input data deteriorates the resolution of simulations under the planetary boundary layer.

Vertical transport across the boundary layer may have a significant role in the model underestimation given the trajectory of our aircraft campaign with large changes in altitude. Figure 12 shows the simulated CH_4 and measured enhancement for the period of the campaign at Barrow, Alaska (Dlugokencky et al., 2021c), which is obviously not subject to vertical movement of the receptor. Simulated mixing ratios are much less biased [JP13][JP14]compared to the measurements (simulated mean CH_4 being at 52 ppb and Barrow CH_4 mean at 37 ppb). There are some occasional overestimates that might be due to fluxes included in both ~~freshwaters~~freshwater and ~~wetlands~~wetland inventories. At this time of the year, Barrow is dominated by freshwater emissions from Alaska and Canada whereas anthropogenic emissions are largely absent in the simulation.

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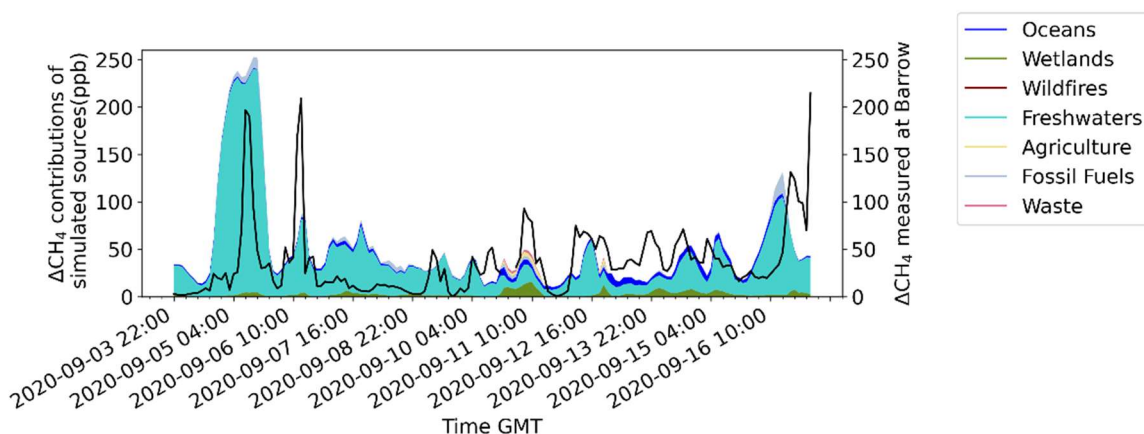
Figure 12: Comparison of enhancement of measured CH_4 (black line) and simulated CH_4 (coloured stacked plot) at Barrow station during the whole campaign.

595 For some inventories, the specific month of September 2020 was not available in the data. Previous years were used instead. This could lead to some underestimates as we can expect that emissions from some sources may possibly increase with time (e.g. CH_4 from anthropogenic activities) or subject to high temporal variability (CH_4 from natural biogenic sources). Two inventories, EDGAR (anthropogenic fluxes) and ORCHIDEE (fluxes from wetlands), were available for different years. We produced a sensitivity analysis by taking the previous years of these two inventories and producing the simulated ~~mole fractions~~mixing ratios for the same period. The time of reference is set to the month of September for the last year available of each inventory (2018 for EDGAR and 2017 for ORCHIDEE) and the ~~mole fractions~~mixing ratios of the two respective years before were also simulated. The simulations ~~exhibit~~exhibited similar patterns, and we extracted the simulated values of the four peaks studied in Sect. 3.4.1. Enhancement's variations induced by changing the year of EDGAR inventories don't exceed 0.4 ppb. Enhancement's variations induced by changing the year of ORDCHIDEE inventories can go up to 5 ppb due

to the variability of ~~wetlands~~wetland emissions. This parameter ultimately appears to have little influence in our simulations that only cover a short period.

Another source of error could be the underestimation of fluxes from poorly-known natural sources such as the marine ones. The estimation of CH_4 emissions by the ocean largely depends on the approach as reported in Weber et al. (2019). However, values vary significantly among the few available studies. For example, methane emissions in East Siberian Arctic Shelf are still uncertain and are estimated to range between 8 and 17 Tg CH_4 yr^{-1} (Berchet et al., 2016) by oceanographic approach, while an atmospheric inverse estimation provides a flux estimate not higher than 4.5 Tg CH_4 yr^{-1} (Berchet et al., 2016). As discussed previously, they are influencing CH_4 variability but do not appear to be a dominant term in any part of our flights. Freshwater estimations are also subject to significant variability as it is difficult to be exhaustive in the distribution of the sources and there are ~~fundamentals~~fundamental gaps in the lakes model (Matthews et al., 2020). The "snapshot" effect of methane flux measurements in field campaigns may also ~~leads~~lead to underestimates (Wik et al., 2016) or biases. Further works may focus on performing simulations with other inventories such as the one used in Matthews et al. (2020) for high-latitude lakes or the submarine seep estimates from Etiope et al. (2019). On the other hand, it is reported in Matthews et al. (2020) that small wetlands are often interwoven with lakes causing difficulties to distinguish them which leads to overestimates. ORCHIDEE also may overestimate CH_4 net primary production by wetlands due to a lack of wetland specific plant functional types representation (Ringeval et al., 2012; Wania et al., 2013). It could be relevant to test other flux estimated on wetlands and quantify the associated variation to check possible overestimates. The present study only captures a precise moment of the atmospheric condition in Siberia, giving an overview of main phenomena at the end of the summer in the year 2020. Therefore, observations should be completed with medium-term and long-term studies, and the aircraft data could be used as validation data for inverse modelling studies.

4 Conclusion

We have investigated the latest data of September 2020 aircraft campaign over the Russia's Arctic, including Siberia. It comprised 47 vertical profiles split in 13 flights across all of Siberia giving opportunities to observe CO_2 and CH_4 emissions and transport in different locations of an imperfectly known region that has a serious impact in global carbon budget. In-situ measurements of CO and O_3 have also been performed and are used as complementary tracers in this work.

CO_2 mixing ratios (median value of the campaign at 410.83±3.29 ppm) were slightly lower than Mauna Loa average (411.52 ppm) due to the passage of air masses above CO_2 sinks, and CH_4 mixing ~~ratio~~ratios (median value of the campaign at 1939±45 ppb) were higher (Mauna Loa average at 1888 ppb), indicating an accumulation of methane from different sources. Both gas show a high variability over Siberia, especially in the lower and the upper troposphere. Western Siberia exhibits steep ~~mole fraction~~mixing ratio gradients in CO_2 explained by the presence of effective sinks close to the ground (dense vegetation with taiga) while higher altitudes are characterized by the long-range transport of air masses polluted by anthropogenic activities~~.~~, mainly related to fossil fuel emissions. Eastern Siberia is subject to local pollution and CH_4 emissions from natural sources

with less atmospheric transport. Individual vertical profiles also revealed more unique patterns such as the stratification of CO_2 and CO mixing ratios with altitude, the O_3 depletion at top altitude in air masses that crossed Norwegian Sea, and the excess of CO_2 in Bering Strait region. As we wanted to determine if ocean fluxes (and hydrate gas) had a significant role in methane emissions, we simulated the CH_4 enhancement by different types of sources ~~presents~~present in Siberia. It appeared that emissions are dominated by wetlands, fossil fuel ~~exploitation and~~ emissions, agriculture and waste management in the West, and by freshwaters and wetlands in the East. Aquatic sources may be underestimated but our measurements are not sufficient to confront existing inventories due to limitations in the numerical models and observational strategy. However, our data suggests that poorly estimated aquatic emissions at regional scale in Arctic Siberia deserve further research and more measurements. With this insight on the main methane sources in northern Russia at the end of summer, we advocate for further research on aquatic CH_4 sources in Siberia to better predict potential positive feedbacks between regional and global warming.

Data availability. In flight measurement data is currently available from the authors upon request (at jean-daniel.paris@lsce.ipsl.fr). Data will be available online with access through: https://yak.aeris-data.fr/.

Author contributions. CN performed data processing and simulations, and analysed the data. JDP designed the study. CN and JDP wrote the first version of the manuscript. AB, SW and MS helped to perform simulations. BB, MA, and SB designed, organised and together with DD, AF, and AK performed the aircraft campaign and did the trace gas measurements. All contributed to the manuscript.

Competing interests. None

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Appendix A: Additional material concerning CO$_2$ enhancements at low altitude in east Siberia

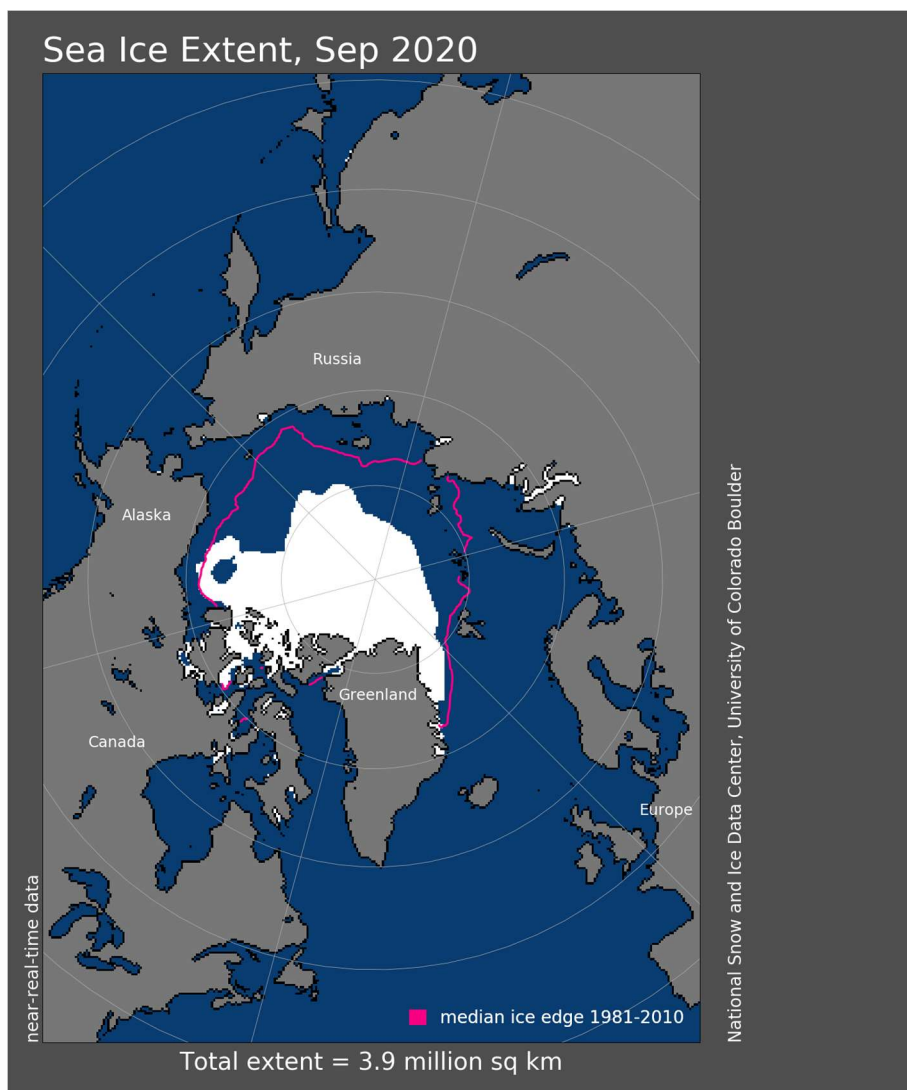



Figure A1: **Sea ice extent in September 2020. As illustrated, there is no sign of ice in Bering Strait (Fetterer et al., 2017).**

980 **Appendix B: Additional material on inventories**

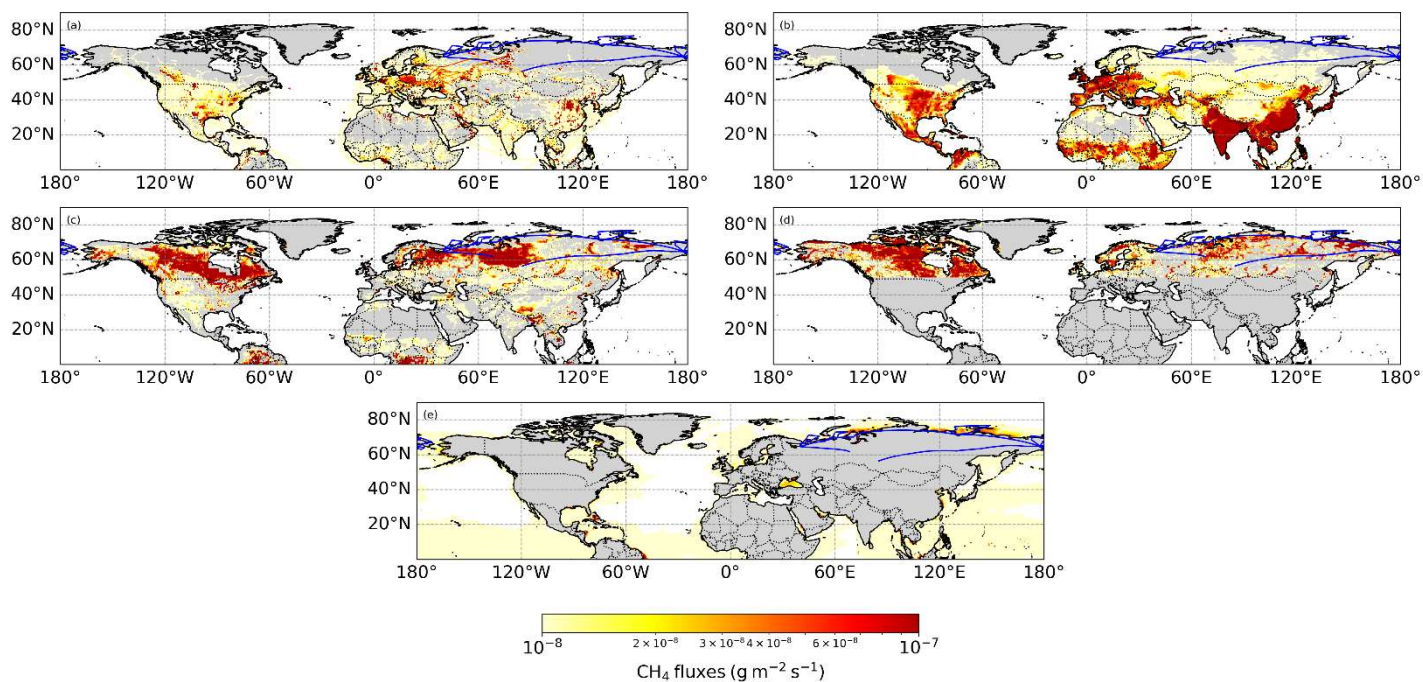
A1B1 EDGAR V6.0 Inventories used and sub-categories created

Table A1B1: The 21 EDGAR V6.0 inventories used for the present study regrouped in three different sub-categories.

Folder name	Name	Our Category
ENE	Power Industry	Fossil ~~fuels exploitation~~fuel emissions
REF_TRF	Oil refineries and Transformation industry	
IND	Combustion for manufacturing	
CHE	Chemical processes	
IRO	Iron and steel production	
TNR_Aviation_CDS	Aviation climbing & descent	
TNR_Aviation_CRS	Aviation cruise	
TNR_Aviation_LTO	Aviation landing & takeoff	
TRO	Road transportation	
TNR_Other	Railways, pipelines, off-road transport	
TNR_Ship	Shipping	
PRO_coal	Fuel exploitation COAL	
PRO_oil	Fuel exploitation OIL	
PRO_gas	Fuel exploitation GAS	
FFF	Fossil fuel fires	
ENF	Enteric fermentation	Agriculture
MNM	Manure management	
AGS	Agricultural soils	

SWD_LDF	Solid waste landfills	Waste management
SWD_INC	Solid waste incineration	
WWT	Wastewater handling	

A2B2 Graphical representation of the main inventories used

CH₄ fluxes (g m⁻² s⁻¹)

985 **Figure A1B1: (a) Graphical representation of CH₄ fluxes due to fossil fuel exploitation as reported in appendix Table A1 by EDGAR inventories. The trajectory of the campaign is represented by the blue line. (b) Same as (a) for agriculture by EDGAR. (c) same as (a) for wetlands by ORCHIDEE. (d) same as (a) for freshwaters by blake4me&GLWD. (e) same as (a) for oceans by Weber.**

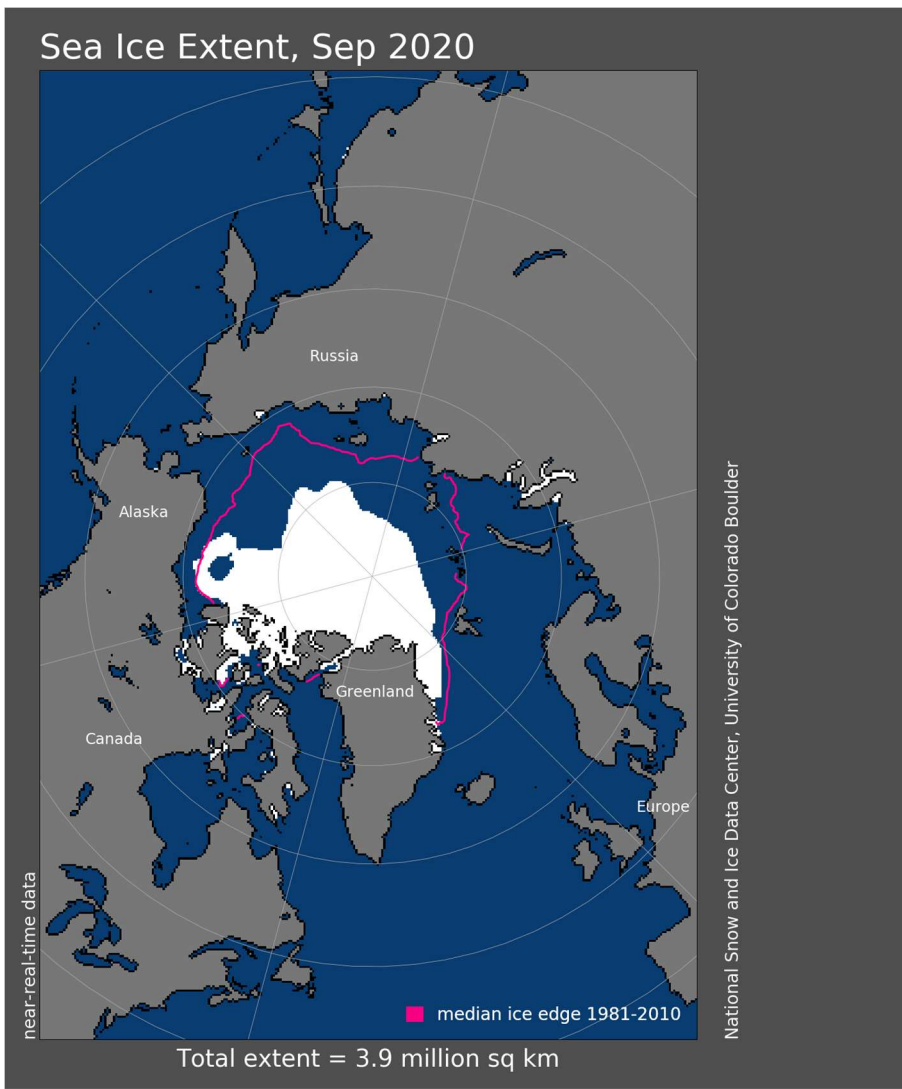



Figure B1: Sea ice extent in September 2020. As illustrated, there is no sign of ice in Bering Strait (Fetterer et al., 2017).

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Appendix C: Additional material concerning flight L3 characterization

C1 Potential emission sensitivity (PES) under 500 m map for flight L3 at 05:01 UTC
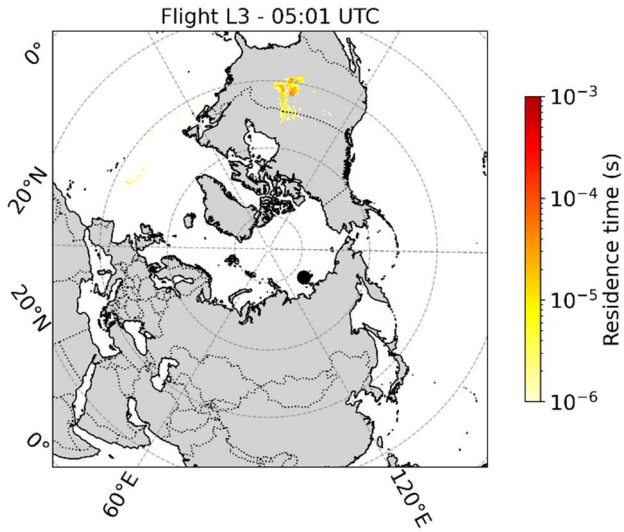
Figure C1: 10 d potential emission sensitivity (PES) with a threshold altitude of 500 m for particles released at 05:01 UTC at an altitude of 5256 m. The black dot represents the position of the receptor.

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Appendix D: Additional material concerning CH₄ freshwater enhancements in Naryan Mar region

~~C1~~**D1 Tundra around Naryan Mar**

Figure C1D1: **An aerial picture of the tundra around Naryan Mar revealing the presence of a huge number of lakes in the region.**
1005 **The picture was provided by Boris D. Belan.**