

Advancing CH₄ and N₂O retrieval strategies for NDACC/IRWG FTIR observations with the support of airborne in situ measurements

Ivan Ortega¹, James W. Hannigan¹, Bianca C. Baier², Kathryn McKain², and Dan Smale³

¹Atmospheric Chemistry Observations & Modeling, NSF National Center for Atmospheric Research, Boulder, Colorado, USA

²Global Monitoring Laboratory, National Oceanic and Atmospheric Administration, Boulder, Colorado, USA

³National Institute of Water and Atmospheric Research (NIWA), Lauder, New Zealand

Correspondence: Ivan Ortega (iortega@ucar.edu)

Abstract.

Atmospheric methane (CH₄) and nitrous oxide (N₂O) are potent greenhouse gases with significant impacts on climate change. Accurate measurement of their atmospheric abundance is essential for understanding their sources, sinks, and the impact of human activities on the atmosphere. Ground-based high-resolution Fourier Transform Infrared (FTIR) observations, employed by collaborative international initiatives like the Infrared Working Group (IRWG) within the Network for the Detection of Atmospheric Composition Change (NDACC), play a vital role in retrieving the atmospheric amounts of these gases. Network wide consistent data products rely on consistent observations and retrievals. Recent developments in spectroscopy, a priori data, retrieval software and techniques underscores the necessity to revisit the retrieval strategies for all NDACC/IRWG species currently ongoing. This study investigates various retrieval strategies of CH₄ and N₂O utilizing high-resolution FTIR observations in Boulder, Colorado, and compares them with unique airborne in situ measurements. The initial focus is on characterizing retrieval differences across spectroscopy databases. While it is challenging to identify the best retrievals purely based on spectroscopy, as they produce similar outcomes, notable differences in profile shapes and magnitudes underscore the importance of independent validation. Specifically, when multi-year independent nearby AirCore and aircraft in situ profile measurements are used to evaluate vertical distributions and biases in partial columns, they reveal excellent agreement in relative differences with FTIR retrievals and thereby strengthening confidence in the assessment. The final optimized retrievals for CH₄ and N₂O are presented incorporating quantitative fitting results and comparisons of vertical profiles, partial and total columns. We find that employing a priori profiles using the latest simulations of the Whole Atmosphere Community Climate Model (WACCM) enhances accuracy relative to in situ profiles. While the HITRAN 2020 spectroscopic database is effective for N₂O, ATM 2020 provides better results for CH₄, with slight improvement observed when paired with the water vapor line list from DLR; however, this improvement may be site-dependent. Regarding regularization, both first-order Tikhonov and Optimal Estimation produce comparable outcomes, as long as the fitted profile degrees of freedom remain between 2 and 2.5. Correspondingly, profile results comparisons yield biases of -0.08 ± 0.38 % and 0.89 ± 0.28 % for tropospheric and stratospheric layers of CH₄ relative to AirCore, respectively, and 0.39 ± 0.42 % for aircraft comparisons in the troposphere. For

N₂O, the bias in the troposphere using aircraft measurements is approximately 0.18 ± 0.2 %. Uncertainty budgets combining random and systematic sources are provided. Random errors, mainly stemming from temperature profile uncertainties and measurement noise dominate in the troposphere for both gases with a retrieval random error of 0.5 %. Systematic errors primarily arise from HITRAN based spectral line parameters, predominantly the line intensity and air-broadened half-width. ~~Finally, we present long-term time series of CH₄ derived from the recommended retrieval strategies applied to observations at Boulder. To contrast these findings with the southern hemisphere, we successfully extended this analysis to the site in Lauder, New Zealand.~~ These findings contribute to advancing our understanding of atmospheric composition and will support the improvement of a harmonized approach for all IRWG/NDACC sites.

1 Introduction

Atmospheric methane (CH₄) and nitrous oxide (N₂O) are potent greenhouse gases that significantly contribute to radiative forcing and, consequently, climate change. Accurate and precise measurement of these trace gases is essential for understanding their sources, sinks, and the impact of anthropogenic activities on the Earth's atmosphere. CH₄ is considered a short-lived greenhouse gas with an atmospheric lifetime of approximately 10 years. Despite its relatively short atmospheric lifetime compared to other greenhouse gases, CH₄ has a high global warming potential, 27 to 30 times that of CO₂ over a 100-year timescale, making it a significant contributor to near-term climate forcing. Its sources are diverse, including natural processes such as wetlands and anthropogenic activities such as livestock management and fossil fuel extraction (Canadell et al., 2021). In contrast, N₂O is classified as a long-lived greenhouse gas, with a lifetime of approximately 114 years. While less abundant, N₂O exhibits a remarkable warming potential, approximately 270 times greater per ton than CO₂ over a 100-year time horizon. This amplifies its impact on climate change and its role in stratospheric ozone depletion (Prather et al., 2015; Tian et al., 2020). Understanding the sources and sinks of both CH₄ and N₂O is crucial for developing effective climate mitigation strategies and policies. Moreover, as both gases are intricately linked to the global carbon and nitrogen cycles, studying their behavior aids in comprehending broader ecological processes and human-induced perturbations to the Earth's atmosphere (Canadell et al., 2021).

Ground-based Fourier Transform Infrared (FTIR) spectroscopy has proven to be a reliable method for extracting atmospheric concentrations of both CH₄ and N₂O through direct-sun solar absorption measurements. Collaborative international initiatives actively use FTIR observations to retrieve information on these species. The Infrared Working Group (IRWG) within the Network for the Detection of Atmospheric Composition Change (NDACC) has demonstrated the capability to retrieve both gases, along with numerous others, in the mid-infrared with higher spectral resolution and for long-term (De Mazière et al., 2018). Simultaneously, the Total Column Observation Network (TCCON) focuses on precision measurements of column-averaged abundances for CH₄ and N₂O in the near-infrared spectral region (Wunch et al., 2011). Additionally, the newly formed Collaborative Carbon Column Observing Network (COCCON) has employed portable low-resolution FTIR spectrometers to measure greenhouse gases, including CH₄ and N₂O (Hase et al., 2016; Alberti et al., 2022). While TCCON and COCCON

have optimized their retrievals through profile scaling to derive total columns, the IRWG/NDACC is able to obtain vertical profile information with higher spectral resolution, up to two or three degrees of freedom (DOF).

Presently, sites within IRWG/NDACC generally adhere to the recommendations outlined in Sussmann et al. (2011). However, as evidenced by Sepúlveda et al. (2014); Bader et al. (2017); and Olsen et al. (2017), a significant variability exists in the final retrieval strategies employed at each site. The retrieval strategy proposed by Sussmann et al. (2011) for CH₄ involved using the High-resolution TRANsmision molecular absorption database (HITRAN) 2001, which exhibited better agreement between measured and simulated spectra compared to newer databases of that time (e.g., HITRAN 2008, (Rothman et al., 2009)). This strategy utilized three out of five possible micro windows and incorporated Tikhonov L1 regularization, which focuses on stabilizing the solution by adding constraints to the optimization problem. In a more recent study, Chesnokova et al. (2020) analyzed total columns derived from various line lists, concluding that HITRAN 2001 offers better CH₄ retrievals. However, ATM19 (Toon, 2015; Toon et al., 2016) produced slightly different, yet comparable, results to HITRAN 2001. This conclusion is based on the optimal agreement observed between measured and simulated spectra. Considering the availability of updated spectroscopy, diverse retrieval strategies at each site, and new validation data, there is a pressing need to explore an optimal and harmonized strategy within the IRWG/NDACC. Additionally, while previous studies have focused mainly on total columns, IRWG/NDACC observations have the potential to separate both tropospheric and stratospheric columns for N₂O and CH₄. However, this has not yet been extensively validated.

This study investigates diverse retrieval strategies for CH₄ and N₂O, employing several micro-windows, a priori profiles, and regularization methods. High-resolution Fourier-transform infrared (HR-FTIR) observations in Boulder, Colorado form the basis of our analysis (Ortega et al., 2021). Initially, we focus on characterizing residuals, representing differences between simulated and observed spectra using different spectroscopy databases. A key aspect of our investigation involves integrating multi-year independent observations from AirCore (CH₄) and aircraft (CH₄ and N₂O) profiles obtained in close proximity to the FTIR site. These independent datasets are crucial for evaluating the vertical distribution (profile shapes) of the retrieval strategies. For CH₄, we examine the bias in tropospheric and stratospheric partial columns, while for N₂O, the assessment primarily focuses on tropospheric columns. Boulder offers a unique opportunity with its extensive, multi-year coverage of ~~semi-co-located-AirCore~~AirCore, aircraft, and IRWG/NDACC profiles, which sample CH₄ in close proximity, allowing for the integration of numerous observations into our analysis.

2 Observations

2.1 Ground-based FTIR

High-resolution (HR) direct-solar infrared absorption spectra have been recorded at the NSF National Center for Atmospheric Research (NSF NCAR) in Boulder, Colorado (40.40° N, 105.24° W, 1600 m.a.s.l) since 2010 to present, from 2010 to 2017 utilizing a Bruker 120 HR-FTIR spectrometer. Subsequently, the instrument underwent an upgrade throughout a significant portion of 2018, incorporating the latest electronics and optics from the Bruker 125 HR-FTIR. Measurements are conducted under clear-sky conditions. The site officially became a part of the Infrared Working Group (IRWG) within NDACC in 2020, and the

observations adhere to the standard measurement protocols established by the network. Optical bandpass filters are employed to optimize the signal-to-noise ratio (SNR) within the near and mid-infrared spectral range. [Hannigan et al. \(2009\) provide a list of typical band limits for each NDACC filter, along with the gases exhibiting absorption features and typically retrieved, covering a spectral range of 750–5000 cm⁻¹.](#) For CH₄~~NDACC filter number~~, NDACC filter 3 is used~~while for and~~, while for N₂O, ~~both~~ filters 3 and 4 can be used, ~~though filter 3 is predominantly used~~. The nominal spectral resolution achieved is 0.004 cm⁻¹ (~~maximum~~ optical path difference (OPD) of ≈ 257 cm), utilizing liquid nitrogen-cooled InSb and Mercury Cadmium Telluride (MCT) detectors and a KBr beam splitter. For a detailed description of the FTIR operations at Boulder see Ortega et al. (2019, 2021).

The spectra undergo analysis using the SFIT4 algorithm, an advancement from SFIT2 (Pougatchev et al., 1995; Rinsland et al., 1998; Hase et al., 2004; Hannigan et al., 2024), to extract profiles of CH₄ and N₂O. Key input and common parameters for the retrieval of all gases using SFIT4 include vertical profiles of pressure, temperature, and volume mixing ratios (VMR) of the atmospheric gases absorbing in the specific micro-windows. The input pressure and temperature vertical profiles are obtained from National Center for Environmental Prediction (NCEP) (Wild et al., 1995) and are standardized for use in all NDACC retrievals. These are daily average profiles that extend to up approximately 50 km and above that we use monthly mean pressure and temperature profiles from simulations using the Whole Atmosphere Community Climate Model (WACCM) (Garcia et al., 2007). As an initial step, and to mitigate water vapor interference, the retrieval of water vapor is conducted first. Subsequently, this water vapor information is utilized in the retrieval process for CH₄ and N₂O to prevent potential bias arising from inaccuracies in the water vapor profile shape (Ortega et al., 2019). Sections 3.1 and 3.2 describe in more detail the specific retrievals for CH₄ and N₂O, respectively.

2.2 AirCore ~~and aircraft~~ CH₄ profiles

We employ CH₄ AirCore data acquired from the National Oceanic and Atmospheric Administration (NOAA) Global Monitoring Laboratory (GML) to examine and assess the vertical distribution of CH₄ in the troposphere and stratosphere. Regular monthly AirCore flights are initiated in Boulder, Colorado, many of these flights occur at approximately 1 pm local time to synchronize with the OCO-2/A-train satellite overpass with air samples collected anywhere from 17 km to 121 km nearby to the FTIR (measured from the descent point). The AirCore technique employed by NOAA involves collecting air samples within a stainless steel tube (approx. 100 m in length and 1 cm in diameter). This tube passively collects atmospheric air during balloon descents from the middle stratosphere to the ground. Subsequently, the collected air is analyzed using a cavity ring-down spectrometer ensuring a trace mole fraction precision of better than 0.4 ppb for CH₄ (Karion et al., 2010). Because storage of the air sample in the tubing does not diffuse quickly between balloon payload landing and analysis, approximately 100 discrete samples are measured in the AirCore tubing in a 4-hour storage time, offering detailed insights into the vertical distribution of CH₄, carbon monoxide (CO), and carbon dioxide (CO₂). For our study, we leverage Level 2 CH₄ from the current NOAA AirCore dataset v20230831, which includes dry air mole fraction profiles retrieved since 2012 (Baier et al., 2021). ~~Our investigation encompasses~~ [Our investigation includes](#) profiles collected from 2018 to 2022, ~~allowing us to evaluate various retrieval strategies with a substantial number of observations, thereby facilitating the creation of statistically meaningful~~

findings enabling the evaluation of different retrieval strategies using a substantial set of coincident FTIR, AirCore, and aircraft observations.

125 Figure 1a shows individual CH₄ AirCore profiles obtained in Boulder, Colorado from 2018 to 2022. As expected, there is a noticeable rise in CH₄ mole fraction from 2018 onwards, reflecting the increase in the global background (https://gml.noaa.gov/ccgg/trends_ch4/). These profiles exhibit distinct features, such as elevated near-ground values in certain instances, suggestive of local sources. Further, within the boundary layer and free troposphere, the profiles demonstrate well-mixed characteristics with some variations in the upper troposphere and the lowest stratosphere of the AirCore profiles, culminating
130 in a sharp decrease at the tropopause. Figure 1b presents the mean vertical profiles from 2018 to 2022. To derive the mean profile, the individual profiles are re-gridded to a common altitude [scheme](#) levels within a range from 1.6 to 25 km and mean values are then estimated within 0.2 km resolution. Ultimately, mean and standard deviation profiles are calculated. The blue shaded area in Figure 1b represents the standard deviation in the AirCore profile dataset used here, while the red circle at the bottom represents the mean values from an in-situ sensor located at the Boulder reservoir about 5 km from the HR-FTIR using
135 the overlap period between 2020 to 2022 (<https://www.bouldair.com/boulder.htm>), providing insight into the variability around the Boulder region.

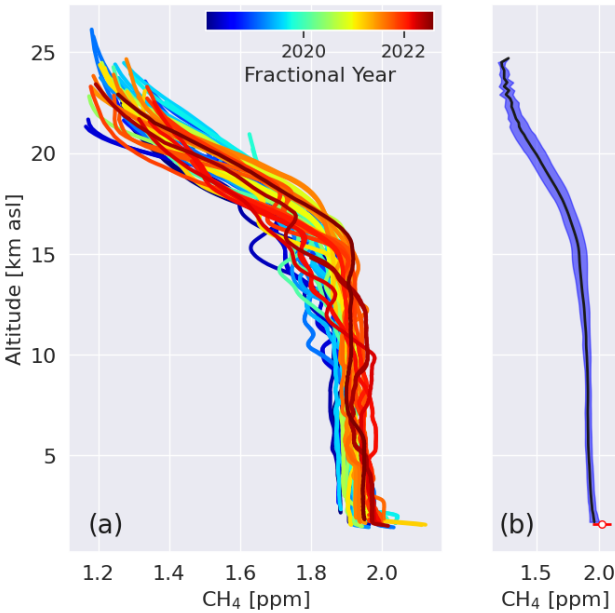


Figure 1. (a) Individual AirCore CH₄ profiles in Boulder, Colorado, color-coded to indicate fractional years spanning from 2018 to 2022. (b) Mean vertical profile of CH₄ accompanied by standard deviation, with the red circle at the bottom representing the mean volume mixing ratio from an in-situ sensor near the HR-FTIR (see text for details). A total of 61 vertical profiles are included in these plots.

2.3 Aircraft-based observations of CH₄ and N₂O profiles

To further evaluate CH₄ and N₂O retrievals we use the vertical profiles obtained in a remote location nearby Briggsdale, Colorado (40.6347 N, 104.3269 W), approximately 100 km northeast of the HR-FTIR, as part of the NOAA/ESRL Global Greenhouse Gas Reference Network Aircraft Program (Sweeney et al., 2015; McKain et al., 2023) (<https://gml.noaa.gov/ccgg/aircraft/>). This program employs contracted aircraft equipped with equipment to collect 12 discrete air samples in vertical profiles, primarily over North America, and monitor over 50 trace gases throughout the boundary layer and free troposphere over the long term. The network provides crucial insights into the sources, sinks, and transport mechanisms of greenhouse and other trace gases. Additionally, these profiles are a resource for model evaluation of vertical mixing and independently constrain the mean biospheric uptake of CO₂ throughout the annual cycle. The routine data collection within the program further facilitates the evaluation of satellite retrievals, which is often complemented by ground-based remote sensing observations. Figure 2 illustrates the CH₄ and N₂O aircraft profiles acquired at Briggsdale, Colorado, spanning from 2018 to 2022. In line with CH₄ AirCore profiles, there is minimal variation observed in the profiles up to 6 km, with occasional enhancements near the ground on certain days. In the case of N₂O, there is a clear increase from 2018 onwards, also consistent with the increase in the global background (https://gml.noaa.gov/ccgg/trends_n2o/). The N₂O profiles exhibit a well-mixed pattern, with no notable enhancements near the ground. Similar to Figure 1, the mean profiles are presented on the right side for each gas, depicting mainly the increase in the global background. However, the mean profile is displayed within the altitude range of 1.6 to 7 km asl, with a resolution of 0.4 km due to the original vertical resolution being lower than that of the AirCore.

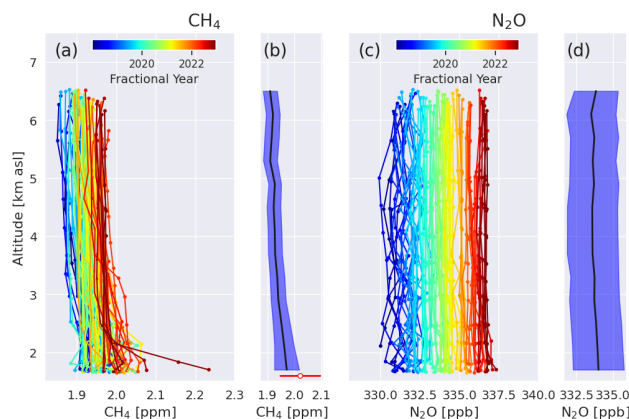


Figure 2. (a) NOAA aircraft profiles of CH₄ at Briggsdale, Colorado, with different colors indicating fractional years between 2018 and 2022. (b) Mean and standard deviation of CH₄ profiles derived from the NOAA aircraft, along with surface CH₄ volume mixing ratio (vmr) from the Boulder reservoir, including standard deviation. (c) Same as (a) but for N₂O. (d) Same as (b) but for N₂O. A total of 81 vertical profiles are included in these plots.

3 Retrieval strategies

155 3.1 CH₄

Multiple retrieval strategies have been devised, encompassing various combinations of micro-windows, spectroscopy, and regularization techniques. The fitting results are employed to assess different spectroscopy databases, and all retrieval strategies are compared with AirCore and aircraft observations to determine the most effective approach. An optimized selection of micro-windows (MW) in the mid-infrared has been carried out by Sussmann et al. (2011) and a summary of these retrieval strategies is provided in Table 2. In alignment with Sussmann et al. (2011), we employed the same five potential micro-windows for our analysis, detailed in Table 1.

The molecular absorption databases employed in this study include the latest official release of HITRAN 2020 (HIT20) (Gordon et al., 2022), the ATM 2020 (ATM20) spectroscopic database used by TCCON (Toon, 2015; Toon et al., 2016), which is available at <https://mark4sun.jpl.nasa.gov/toon/linelist/linelist.html> (last accessed: 22 Dec 2023) (last accessed: 22 Dec 2023), and HITRAN 2000 (HIT00) (including the 2001 updates), as recommended by Sussmann et al. (2011). In their study, Sussmann et al. (2011) also tested HITRAN 2008, which was the most recent database at the time, and found significant residuals with HITRAN 2008, while HIT00 showed much lower residuals. More recently, Chesnokova et al. (2020) examined the impact of different databases and similarly concluded that HIT00 provides superior fitting results, with the exception of ATM, 2019 at the time. Additionally, we utilized combinations of these databases with others for specific purposes. For instance, a water vapor spectroscopy is based on the line list provided by the German Aerospace Center (DLR) (Zhou et al., 2018), while ATM20 is used in conjunction with the HIT00 linelist for CH₄. Section 4.1 provides a more detailed overview of the various spectroscopy databases and their corresponding results.

For the current IRWG gas retrievals, mean a priori profiles were primarily obtained using WACCM version 4 for the 1980 - 2020 period (referred to as WACCM4). WACCM simulations underwent enhancements in physical, chemical, and aerosol parameterizations (Gettelman et al., 2019), resulting in version 6 (WACCM6) and new mean a priori profiles have been created spanning the 1980 - 2040 period (<https://wiki.ucar.edu/pages/viewpage.action?pageId=452264118>). Comparing CH₄ between these two WACCM versions at different sites (e.g., Mauna Loa, Hawaii; Thule, Greenland) with independent profiles obtained during the airborne campaign Pole-to-Pole Observations (HIPPO; www.eol.ucar.edu/field_projects/hippo) (Wofsy, 2011), and Atmospheric Tomography Mission (ATom; <https://daac.ornl.gov/ATOM/campaign/>) (Thompson et al., 2022) field campaigns indicates that the latest version 6 profiles are more suitable for recent years (not shown). To streamline the retrieval process all retrievals utilized the most recent version of WACCM profiles. For regularization, the process of adding constraints or penalties to the optimization problem to ensure a more stable and reliable solution, we employed both the first-order Tikhonov regularization, following the recommendation of Sussmann et al. (2011), and the conventional optimal estimation (OEM) approach. In Tikhonov regularization, the regularization strength (α) is optimized to maintain degrees of freedom (DOF) at approximately 2, with potential variations depending on conditions. Optimization is achieved through the typical L-curve method, as proposed by Steck (2002). The Tikhonov regularization procedure is directly implemented in the latest version of SFIT4, accessible at <https://wiki.ucar.edu/display/sfit4/SFIT+Core+Code> (last accessed: 22 Dec 2023). For the OEM approach,

regularization involved diagonal a priori covariance matrices. Variances magnitudes were empirically tuned to prevent profile oscillations and similarly to achieve a target DOF of around 2. The EOM technique has the advantage that the retrieved vertical profile relaxes back to the a priori profile shape, as spectral information diminishes with altitude. In contrast, the Tikhonov constraint tends to preserve the deviation from the a priori at higher altitudes where spectral information is still present. Additional details on the retrieval workflow in Boulder are provided in Ortega et al. (2021).

We establish a baseline retrieval strategy (r1) by employing three micro-windows (1, 3, and 5), along with the first-order Tikhonov regularization approach, and the HIT00 spectroscopy database (Sussmann et al., 2011). A total of ten retrieval strategies have been developed to explore various approaches and determine whether different methods impact the results when compared with AirCore and aircraft profiles. As shown in Section 4.1, HIT20 was excluded due to its significantly higher residuals in the fitting results. Detailed information on each retrieval strategy is provided in Table 2.

Table 1. Summary of the 5 micro windows tested in the the retrieval of CH₄.

	Micro-windows (cm ⁻¹)	Main species
1	2613.70 – 2615.40	CH ₄ , HDO, CO ₂
2	2650.60 – 2651.30	CH ₄ , HDO, CO ₂
3	2835.50 – 2835.80	CH ₄ , HDO
4	2903.60 – 2904.03	CH ₄ , NO ₂ , HDO, H ₂ O
5	2921.00 – 2921.60	CH ₄ , H ₂ O, HDO, NO ₂

Table 2. Different strategies tested in the retrieval of CH₄.

Retrieval	MWs	Spectroscopy details	Regularization	Additional details
r1	1,3,5	HIT00	Tikhonov L1	Baseline following Sussmann et al. (2011)
r2	1,3,5	ATM20	Tikhonov L1	ATM20 for all gases
r3	1,3,5	ATM20 and HIT00 (CH ₄)	Tikhonov L1	Same as r2 but HIT00 for CH ₄
r4	1,2,3,4,5	ATM20 and HIT00 (CH ₄)	Tikhonov L1	5 Micro-windows
r5	1,2,3,4,5	ATM20	Tikhonov L1	5 Micro-windows and ATM20 for all gases
r6	1,2,3,5	ATM20	Tikhonov L1	4 Micro-windows and ATM20 for all gases
r7	1,2,3	ATM20	Tikhonov L1	3 Micro-windows and ATM20 for all gases
r8	1,2,3,4,5	ATM20 and DLR (H ₂ O)	Tikhonov L1	DLR water vapor and ATM20 all others
r9	1,2,3,4,5	ATM20 and DLR (H ₂ O)	OEM	same as r8 but using OEM
r10	1,2,3,4,5	ATM20	Tikhonov L1	Same as r5 but using monthly mean WACCM6

3.2 N₂O

The N₂O retrievals suggested by the IRWG involve the use of four micro-windows within the 2480 – 2540 cm⁻¹ spectral range. The specific micro-windows and associated interfering species are detailed in Table 3. Our focus is not directed towards

the identification or refinement of micro-windows since they are well-harmonized across the network, yielding satisfactory results. Instead, our objective is to assess the performance of the latest spectroscopy. Similarly, the spectroscopy evaluation includes testing HITRAN 2008 (HIT08) (the current official IRWG recommendation), HIT20, and ATM20. Within the retrieval strategies outlined in Table 4, we incorporate both the Tikhonov L1 and OEM approaches since the regularization approach is not uniformly harmonized (Zhou et al., 2019). Additionally, two optical band pass filters, denoted as filters 3 and 4 within the IRWG, are available for utilization in the N₂O retrieval process. In this case, we conducted separate N₂O retrievals using each filter independently, as well as ~~employing a retrieval strategy~~ a retrieval approach where both filters ~~are utilized simultaneously~~ were used independently, but the results were combined without distinguishing between filters. In this instance, we incorporate retrieval strategies utilizing both WACCM4 and v6, as there is a comparatively less number of sensitivity studies compared to CH₄.

Table 3. Summary of the 4 micro windows used in the the retrieval of N₂O.

	Micro-windows (MW)	Main species
1	2481.30 – 2482.60	N ₂ O, H ₂ O, CO ₂ , CH ₄
2	2526.40 - 2528.20	CH ₄ , N ₂ O, CO ₂ , HDO, CH ₄
3	2537.85 - 2538.80	N ₂ O, CH ₄
4	2540.10 - 2540.70	N ₂ O

Table 4. Different strategies tested in the retrieval of N₂O.

Retrieval	Spectroscopy details	Regularization	Additional details
r1*	HIT08	OEM	filter 3 and WACCM4
r2	HIT20	OEM	filter 3 and WACCM4
r3	ATM20	OEM	filter 3 and WACCM4
r4	HIT20	OEM	filter 4 and WACCM4
r5	HIT08	OEM	filter 3 and WACCM6
r6	HIT20	Tikhonov L1	filter 3 and WACCM6
r7	HIT20	Tikhonov L1	filter 4 and WACCM6
r8	HIT20	Tikhonov L1	filters 3, 4 and WACCM6

*Current retrieval strategy as suggested by the IRWG

4 Results

4.1 Inter-comparison among spectroscopy databases

This section addresses the quality of fits resulting from various spectroscopy databases. Specifically, for CH₄ we compare five databases: (1) HIT00; (2) HIT20; (3) ATM20; (4) ATM20 with DLR water vapor (ATM20_DLRWV); and (5) ATM20 with HIT00 for CH₄ (ATM20_HIT00CH4). To maintain focus on spectroscopy, all five retrievals follow the same procedure, employing Tikhonov regularization and WACCM6 as the a priori, with the only variation being the choice of spectroscopy. Figure 3 visually shows the spectral contributions of various species during the CH₄ retrieval across the five micro-windows. As an example, the top panel displays these contributions for each micro-window on November 18th, 2019, at a solar zenith angle of 60.0°. The observed and fitted lines, presented in blue and green, respectively, provide insights into the quality of the retrieval. In the corresponding bottom plots, the residuals (observed minus fitted) are presented, with distinct colors indicating the different spectroscopy databases. There are clear differences for some micro-windows. For instance, HIT20 exhibits substantial systematic residuals for micro-windows 2 and 5. The large residual at 2921.33 cm⁻¹ in HIT20 for micro-window 5, previously identified by Sussmann et al. (2011) for HIT08, suggests that HIT20 still needs to be improved for this line strength. To further identify differences, the numbers within the labels provide the root mean square fitting for each micro-window across diverse spectroscopy databases. Micro-window 1 stands out as the only one where all databases perform similarly, while others exhibit variations. Nonetheless, ATM20 demonstrates slightly better results within micro-window 1. Apart from HIT20, which displays high residuals in micro-window 2, all other databases show similar residuals. In micro-window 3, ATM20, ATM20_DLRWV, and HIT20 ~~yield-produce~~ improved results, ~~while-whereas in~~ micro-window 4 ~~shows superior-residuals-~~ lower residuals are observed with ATM20_ ~~HIT04CH4~~ HIT00CH4, HIT20_ ~~HIT04CH4~~ HIT00CH4, and HIT00, ~~suggesting-indicating~~ that HIT00 remains ~~preferable-the preferred choice~~ for this micro-window. As previously noted, HIT20 continues to show large residuals in micro-window 5, whereas HIT00 still provides better residuals. A summary of root mean square values (RMS) for all micro-windows for a full-year (2021) of retrievals is provided in Table 5. Overall, HIT20 displays larger RMS, suggesting it may not be an optimal choice, while all other spectroscopy databases show similar RMS values within the standard deviation. These findings align with Chesnokova et al. (2020), where HIT00 yields better RMS but is similar to ATM20. However, there are notable differences in the profile shapes and magnitudes (see section 4.2), highlighting the importance of the assessment conducted with independent observations.

For N₂O, we compare the databases HIT08, HIT20, and ATM20. Our objective is to assess whether HIT20 or ATM20 exhibits better residual quality than HIT08, currently employed in the IRWG/NDACC strategy. Figure 4 illustrates the spectral contributions of various species during the N₂O retrieval across four micro-windows. Similar to CH₄, the top panels showcase an example, specifically on January 31, 2019, at a zenith angle of 63.8°. The bottom residual plots do not reveal a significant difference among the three databases. While the RMS suggests a slight improvement with HIT20, an overview table (6) presenting results for the entire year (2021) indicates no significant improvement. Nonetheless, this suggests that either HIT20 or ATM20 could be considered in the IRWG recommendation.

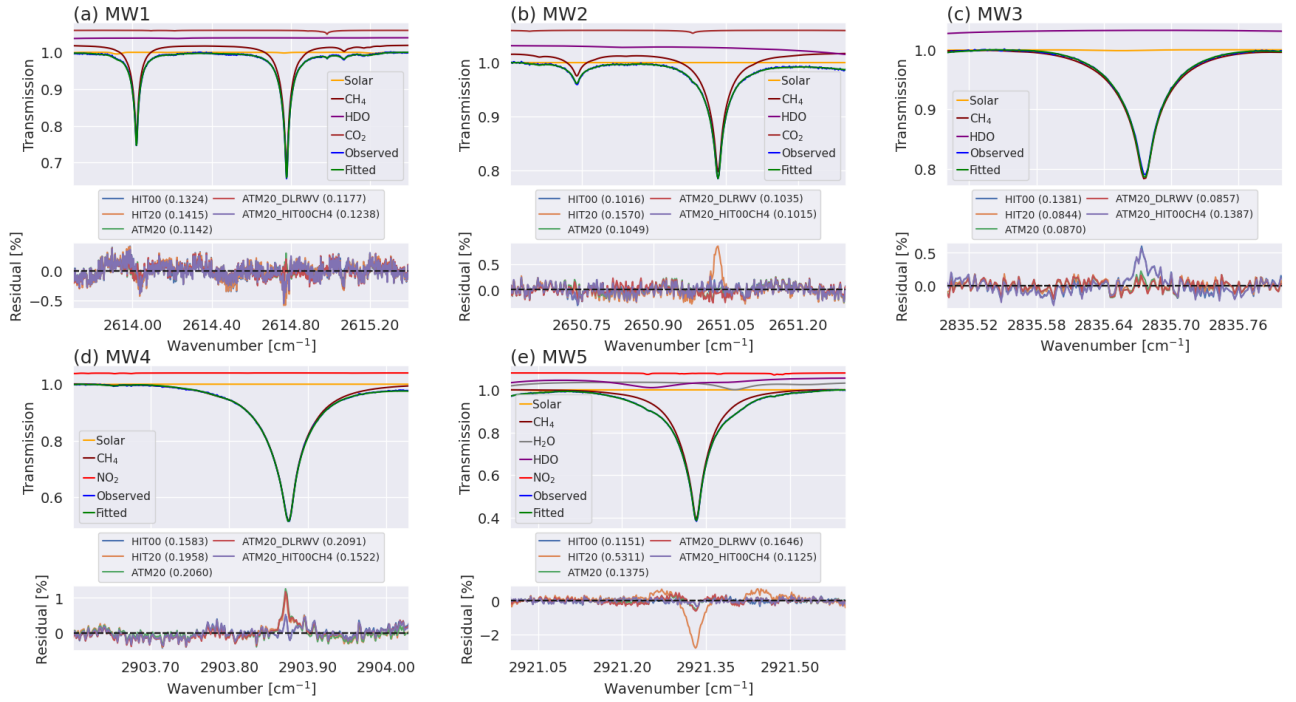


Figure 3. Retrieval fit example of CH_4 for November 18th, 2019, with a solar zenith angle of 60.0° . Spectral contributions for various species are displayed above each micro-window, spanning from micro-window 1 (a) to micro-window 5 (e) employing Tikhonov regularization and WACCM6 as the a priori. The observed and fitted lines are represented in blue and green, respectively. The bottom plots shows the residuals (observed minus fitted), with distinct colors indicating different spectroscopy types as labeled. The numbers within the labels denote the root mean square fitting for each micro-window across various spectroscopy databases.

Table 5. Overview of the mean root mean square (RMS) values across all five micro-windows and the overall RMS for CH_4 retrieval using various spectroscopy databases for 2021. The last column displays the degrees of freedom (DOF). Values in parenthesis are standard deviation.

Spectroscopy	RMS						DOF
	MW1	MW2	MW3	MW4	MW5	Overall	
HIT00	0.140 (0.022)	0.125 (0.022)	0.160 (0.038)	0.169 (0.033)	0.168 (0.038)	0.156 (0.030)	2.46 (0.15)
HIT20	0.148 (0.024)	0.164 (0.035)	0.115 (0.016)	0.205 (0.042)	0.504 (0.058)	0.270 (0.039)	2.31 (0.13)
ATM20	0.139 (0.018)	0.122 (0.017)	0.116 (0.018)	0.204 (0.036)	0.183 (0.032)	0.161 (0.025)	2.42 (0.16)
ATM20_DLRWV	0.140 (0.017)	0.134 (0.022)	0.117 (0.018)	0.206 (0.039)	0.210 (0.051)	0.170 (0.032)	2.41 (0.16)
ATM20_HIT00CH4	0.143 (0.022)	0.120 (0.019)	0.160 (0.037)	0.164 (0.031)	0.142 (0.024)	0.150 (0.027)	2.47 (0.14)

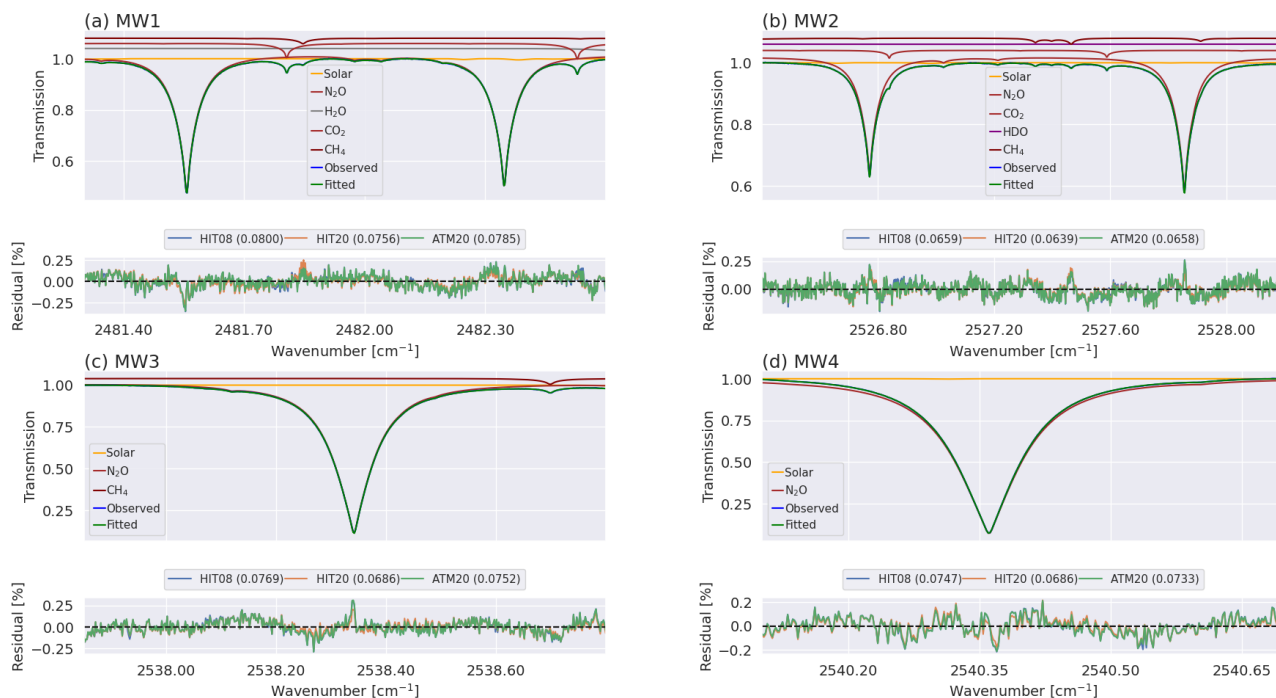


Figure 4. Retrieval fit example of N_2O on January 31th, 2019 with a solar zenith angle of 63.8° . Spectral contributions for various species are displayed above each micro-window, spanning from micro-window 1 (a) to micro-window 4 (d) employing Tikhonov regularization, WACCM6 as the a priori, and filter 4. The observed and fitted lines are represented in blue and green, respectively. The bottom plots shows the residuals (observed minus fitted), with distinct colors indicating different spectroscopy types as labeled. The numbers within the labels denote the root mean square fitting for each micro-window across various spectroscopy databases.

Table 6. Summary of the RMS values for N_2O retrieval in 2021, encompassing all five micro-windows and the overall RMS. The last column presents the DOF, with standard deviation values shown in parentheses.

Spectroscopy	RMS					DOF
	MW1	MW2	MW3	MW4	Overall	
HIT08	0.149 (0.031)	0.130 (0.019)	0.133 (0.029)	0.132 (0.032)	0.148 (0.030)	2.56 (0.11)
HIT20	0.148 (0.031)	0.131 (0.019)	0.131 (0.028)	0.128 (0.032)	0.147 (0.029)	2.57 (0.11)
ATM20	0.149 (0.031)	0.130 (0.019)	0.133 (0.029)	0.131 (0.032)	0.148 (0.029)	2.57 (0.11)

4.2 Inter-comparison with AirCore and aircraft profiles

Between 2018 and 2022, a total of 61 AirCore sampling systems were launched in Boulder, 36 which were coincident FTIR measurements. The total number of coincident dates between FTIR and aircraft profiles is 51. To enable a quantitative comparison of insitu profiles and partial columns, vertical profile measurements from AirCore and aircraft are re-gridded onto the

FTIR retrieval's altitude grid using spline order 1 interpolation. Subsequently, these profiles are smoothed with the daily mean FTIR averaging kernels to account for the FTIR altitude sensitivity (Rodgers and Connor, 2003). Figure 5a presents an illustrative comparison between FTIR profiles obtained on a given day and an AirCore profile. The high vertical resolution AirCore profile is depicted with black triangles, while the smoothed AirCore profile is shown with blue triangles. FTIR retrievals from the same day are represented by circles in various colors. In Figure 5b, the relative difference in percentage (or bias), defined as the difference between FTIR and AirCore values divided by the AirCore values, is shown for each corresponding time. To maximize the number of profile comparisons, and considering the relatively stable nature of CH_4 (and N_2O), we compare FTIR profiles obtained within a 6-hour time window of the AirCore descent time. The same approach is undertaken when comparing with aircraft observations but the aircraft profiles generally extend up to around 6 km asl altitude whereas the aircore profiles extend up to 20 km or higher.

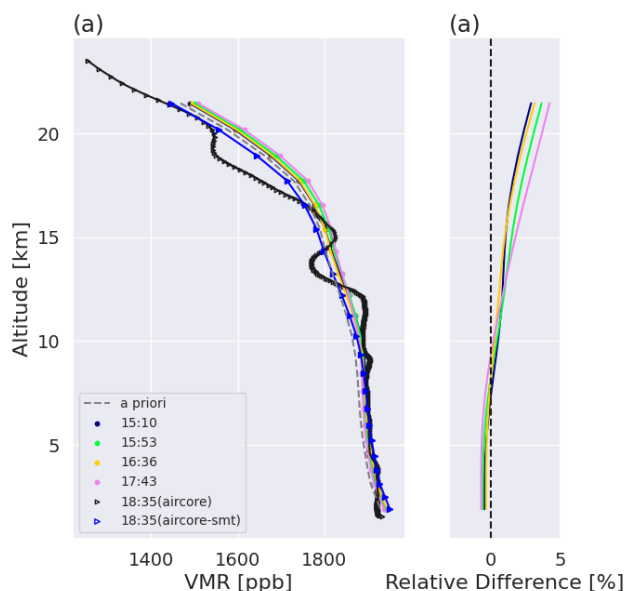


Figure 5. (a) Illustrative comparison between FTIR and AirCore CH_4 profiles on May 14th, 2019, utilizing ATM20, WACCM6, and Tikhonov regularization (retrieval r4 in Table 2). Various colors denote different times (UT) of HR-FTIR retrievals, with the high-resolution AirCore profile presented in black and the smoothed profile in blue. (b) Relative difference (FTIR minus AirCore) for each corresponding time.

4.2.1 CH_4

Figure 6 shows the relative differences in CH_4 vertical profiles obtained for each retrieval strategy when compared with AirCore profiles. The sub-panel names correspond to the retrievals listed in Table 2. The individual profile relative differences are displayed in gray, while the mean and median profiles are depicted in red and blue, respectively. To highlight the similarity between AirCore and aircraft profile results, the median relative differences in the troposphere from aircraft profiles are displayed in green in the same figure. The relative differences with aircraft profiles in the troposphere (up to 6 km) are further

detailed in Figure 7. Despite the absence of coincidence between AirCore and aircraft profiles, the relative differences exhibit remarkable similarity in both magnitude and shape, enhancing confidence in the assessment of multiple retrievals. It is interesting to observe that changing only the CH₄ database spectroscopy leads to a substantial change in the magnitude of profile differences. For example, retrievals r1 and r2 show opposite signs in their relative differences within both the troposphere and stratosphere, despite having identical retrieval parameters, apart from the spectroscopy. The key distinction between retrievals r1 and r2 is that r1 employs HIT00 and r2 relies on ATM20. Interestingly, using a combination of ATM20 and HIT00 for CH₄ yields profile differences that are very similar to those obtained using only HIT00, suggesting that the line parameters for CH₄ primarily dictate the profile shape. The inclusion of 3 or 5 micro-windows does not change the shape of the profile difference; however, r5, which uses 5 micro-windows, shows smaller relative differences in both the troposphere and stratosphere compared to r2, which uses the same settings but with only 3 micro-windows. Retrieval strategies showing favorable bias outcomes are encompassed in r7 to r10. Strategy r7 utilizes micro-windows 1, 2, and 3, while r8 employs all five micro-windows, deviating from the micro-windows 1, 3, and 5 recommended by (Sussmann et al., 2011) Sussmann et al. (2011). We observe slightly higher variability in r7, whereas r8 and r9 exhibit more compact variability, particularly in the troposphere. Retrieval r8 is identical to r9, except that r9 uses the OEM approach. Additionally, using monthly mean CH₄ a priori profiles does not significantly improve biases, as seen when comparing r10 to r5. It's important to note that past studies have predominantly focused on total column analysis. In such cases, retrieval strategies might produce biases in the troposphere and stratosphere with opposite signs, potentially canceling each other out and reducing the overall bias in total column measurements. In contrast, this work focuses on examining the differences at various altitudes within the vertical profiles, which, due to vertical sensitivity, ultimately lead to distinct biases in tropospheric and stratospheric columns, as well as in total column measurements.

The retrieval methodology implemented in SFIT4 encompasses the vertical profile but is confined in sensitivity to specific altitude ranges. Figure 8a shows typical averaging kernels (row kernels) employed in the CH₄ retrieval, while Figure 8b portrays the total column averaging kernel and the cumulative sum of DOF, denoting the number of independent layers in the retrieved profile. Shaded rectangles separate the cumulative sum of DOF, with the first DOF spanning 1.6 - 9.8 km and the second DOF ranging from 9.8 - 22.1 km. Note, that all profile regularization constraints of the retrieval strategies have been adjusted to attain similar DOF, although averaging kernels may differ, particularly between those utilizing Tikhonov and OEM due to the inherent nature of the approaches. ~~For a comprehensive evaluation of~~ To comprehensively evaluate these layers, ~~we conduct~~ a comparative analysis with AirCore and aircraft data ~~is conducted, emphasizing the mixing ratio, using mixing ratios~~ weighted by air mass density within the specified altitude ranges mentioned above ~~(see Ortega et al. (2021) for details on weighted mixing ratios)~~. Tables 7 and 8 list the mean and median CH₄ relative differences among various retrieval strategies for different partial columns using AirCore and aircraft profiles, respectively. The last column represents the 95 % confidence interval (CI) calculated. As indicated earlier, tropospheric biases estimated from both AirCore and aircraft observations reveal congruent findings. ~~An integral facet is the assessment of the a priori profile when contrasted with AirCore and aircraft profiles. When comparing WACCM6 across different partial columns, the biases are -1.9 ± 1.0 % and -1.5 ± 1.3 % for the troposphere and stratosphere, respectively. In principle, one might anticipate that retrievals would yield improved and lower biases. Correspondingly, in profile results, Strategy r8 demonstrates satisfactory values~~ performance, with mean biases of -0.08

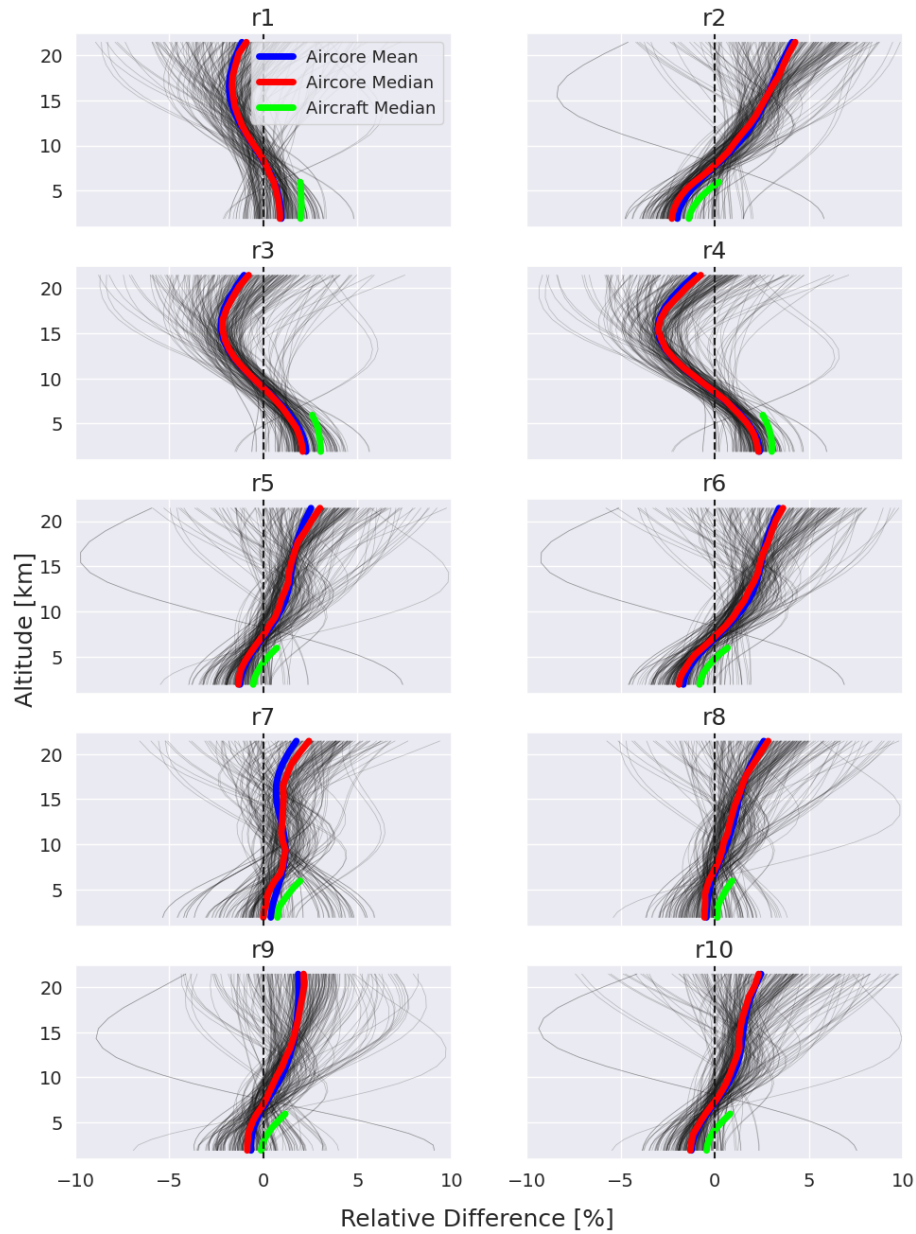


Figure 6. Relative differences in CH₄ profiles, calculated as the difference between FTIR and AirCore values divided by the AirCore values, employing diverse retrieval strategies. The mean and median profiles are shown in blue and red lines, respectively. The green lines illustrates the relative differences obtained in the lower troposphere using aircraft comparisons, as shown in Figure 7.

300 $\pm 0.38 \%$ and $0.89 \pm 0.28 \%$ for the tropospheric and stratospheric layers, respectively, and $0.39 \pm 0.42 \%$ for aircraft comparisons in the troposphere. Strategy r9, which is identical to r8 but utilizes the OEM approach, produces very similar results. Additionally, we assessed the a priori profile by comparing it with AirCore and aircraft profiles. When evaluating

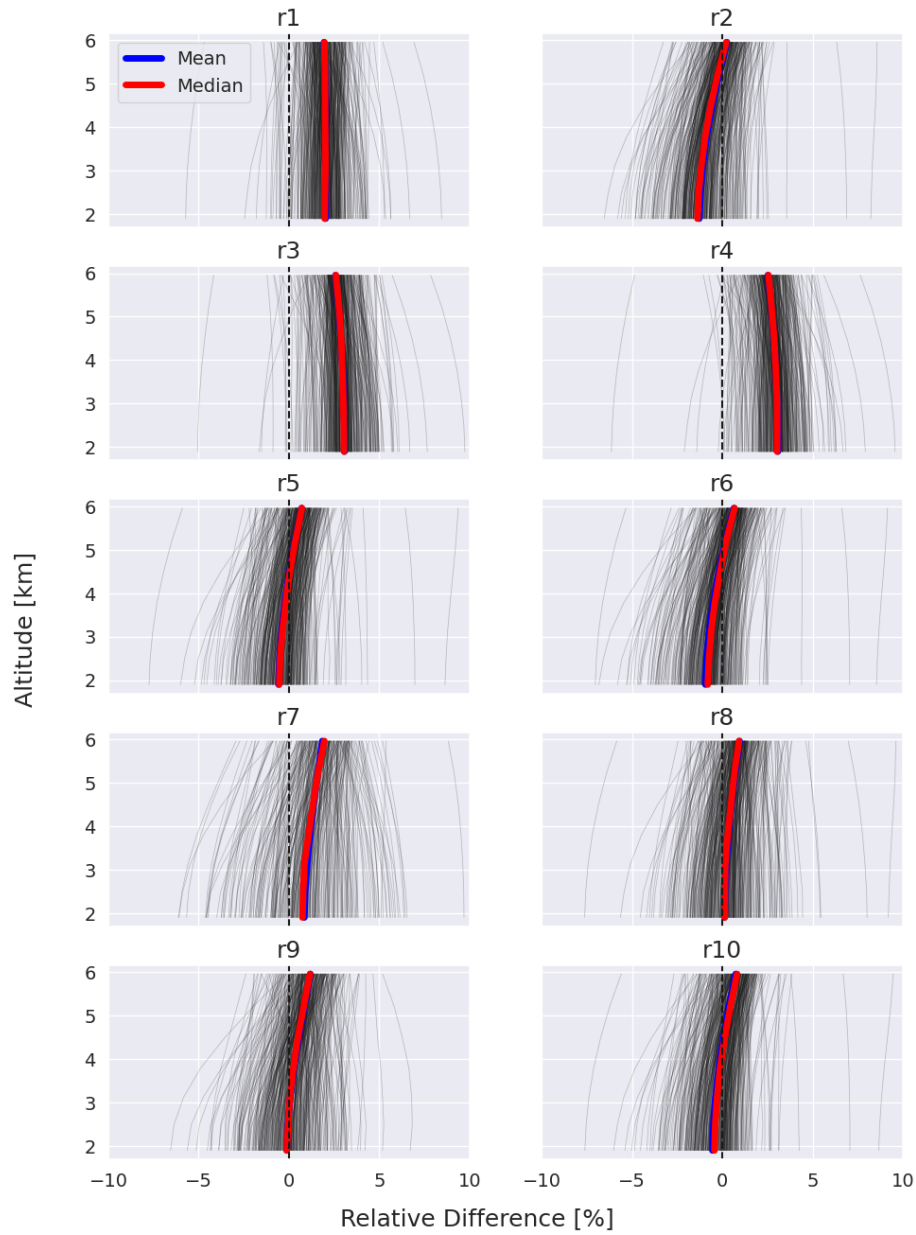


Figure 7. Relative difference between FTIR and NOAA aircraft CH_4 profiles using several retrieval strategies.

WACCM6 across different partial columns, the biases are -1.9 ± 1.0 % and -1.5 ± 1.3 % for the troposphere and stratosphere, respectively. Retrievals yield improved and lower biases, with r8 and r9 showing significant reductions.

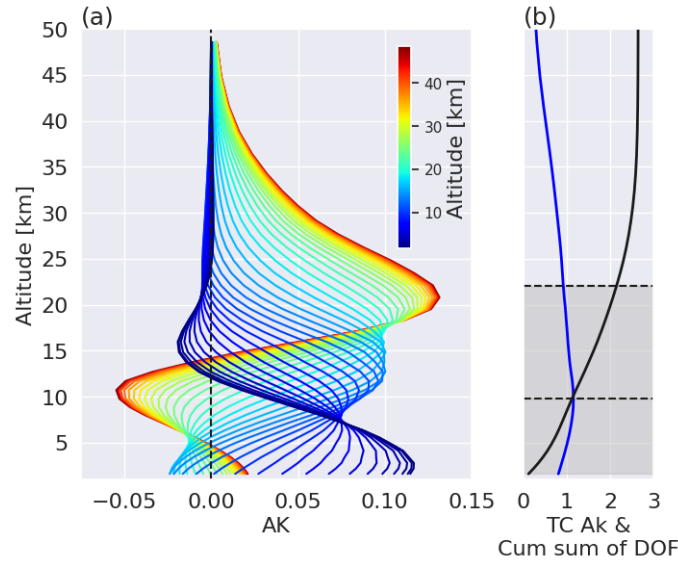


Figure 8. (a) Averaging kernel for CH₄, representative of November 18th, 2019, at a solar zenith angle of 60.0°. (b) Total column averaging kernel in blue and cumulative sum of DOF in black. The shaded rectangles indicate the cumulative sum of degrees of freedom (DOF). The first DOF spans 1.6 - 9.8 km, while the second DOF ranges from 9.8 - 22.1 km. This example employs 5 micro-windows, Tikhonov regularization, and WACCM6 as the a priori.

4.2.2 N₂O

305 To evaluate N₂O, we employ a comparable approach, but use only low to mid tropospheric values from aircraft profiles. Figure 9 shows the vertical profile differences across all retrieval strategies, and a summary of tropospheric weighted mixing ratio biases is presented in Table 9. In the current retrieval strategy (r1) recommended by the IRWG, there is a systematic positive bias of 1.58 ± 0.28 % at all tropospheric altitudes. Transitioning to the latest HIT20 spectroscopy (r2) reduces the overall tropospheric bias to 0.12 ± 0.27 %, although the profile's relative differences shift from negative values below 3 km to positive values above 3 km. Interestingly, employing ATM20 (r3) also results in a systematic positive bias of 1.42 ± 0.27 %, similar to HIT08. Similar outcomes are observed when using both filters independently. Switching to WACCM6 yields a slight improvement. For ~~instance, in example, in retrieval~~ r5, which ~~employs utilizes~~ HIT08 with the ~~new updated~~ WACCM6 profile, the difference is 1.27 ± 0.28 %. ~~Further improvement is achieved by applying the Tikhonov approach (The combination of HIT20 and WACCM6 further improves the comparison, as seen in retrievals r6 (-0.02 ± 0.25 %) for filter 3 and r7)-using~~ ~~HIT20~~ (0.36 ± 0.22 %) for filter 4, both employing the Tikhonov approach. Retrieval r8, ~~combining which uses~~ both filters, ~~demonstrates exhibits~~ minimal bias ($0.10-0.18 \pm 0.20$ %), ~~suggesting the successful applicability of both demonstrating the feasibility of using~~ filters 3 and 4 across the IRWG/NDACC network. ~~The Additionally, combining both filters maximizes the number of available retrievals. The OEM retrieval approach was also tested with HIT20 and WACCM6, yielding similar results to the Tikhonov approach. Specifically, the OEM-based retrieval showed a bias of 0.02 ± 0.23 %, which falls within~~

Table 7. CH₄ bias results among the different retrieval strategies for the different partial columns using AirCore profiles as the reference.
The results in bold indicate the best performance.

Retrieval #	Partial Column [km asl]	Mean bias [%]	Median bias [%]	CI [%]
r1	1.6 - 10.0	0.88	0.84	0.31
	10.0 - 20.0	-1.39	-1.26	0.32
	1.6 - 20.0	0.48	0.34	0.23
r2	1.6 - 10.0	-0.97	-1.32	0.36
	10.0 - 20.0	1.91	1.99	0.27
	1.6 - 20.0	-0.46	-0.66	0.27
r3	1.6 - 10.0	1.79	1.70	0.30
	10.0 - 20.0	-1.70	-1.60	0.32
	1.6 - 20.0	1.18	1.15	0.23
r4	1.6 - 10.0	1.71	1.80	0.28
	10.0 - 20.0	-2.37	-2.30	0.36
	1.6 - 20.0	0.99	1.05	0.21
r5	1.6 - 10.0	-0.64	-0.49	0.27
	10.0 - 20.0	1.20	1.19	0.25
	1.6 - 20.0	-0.32	-0.18	0.21
r6	1.6 - 10.0	-0.75	-0.99	0.34
	10.0 - 20.0	1.86	1.99	0.28
	1.6 - 20.0	-0.29	-0.40	0.26
r7	1.6 - 10.0	0.97	0.52	0.61
	10.0 - 20.0	0.52	1.00	0.60
	1.6 - 20.0	0.89	0.61	0.43
r8	1.6 - 10.0 <u>1.6 - 10.0</u>	-0.08 <u>-0.08</u>	-0.17 <u>-0.17</u>	0.38 <u>0.38</u>
	10.0 - 20.0 <u>10.0 - 20.0</u>	0.89 <u>0.89</u>	0.86 <u>0.86</u>	0.28 <u>0.28</u>
	1.6 - 20.0 <u>1.6 - 20.0</u>	0.09 <u>0.09</u>	0.12 <u>0.12</u>	0.29 <u>0.29</u>
r9	1.6 - 10.0 <u>1.6 - 10.0</u>	-0.15 <u>-0.15</u>	-0.18 <u>-0.18</u>	0.37 <u>0.37</u>
	10.0 - 20.0 <u>10.0 - 20.0</u>	1.19 <u>1.19</u>	1.26 <u>1.26</u>	0.33 <u>0.33</u>
	1.6 - 20.0 <u>1.6 - 20.0</u>	0.08 <u>0.08</u>	0.05 <u>0.05</u>	0.28 <u>0.28</u>
r10	1.6 - 10.0	-0.64	-0.49	0.27
	10.0 - 20.0	1.19	1.07	0.26
	1.6 - 20.0	-0.32	-0.18	0.21

320 the uncertainty of r8 using Tikhonov. The averaging kernels and cumulative sum of degrees of freedom (DOF) for N₂O are shown in Figure 10. The first DOF is captured from the surface to about 8 km, while the second DOF is between 8 to 17 km.

Table 8. CH₄ bias results among the different retrieval strategies in the lower troposphere using aircraft profiles as the reference. [The results in bold indicate the best performance.](#)

Retrieval #	Mean bias [%]	Median bias [%]	CI [%]
r1	1.90	1.92	0.32
r2	-0.76	-0.79	0.40
r3	2.89	2.88	0.34
r4	2.84	2.84	0.31
r5	-0.26	-0.10	0.36
r6	-0.47	-0.43	0.39
r7	1.08	1.24	0.64
r8 r8	0.39 0.39	0.30 0.30	0.42 0.42
r9 r9	0.35 0.35	0.41 0.41	0.41 0.41
r10 r10	-0.24 -0.24	-0.22 -0.22	0.37 0.37

Because N₂O appears to be well-mixed among all tropospheric altitudes, the results shown above are representative of the entire troposphere. Unfortunately, long-term stratospheric [AirCore](#) N₂O data is not yet available in the same fashion as CH₄ to evaluate the stratospheric component [using the same approach.](#)

Table 9. N₂O bias results among the different retrieval strategies in the lower troposphere using aircraft profiles as the reference. [The results in bold indicate the best performance.](#)

Retrieval #	Mean bias [%]	Median bias [%]	CI [%]
r1	1.58	1.62	0.28
r2	0.12 0.12	0.15 0.15	0.27 0.27
r3	1.42	1.50	0.27
r4	0.56	0.54	0.26
r5	1.27	1.35	0.28
r6	-0.02 -0.02	0.15 0.15	0.25 0.25
r7	0.36 0.36	0.36 0.36	0.22 0.22
r8	0.18 0.18	0.30 0.30	0.20 0.20

325

4.3 Recommended retrieval strategies and error budget

The final suggested retrievals for CH₄ and N₂O are presented in Table 10, based on both fitting results using various spectroscopy line parameters and especially by the assessment of vertical profiles and columns using independent AirCore and aircraft profiles. These retrievals correspond to r8 and r9 for CH₄ (Table 2), which are identical except that r8 employs Tikhonov regularization while r9 uses OE. For N₂O, retrievals follow the r8 strategy (~~r~~Table 4). Although Table 2 lists DLR water vapor

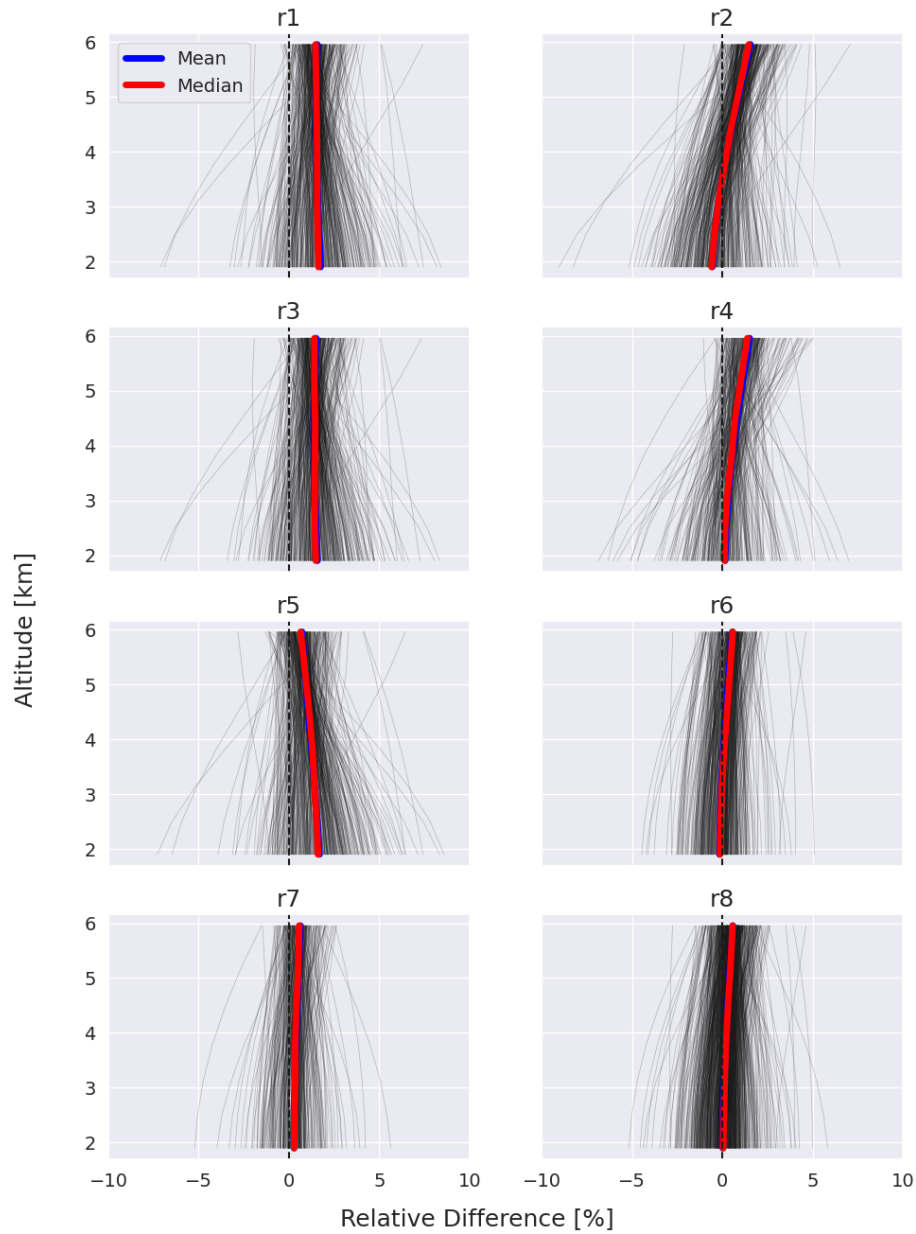


Figure 9. Relative difference between FTIR and NOAA aircraft N_2O profiles using several retrieval strategies.

330 for CH_4 retrievals, we believe that a standardized approach using ATM20 for all gases would provide satisfactory results, while also being more practical and easier to implement on a broader scale. SFIT4 provides uncertainty budgets that combine random and systematic sources following the formalism outlined in Rodgers and Connor (2003). The primary random and systematic error profiles for CH_4 and N_2O are illustrated in Figures 11 and 12, respectively. These vertical profile uncertainties are ex-

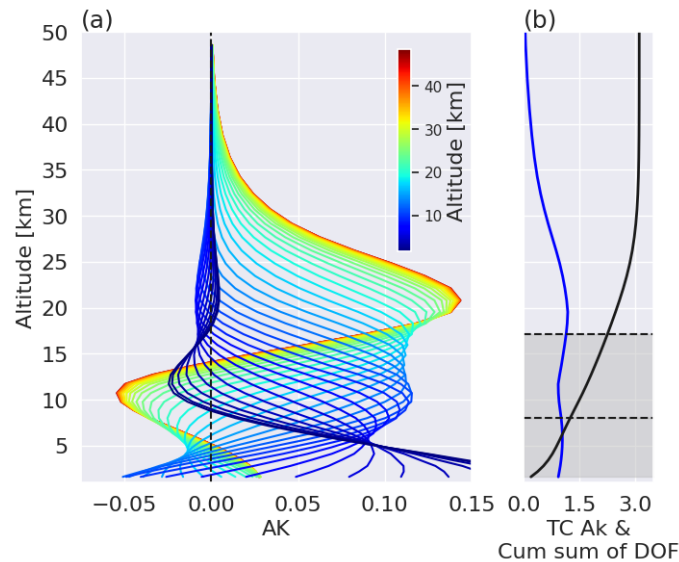


Figure 10. Left: N₂O FTIR mean row averaging kernels. Right: total column averaging kernel and cumulative sum of DOF on 20190131.171426 using 5 mw, Tikhonov regularization, and WACCM6 a priori. First DOF is between 1.6 - 8km, second DOF is between 8.0 - 17.12km

pressed as percentages relative to the mean mixing ratio, with the total respective errors represented by black dotted lines. The error calculation in SFIT4 is detailed in Ortega et al. (2019), with a specific analysis for water vapor at Boulder. The dominant random error in the troposphere for both gases stem from uncertainties in the temperature profile and measurement noise characterized by SNR in the spectral region of interest. Additional error parameters, such as interfering species, apodization function, and solar zenith angle, have a comparatively lesser impact. Major systematic error components arise from absorption line parameters, specifically line intensity (S), air-broadened half-width (γ), and temperature dependence of γ (n). The lower limit uncertainties reported in HIT20 are used for systematic error retrieval, with values of 5 %, 5 %, and 10 % for S , γ , and n respectively for CH₄, and 2 %, 10 %, and 5 % respectively for N₂O. It should be noted that these errors are expected to be significantly smaller for CH₄ due to the use of the empirically modified linelist ATM20. Smoothing error is treated separately and excluded from the total error analysis because it is often not well-known and is thus frequently simplified. Tables 11 and 12 provides a summary of mean values of summarize the mean random, systematic, and total uncertainties in the retrieval of both-CH₄ and N₂O, respectively. These values for three partial columns. These uncertainties are expressed as percentages in the relative to the mean weighted mixing ratio of the corresponding partial column as well as (using weighted mean errors) and the total column errors. Random error, associated with precision or statistical uncertainty, reflects the inherent variability in measurements and is crucial for meeting precision requirements in observations for CH₄, such as growth rate, seasonal cycle amplitudes, inter-hemispheric gradients, among others. This type of uncertainty is typically quantified as the standard deviation or standard error of a set of measurements. For example, (Sussmann et al., 2011) optimized precision to about 0.3 % using

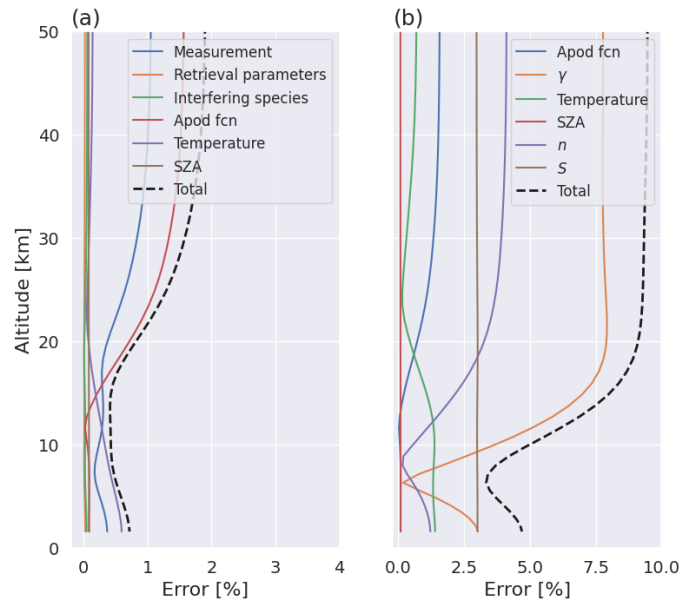


Figure 11. Mean vertical profiles of the primary random (a) and systematic (b) uncertainty components in the retrieval of CH₄ for the year 2021.

1-sigma standard deviation and diurnal variation in a 7-minute integration, aligning with the retrieval random error of 0.5 % shown in Table 11.

Table 10. Suggested CH₄ and N₂O final retrieval strategies based on fitting results and assessment with AirCore and aircraft profiles.

	CH ₄	N ₂ O
Micro-windows	1,2,3,4,5 (Table 1)	1,2,3,4 (Table 3)
Spectroscopy	ATM 2020 and DLR (H ₂ O)	HIT20
A priori	WACCM6 (all main species)	
Retrieval constraint	Tikhonov L1 or OEM	
IRWG Filter	Filter 3	Filter 3 and 4
Additional details	Regularization strength optimized for DOF $\approx$ 2.0-2.5	
	pre-retrieved H ₂ O, 257 cm OPD	
	Pressure and Temperature NCEP (at least daily)	

4.4 Seasonal variations and trends of CH₄: Boulder and Lauder NDACC sites

The retrieval strategies outlined in Table 10 have been applied to Boulder measurements spanning 2010–2023. To contrast and evaluate the CH₄ signal in the Southern Hemisphere, we extended the analysis to the NDACC site in Lauder, New Zealand;

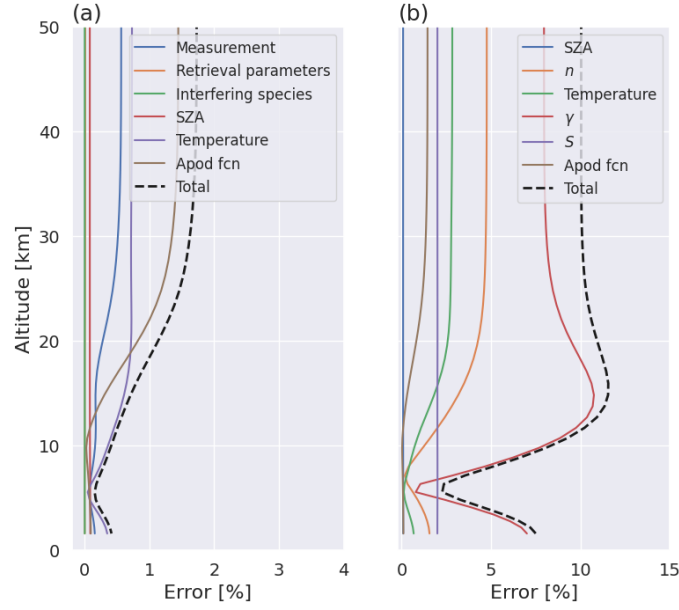


Figure 12. Same as 11 but for N_2O .

Table 11. Mean values of random, systematic, and total uncertainties in the CH_4 retrieval for the entire year of 2021 using the final suggested retrieval strategy. These values represent the weighted errors in the corresponding partial column, with the total column errors presented in the bottom row.

Altitude [km]	Random [%]	Systematic [%]	Total [%]
1.6-9.8	0.58 ± 0.06	4.07 ± 0.41	4.12 ± 0.41
10.7-19.6	0.46 ± 0.03	7.22 ± 0.47	7.24 ± 0.47
Total Column	0.44 ± 0.06	3.45 ± 0.05	3.48 ± 0.05

located at the National Institute of Water and Atmospheric Research (NIWA) station in Central Otago, at an altitude of 370 m.a.s.l. (Hausmann et al., 2016). FTIR observations at Lauder have been conducted since 1990, but to facilitate comparison with Boulder, we focus on data from 2010 onwards. Between 2010 and 2018, observations at Lauder were collected using a Bruker 120HR, while from 2018 onward, a Bruker 125HR has been used. To capture both the long-term trend and seasonal variations, we performed a bootstrap re-sampling analysis on the time series $f(t)$ using the following expression:

$$f(t) = a_0 + a_1(t - t_0) + \sum_{n=1}^N b_n \cos(n\pi t) + \sum_{n=1}^N c_n \sin(n\pi t)$$

Here, the first two terms represent the linear trend, where a_0 is the intercept, a_1 is the slope (indicating the long-term trend), and t_0 is the time of the initial observation. The third and fourth terms are a Fourier series that models the seasonal variability, with a second-order Fourier series ($N=2$) adequately capturing the seasonal cycle. The bootstrap analysis was

Table 12. Mean values of random, systematic, and total uncertainties in the N₂O retrieval for the entire year of 2021 using the final suggested retrieval strategy (r8). These values represent the weighted errors in the corresponding partial column, with the total column errors presented in the bottom row.

Altitude [km]	Random [%]	Systematic [%]	Total [%]
1.6-9.8	0.29 ± 0.09	5.21 ± 0.85	5.22 ± 0.85
10.7-19.6	0.66 ± 0.13	10.74 ± 0.75	10.76 ± 0.75
Total Column	0.16 ± 0.05	2.58 ± 0.09	2.59 ± 0.09

conducted 5×10^3 times, and the mean slope was used to quantify the annual rate of change, while the standard deviation provided an estimate of the uncertainty. An advantage of NDACC’s high-resolution FTIR measurements is that these retrievals yield 2 DOFs, allowing us to distinguish between the tropospheric and stratospheric CH₄ columns. Figure ?? displays the time-series of CH₄ tropospheric, stratospheric, and total columns at Boulder and Lauder, derived using the retrieval strategies recommended in this study. The altitude ranges for the tropospheric and stratospheric columns extend from the site altitude to the altitude where the DOF is closest to one, and from that altitude up to 25 km, respectively. For Boulder, this altitude is 9.3 km, and for Lauder, it is 7.6 km. While our goal is to differentiate the tropospheric and stratospheric signals, we opted not to exclude the tropopause in this analysis, though this might be an important factor for more detailed assessments in future studies. To enhance clarity, only data points within 5 % of the seasonal modulation’s amplitude are shown in gray. Monthly means are represented by circles with associated standard deviations, while solid lines represent the seasonal modulation and trend components derived from the bootstrap analysis. The annual rate of change for each column, relative to the mean of the full time series, is presented in Table ?. Another important assessment is the characterization of the seasonal variation. For a closer look at seasonal variation, we removed the linear trend using:-

$$f(t)_d = f(t) - (a_0 + a_1 (t - t_0))$$

The seasonal variation is then represented by the monthly means of this de-trended data, along with their uncertainties, in Figure ?? for both Boulder and Lauder over the period 2010–2023. This analysis is not intended to validate the trends or seasonal variations but to present the data, serving as a basis for future studies aiming to investigate these aspects in more detail. For additional insights into the comparison between various retrievals and AirCore and surface in situ trends and seasonal variation, see supplemental information. Overall, the linear trends and seasonal variations remain consistent across different retrieval strategies, with only the bias varying, as demonstrated in the comparisons with AirCore and aircraft measurements.

Time-series of CH₄ columns at Boulder (black) and Lauder (red) derived using the suggested retrieval strategy, separated into (a) tropospheric, (b) stratospheric, and (c) total columns. Gray dots represent individual data points as described in the main text, circles denote monthly mean values, and the solid lines show the combined seasonal modulation and trend component derived from the entire dataset.

hLinear trend analysis of CH₄ time series at Boulder and Lauder for tropospheric, stratospheric, and total columns as shown
390 in Figure ??.

Column	Boulder %/year	Lauder %/year
Tropospheric	-0.702 ± 0.007	-0.510 ± 0.005
Stratospheric	-0.390 ± 0.012	-0.715 ± 0.009
Total	-0.588 ± 0.005	-0.581 ± 0.004

Monthly mean values of CH₄ derived from the detrended analysis are presented for Boulder (black) and Lauder (red) across the troposphere (a), stratosphere (b), and total columns (c). Error bars represent the standard deviations of the multi-annual monthly means.

Over recent years, there has been noticeable variability in the retrieval strategies for CH₄ among different ground-based FTIR instruments within the IRWG of NDACC. Acknowledging advancements in spectroscopy and the need for retrieval consistency among the IRWG of NDACC, our study evaluates various retrieval strategies, encompassing not only CH₄ but also N₂O. Using observations from Boulder, Colorado, we initially assess fitting results across diverse spectroscopy databases, including the latest HITRAN databases, the ATM spectroscopy utilized by TCCON and IRWG, a combination of them, and a water vapor line list from DLR. We also evaluate previously identified suitable micro-windows. While some cases exhibit clear improvements in residuals for certain spectroscopy, in other instances, residual differences are less distinct, despite significant variations in profile retrievals. Thus, we emphasize the importance of the subsequent step: comparing independent long-term AirCore and aircraft insitu profile observations with the various ~~retrieval~~ FTIR retrieval strategies. By utilizing this dataset, we strengthen the evaluation of various retrieval strategies, ~~, i. e., 10 different retrieval strategies~~ for CH₄ ~~utilizing different~~ and N₂O, an approach rarely undertaken within the IRWG. In total, we assess 10 CH₄ retrieval strategies, varying micro-windows, spectroscopy, ~~regularization approach and regularization approaches~~ (Tikhonov or optimal estimation) and, as well as 8 retrievals for N₂O mainly retrievals, primarily focusing on spectroscopy, regularization ~~approach~~, and a priori profiles. Notably, both AirCore and aircraft profiles exhibit remarkable similarity in their relative differences compared to all FTIR retrievals, strengthening our

confidence in the assessment. Moreover, bias analyses across various atmospheric layers for both CH₄ and N₂O contribute to refining retrieval methodologies. Clear biases were observed when comparing with independent profiles, making the exclusion of certain retrievals straightforward. We propose optimized retrieval settings for CH₄ and N₂O, leveraging a first-order Tikhonov method and incorporating a priori profiles from the latest WACCM model simulations to enhance accuracy. While the HITRAN 2020 spectroscopic database is effective for N₂O, it exhibits suboptimal residuals for CH₄, indicating the need for further refinements. In contrast, ATM 2020 ~~provides yields~~ better results for CH₄, with ~~slight improvement observed when paired with the~~ a slight improvement when combined with the DLR water vapor line list ~~from DLR; however, However,~~ this enhancement may be site-dependent. Furthermore, profile comparisons reveal biases within different atmospheric layers. For the proposed retrievals, we obtain a mean bias of -0.08 ± 0.38 % and 0.89 ± 0.28 % for the tropospheric and stratospheric layers of CH₄ using AirCore, respectively, and 0.39 ± 0.42 % for aircraft comparisons in the troposphere. For N₂O, the bias

in the troposphere is approximately 0.18 ± 0.2 %. Furthermore, we provide uncertainty budgets, including random and systematic sources, to comprehensively understand error sources, guiding future refinement efforts. Random errors of about 0.5 % are found, mainly attributed to temperature profile uncertainties and measurement noise, dominate in the troposphere for both gases. Overall, our findings significantly contribute to advancing understanding of atmospheric composition and advocate for an effective harmonized approach among all IRWG/NDACC sites. ~~The accuracy of our retrievals allows us measurements of growth rates, seasonal cycle amplitudes, interhemispheric gradients, and supports satellite mission characterization.~~ Although Boulder benefits from unique long-term AirCore and aircraft profile observations, these capabilities are not widespread across IRWG/NDACC sites, underscoring the importance of deploying similar observations systems elsewhere in the NDACC network.

Code and data availability. The profile retrieval algorithm, SFIT4 (Hannigan et al., 2024), is freely available and can be obtained at <https://wiki.ucar.edu/display/sfit4/SFIT+Core+Code>. The Level 2 NOAA AirCore dataset (v20230831) (Baier et al., 2021) is publicly available via the NOAA data repository: <https://doi.org/10.15138/6AV0-MY81>. Atmospheric observations from the NOAA GML Carbon Cycle Aircraft Vertical Profile Network (Sweeney et al., 2015; McKain et al., 2023) can be obtained at <https://gml.noaa.gov/ccgg/aircraft/>. Additional data used in this study can be provided upon request from the corresponding author.

Author contributions. IO and JH conceptualized the project. IO carried out the data analysis, generated all figures and plots, and prepared the initial draft of the manuscript. BB, KM, and DS contributed the AirCore, aircraft, and Lauder FTIR datasets, respectively, and provided substantial input throughout the study. All co-authors contributed to discussions, offered feedback on the manuscript, and assisted in its preparation.

Competing interests. The authors declare no competing interests.

Acknowledgements. This material is based upon work supported by the National Center for Atmospheric Research (NCAR), which is a major facility sponsored by the National Science Foundation under Cooperative Agreement No. 1852977. This study has been supported under contract by the National Aeronautics and Space Administration (NASA) award No. NNX17AE38G. This study was supported by NOAA Climate Program Office's Atmospheric Chemistry, Carbon Cycle, and Climate program, grant number NA23OAR4310285. We thank the many people at NOAA GML that contributed to the AirCore and aircraft datasets used in this study, including Andy Crotwell, Molly Crotwell, Ed Dlugokencky, Jack Higgs, Xin Lan, Pat Lang, Thomas Legard, Monica Madronich, Eric Moglia, John Mund, Don Neff, Tim Newberger, Colm Sweeney, Pieter Tans, and Sonja Wolter.

References

- Alberti, C., Hase, F., Frey, M., Dubravica, D., Blumenstock, T., Dehn, A., Castracane, P., Surawicz, G., Harig, R., Baier, B. C., Bès, C., Bi, J., Boesch, H., Butz, A., Cai, Z., Chen, J., Crowell, S. M., Deutscher, N. M., Ene, D., Franklin, J. E., García, O., Griffith, D., Grouiez, B., Grutter, M., Hamdouni, A., Houweling, S., Humpage, N., Jacobs, N., Jeong, S., Joly, L., Jones, N. B., Jouglet, D., Kivi, R., Kleinschek, R., Lopez, M., Medeiros, D. J., Morino, I., Mostafavipak, N., Müller, A., Ohyama, H., Palmer, P. I., Pathakoti, M., Pollard, D. F., Raffalski, U., Ramonet, M., Ramsay, R., Sha, M. K., Shiomi, K., Simpson, W., Stremme, W., Sun, Y., Tanimoto, H., Té, Y., Tsidu, G. M., Velazco, V. A., Vogel, F., Watanabe, M., Wei, C., Wunch, D., Yamasoe, M., Zhang, L., and Orphal, J.: Improved calibration procedures for the EM27/SUN spectrometers of the COllaborative Carbon Column Observing Network (COCCON), *Atmospheric Measurement Techniques*, 15, 2433–2463, <https://doi.org/10.5194/amt-15-2433-2022>, publisher: Copernicus GmbH, 2022.
- 455 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, *Atmospheric Chemistry and Physics*, 17, 2255–2277, <https://doi.org/10.5194/acp-17-2255-2017>, publisher: Copernicus GmbH, 2017.
- 460 Baier, B., Sweeney, C., Newberger, T., Higgs, J., and Wolter, S.: NOAA AirCore atmospheric sampling system profiles (Version 20230831), <https://doi.org/https://doi.org/10.15138/6AV0-MY81>, 2021.
- Canadell, J. G., Monteiro, P. M. S., Costa, M. H., Cotrim da Cunha, L., Cox, P. M., Eliseev, A. V., Henson, S., Ishii, M., Jaccard, S., Koven, C., Lohila, A., Patra, P. K., Piao, S., Rogelj, J., Syampungani, S., Zaehle, S., and Zickfeld, K.: Global Carbon and other Biogeochemical Cycles and Feedbacks, in: *Climate Change 2021: The Physical Science Basis. Contribution of Working Group I to the Sixth Assessment Report of the Intergovernmental Panel on Climate Change*, edited by Masson-Delmotte, V., Zhai, P., Pirani, A., Connors, S. L., Péan, C., Berger, S., Caud, N., Chen, Y., Goldfarb, L., Gomis, M. I., Huang, M., Leitzell, K., Lonnoy, E., Matthews, J. B. R., Maycock, T. K., Waterfield, T., Yelekçi, O., Yu, R., and Zhou, B., pp. 673–816, Cambridge University Press, Cambridge, United Kingdom and New York, NY, USA, <https://doi.org/10.1017/9781009157896.007>, 2021.
- 465 Chesnokova, T. Y., Makarova, M. V., Chentsov, A. V., Kostsov, V. S., Poberovskii, A. V., Zakharov, V. I., and Rokotyan, N. V.: Estimation of the impact of differences in the CH₄ absorption line parameters on the accuracy of methane atmospheric total column retrievals from ground-based FTIR spectra, *Journal of Quantitative Spectroscopy and Radiative Transfer*, 254, 107187, <https://doi.org/10.1016/j.jqsrt.2020.107187>, 2020.
- De Mazière, M., Thompson, A. M., Kurylo, M. J., Wild, J. D., Bernhard, G., Blumenstock, T., Braathen, G. O., Hannigan, J. W., Lambert, J.-C., Leblanc, T., McGee, T. J., Nedoluha, G., Petropavlovskikh, I., Seckmeyer, G., Simon, P. C., Steinbrecht, W., and Strahan, S. E.: The Network for the Detection of Atmospheric Composition Change (NDACC): history, status and perspectives, *Atmospheric Chemistry and Physics*, 18, 4935–4964, <https://doi.org/10.5194/acp-18-4935-2018>, publisher: Copernicus GmbH, 2018.
- 475 Garcia, R. R., Marsh, D. R., Kinnison, D. E., Boville, B. A., and Sassi, F.: Simulation of secular trends in the middle atmosphere, 1950–2003, *Journal of Geophysical Research: Atmospheres*, 112, <https://doi.org/10.1029/2006JD007485>, <https://agupubs.onlinelibrary.wiley.com/doi/pdf/10.1029/2006JD007485>, 2007.
- 480 Gettelman, A., Mills, M. J., Kinnison, D. E., Garcia, R. R., Smith, A. K., Marsh, D. R., Tilmes, S., Vitt, F., Bardeen, C. G., McInerny, J., Liu, H.-L., Solomon, S. C., Polvani, L. M., Emmons, L. K., Lamarque, J.-F., Richter, J. H., Glanville, A. S., Bacmeister, J. T., Phillips, A. S., Neale, R. B., Simpson, I. R., DuVivier, A. K., Hodzic, A., and Randel, W. J.: The Whole Atmosphere Community Climate Model Ver-

- sion 6 (WACCM6), *Journal of Geophysical Research: Atmospheres*, 124, 12 380–12 403, <https://doi.org/10.1029/2019JD030943>, _eprint: <https://onlinelibrary.wiley.com/doi/pdf/10.1029/2019JD030943>, 2019.
- 485 Gordon, I. E., Rothman, L. S., Hargreaves, R. J., Hashemi, R., Karlovets, E. V., Skinner, F. M., Conway, E. K., Hill, C., Kochanov, R. V., Tan, Y., Wcisło, P., Finenko, A. A., Nelson, K., Bernath, P. F., Birk, M., Boudon, V., Campargue, A., Chance, K. V., Coustenis, A., Drouin, B. J., Flaud, J. Gamache, R. R., Hodges, J. T., Jacquemart, D., Mlawer, E. J., Nikitin, A. V., Perevalov, V. I., Rotger, M., Tennyson, J., Toon, G. C., Tran, H., Tyuterev, V. G., Adkins, E. M., Baker, A., Barbe, A., Canè, E., Császár, A. G., Dudaryonok, A., Egorov, O., Fleisher, A. J., Fleurbaey, H., Foltynowicz, A., Furtenbacher, T., Harrison, J. J., Hartmann, J. Horneman, V. Huang, X., Karman, T., Karns, J.,
- 490 Kassi, S., Kleiner, I., Kofman, V., Kwabia-Tchana, F., Lavrentieva, N. N., Lee, T. J., Long, D. A., Lukashevskaya, A. A., Lyulin, O. M., Makhnev, V. Y., Matt, W., Massie, S. T., Melosso, M., Mikhailenko, S. N., Mondelain, D., Müller, H. S. P., Naumenko, O. V., Perrin, A., Polyansky, O. L., Raddaoui, E., Raston, P. L., Reed, Z. D., Rey, M., Richard, C., Tóbiás, R., Sadiek, I., Schwenke, D. W., Starikova, E., Sung, K., Tamassia, F., Tashkun, S. A., Vander Auwera, J., Vasilenko, I. A., Vigasin, A. A., Villanueva, G. L., Vispoel, B., Wagner, G., Yachmenev, A., and Yurchenko, S. N.: The HITRAN2020 molecular spectroscopic database, *Journal of Quantitative Spectroscopy and*
- 495 *Radiative Transfer*, 277, 107 949, <https://doi.org/10.1016/j.jqsrt.2021.107949>, 2022.
- Hannigan, J., Palm, M., Jones, N., Ortega, I., Langerock, B., Mahieu, E., Zhou, M., and Smale, D.: SFIT4 Line-by-line nonlinear spectral fitting software: version 1.0.21, <https://doi.org/10.18758/CLYG3EUO>, 2024.
- Hannigan, J. W., Coffey, M. T., and Goldman, A.: Semiautonomous FTS observation system for remote sensing of stratospheric and tropospheric gases, *Journal of Atmospheric and Oceanic Technology*, 26, 1814–1828, <https://doi.org/10.1175/2009JTECHA1230.1>, 2009.
- 500 Hase, F., Hannigan, J. W., Coffey, M. T., Goldman, A., Höpfner, M., Jones, N. B., Rinsland, C. P., and Wood, S. W.: Intercomparison of retrieval codes used for the analysis of high-resolution, ground-based FTIR measurements, *Journal of Quantitative Spectroscopy and Radiative Transfer*, 87, 25–52, <https://doi.org/10.1016/j.jqsrt.2003.12.008>, 2004.
- Hase, F., Frey, M., Kiel, M., Blumenstock, T., Harig, R., Keens, A., and Orphal, J.: Addition of a channel for XCO observations to a portable FTIR spectrometer for greenhouse gas measurements, *Atmospheric Measurement Techniques*, 9, 2303–2313, [https://doi.org/10.5194/amt-](https://doi.org/10.5194/amt-9-2303-2016)
- 505 [9-2303-2016](https://doi.org/10.5194/amt-9-2303-2016), publisher: Copernicus GmbH, 2016.
- 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, *Atmospheric Chemistry and Physics*, 16, 3227–3244, <https://doi.org/10.5194/acp-16-3227-2016>, 2016.
- Karion, A., Sweeney, C., Tans, P., and Newberger, T.: AirCore: An Innovative Atmospheric Sampling System, *Journal of Atmospheric and*
- 510 *Oceanic Technology*, 27, 1839–1853, <https://doi.org/10.1175/2010JTECHA1448.1>, publisher: American Meteorological Society Section: *Journal of Atmospheric and Oceanic Technology*, 2010.
- McKain, K., Sweeney, C., Baier, B., Crotwell, A., Crotwell, M., Handley, P., Higgs, J., Legard, T., Madronich, M., Miller, J. B., Moglia, E., Mund, J., Newberger, T., Wolter, S., and Laboratory, . N. G. M.: NOAA Global Greenhouse Gas Reference Network Flask-Air PFP Sample Measurements of CO₂, CH₄, CO, N₂O, H₂, SF₆ and isotopic ratios collected from aircraft vertical profiles [Data set], <https://doi.org/10.15138/39HR-9N34>, 2023.
- 515 Olsen, K. S., Strong, K., Walker, K. A., Boone, C. D., Raspollini, P., Plieninger, J., Bader, W., Conway, S., Grutter, M., Hannigan, J. W., Hase, F., Jones, N., de Mazière, M., Notholt, J., Schneider, M., Smale, D., Sussmann, R., and Saitoh, N.: Comparison of the GOSAT TANSO-FTS TIR CH₄ volume mixing ratio vertical profiles with those measured by ACE-FTS, ESA MIPAS, IMK-IAA MIPAS, and 16 NDACC stations, *Atmospheric Measurement Techniques*, 10, 3697–3718, <https://doi.org/10.5194/amt-10-3697-2017>, publisher: Copernicus GmbH, 2017.
- 520

- Ortega, I., Buchholz, R., Hall, E. G., Hurst, D. F., Jordan, A. F., and Hannigan, J. W.: Tropospheric water vapor profiles obtained with FTIR: Comparison with balloon-borne frost point hygrometers and influence on trace gas retrievals, *Atmospheric Measurement Techniques*, 12, 873–890, <https://doi.org/10.5194/amt-12-873-2019>, 2019.
- Ortega, I., Hannigan, J. W., Buchholz, R. R., and Pfister, G.: Long-term variability and source signature of gases emitted from oil & natural gas and cattle feedlot operations in the Colorado front range, *Atmospheric Environment*, 263, 118663, <https://doi.org/10.1016/j.atmosenv.2021.118663>, 2021.
- Pougatchev, N. S., Connor, B., and Rinsland, C. P.: Infrared measurements of the ozone vertical distribution above Kitt Peak, *Journal of Geophysical Research*, 100, 16 689–16 697, <https://doi.org/10.1029/95JD01296>, 1995.
- Prather, M. J., Hsu, J., DeLuca, N. M., Jackman, C. H., Oman, L. D., Douglass, A. R., Fleming, E. L., Strahan, S. E., Steenrod, S. D., Søvde, O. A., Isaksen, I. S. A., Froidevaux, L., and Funke, B.: Measuring and modeling the lifetime of nitrous oxide including its variability, *Journal of Geophysical Research: Atmospheres*, 120, 5693–5705, <https://doi.org/https://doi-org.cuucar.idm.oclc.org/10.1002/2015JD023267>, 2015.
- Rinsland, C. P., Jones, N. B., Connor, B. J., Logan, J. A., Pougatchev, N. S., Goldman, A., Murcray, F. J., Stephen, T. M., Pine, A. S., Zander, R., Mahieu, E., and Demoulin, P.: Northern and southern hemisphere ground-based infrared spectroscopic measurements of tropospheric carbon monoxide and ethane, *Journal of Geophysical Research: Atmospheres*, 103, 28 197–28 217, <https://doi.org/10.1029/98JD02515>, 1998.
- Rodgers, C. and Connor, B.: Intercomparison of remote sounding instruments, *Journal of Geophysical Research (Atmospheres)*, 108, <https://doi.org/10.1029/2002JD002299>, 2003.
- Rothman, L. S., Gordon, I. E., Barbe, A., Benner, D. C., Bernath, P. F., Birk, M., Boudon, V., Brown, L. R., Campargue, A., Champion, J. P., Chance, K., Coudert, L. H., Dana, V., Devi, V. M., Fally, S., Flaud, J. M., Gamache, R. R., Goldman, A., Jacquemart, D., Kleiner, I., Lacome, N., Lafferty, W. J., Mandin, J. Y., Massie, S. T., Mikhailenko, S. N., Miller, C. E., Moazzen-Ahmadi, N., Naumenko, O. V., Nikitin, A. V., Orphal, J., Perevalov, V. I., Perrin, A., Predoi-Cross, A., Rinsland, C. P., Rotger, M., Šimečková, M., Smith, M. A. H., Sung, K., Tashkun, S. A., Tennyson, J., Toth, R. A., Vandaele, A. C., and Vander Auwera, J.: The *HITRAN* 2008 molecular spectroscopic database, *Journal of Quantitative Spectroscopy and Radiative Transfer*, 110, 533–572, <https://doi.org/10.1016/j.jqsrt.2009.02.013>, 2009.
- Sepúlveda, E., Schneider, M., Hase, F., Barthlott, S., Dubravica, D., García, O. E., Gomez-Pelaez, A., González, Y., Guerra, J. C., Gisi, M., Kohlhepp, R., Dohe, S., Blumenstock, T., Strong, K., Weaver, D., Palm, M., Sadeghi, A., Deutscher, N. M., Warneke, T., Notholt, J., Jones, N., Griffith, D. W. T., Smale, D., Brailsford, G. W., Robinson, J., Meinhardt, F., Steinbacher, M., Aalto, T., and Worthy, D.: Tropospheric CH₄ signals as observed by NDACC FTIR at globally distributed sites and comparison to GAW surface in situ measurements, *Atmospheric Measurement Techniques*, 7, 2337–2360, <https://doi.org/10.5194/amt-7-2337-2014>, publisher: Copernicus GmbH, 2014.
- Steck, T.: *Methods for determining regularization*, 2002.
- Sussmann, R., Forster, F., Rettinger, M., and Jones, N.: Strategy for high-accuracy-and-precision retrieval of atmospheric methane from the mid-infrared FTIR network, *Atmospheric Measurement Techniques*, 4, 1943–1964, <https://doi.org/10.5194/amt-4-1943-2011>, publisher: Copernicus GmbH, 2011.
- Sweeney, C., Karion, A., Wolter, S., Newberger, T., Guenther, D., Higgs, J. A., Andrews, A. E., Lang, P. M., Neff, D., Dlugokencky, E., Miller, J. B., Montzka, S. A., Miller, B. R., Masarie, K. A., Biraud, S. C., Novelli, P. C., Crotnell, M., Crotnell, A. M., Thoning, K., and Tans, P. P.: Seasonal climatology of CO₂ across North America from aircraft measurements in the NOAA/ESRL Global Greenhouse Gas Reference Network, *Journal of Geophysical Research: Atmospheres*, 120, 5155–5190, <https://doi.org/https://doi.org/10.1002/2014JD022591>, 2015.

Thompson, C. R., Wofsy, S. C., Prather, M. J., Newman, P. A., Hanisco, T. F., Ryerson, T. B., Fahey, D. W., Apel, E. C., Brock, C. A., Brune, W. H., Froyd, K., Katich, J. M., Nicely, J. M., Peischl, J., Ray, E., Veres, P. R., Wang, S., Allen, H. M., Asher, E., Bian, H., Blake, D., Bourgeois, I., Budney, J., Bui, T. P., Butler, A., Campuzano-Jost, P., Chang, C., Chin, M., Commane, R., Correa, G., Crounse, J. D., Daube, B., Dibb, J. E., DiGangi, J. P., Diskin, G. S., Dollner, M., Elkins, J. W., Fiore, A. M., Flynn, C. M., Guo, H., Hall, S. R., Hannun, R. A., Hills, A., Hintsa, E. J., Hodzic, A., Hornbrook, R. S., Huey, L. G., Jimenez, J. L., Keeling, R. F., Kim, M. J., Kupc, A., Lacey, F., Lait, L. R., Lamarque, J.-F., Liu, J., McKain, K., Meinardi, S., Miller, D. O., Montzka, S. A., Moore, F. L., Morgan, E. J., Murphy, D. M., Murray, L. T., Nault, B. A., Neuman, J. A., Nguyen, L., Gonzalez, Y., Rollins, A., Rosenlof, K., Sargent, M., Schill, G., Schwarz, J. P., Clair, J. M. S., Steenrod, S. D., Stephens, B. B., Strahan, S. E., Strode, S. A., Sweeney, C., Thames, A. B., Ullmann, K., Wagner, N., Weber, R., Weinzierl, B., Wennberg, P. O., Williamson, C. J., Wolfe, G. M., and Zeng, L.: The NASA Atmospheric Tomography (ATom) Mission: Imaging the Chemistry of the Global Atmosphere, *Bulletin of the American Meteorological Society*, 103, E761 – E790, <https://doi.org/10.1175/BAMS-D-20-0315.1>, 2022.

Tian, H., Xu, R., Canadell, J. G., Thompson, R. L., Winiwarter, W., Suntharalingam, P., Davidson, E. A., Ciais, P., Jackson, R. B., Janssens-Maenhout, G., Prather, M. J., Regnier, P., Pan, N., Pan, S., Peters, G. P., Shi, H., Tubiello, F. N., Zaehle, S., Zhou, F., Arneth, A., Battaglia, G., Berthet, S., Bopp, L., Bouwman, A. F., Buitenhuis, E. T., Chang, J., Chipperfield, M. P., Dangal, S. R. S., Dlugokencky, E., Elkins, J. W., Eyre, B. D., Fu, B., Hall, B., Ito, A., Joos, F., Krummel, P. B., Landolfi, A., Laruelle, G. G., Lauerwald, R., Li, W., Lienert, S., Maavara, T., MacLeod, M., Millet, D. B., Olin, S., Patra, P. K., Prinn, R. G., Raymond, P. A., Ruiz, D. J., van der Werf, G. R., Vuichard, N., Wang, J., Weiss, R. F., Wells, K. C., Wilson, C., Yang, J., and Yao, Y.: A comprehensive quantification of global nitrous oxide sources and sinks, *Nature*, 586, 248–256, <https://doi.org/10.1038/s41586-020-2780-0>, 2020.

Toon, G. C.: Atmospheric Line List for the 2014 TCCON Data Release, <https://doi.org/https://doi.org/10.14291/TCCON.GGG2014.ATM.R0/1221656>, 2015.

Toon, G. C., Blavier, J.-F., Sung, K., Rothman, L. S., and E. Gordon, I.: HITRAN spectroscopy evaluation using solar occultation FTIR spectra, *Journal of Quantitative Spectroscopy and Radiative Transfer*, 182, 324–336, <https://doi.org/https://doi.org/10.1016/j.jqsrt.2016.05.021>, 2016.

Wild, J. D., Gelman, M. E., Miller, A. J., Chanin, M. L., Hauchecorne, A., Keckhut, P., Farley, R., Dao, P. D., Meriwether, J. W., Gobbi, G. P., Congeduti, F., Adriani, A., McDermid, I. S., McGee, T. J., and Fishbein, E. F.: Comparison of stratospheric temperatures from several lidars, using National Meteorological Center and microwave limb sounder data as transfer references, *Journal of Geophysical Research: Atmospheres*, 100, 11 105–11 111, <https://doi.org/10.1029/95JD00631>, _eprint: <https://onlinelibrary.wiley.com/doi/pdf/10.1029/95JD00631>, 1995.

Wofsy, S. C.: HIAPER Pole-to-Pole Observations (HIPPO): fine-grained, global-scale measurements of climatically important atmospheric gases and aerosols, *Philosophical Transactions of the Royal Society A: Mathematical, Physical and Engineering Sciences*, 369, 2073–2086, <https://doi.org/10.1098/rsta.2010.0313>, 2011.

Wunch, D., Toon, G. C., Blavier, J.-F. L., Washenfelder, R. A., Notholt, J., Connor, B. J., Griffith, D. W. T., Sherlock, V., and Wennberg, P. O.: The Total Carbon Column Observing Network, *Philosophical Transactions of the Royal Society A: Mathematical, Physical and Engineering Sciences*, 369, 2087–2112, <https://doi.org/10.1098/rsta.2010.0240>, publisher: Royal Society, 2011.

Zhou, M., Langerock, B., Vigouroux, C., Sha, M. K., Ramonet, M., Delmotte, M., Mahieu, E., Bader, W., Hermans, C., Kumps, N., Metzger, J.-M., Dufлот, 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, *Atmospheric Chemistry and Physics*, 18, 13 881–13 901, <https://doi.org/10.5194/acp-18-13881-2018>, publisher: Copernicus GmbH, 2018.

Zhou, M., Langerock, B., Wells, K. C., Millet, D. B., Vigouroux, C., Sha, M. K., Hermans, C., Metzger, J.-M., Kivi, R., Heikkinen, P., Smale, D., Pollard, D. F., Jones, N., Deutscher, N. M., Blumenstock, T., Schneider, M., Palm, M., Notholt, J., Hannigan, J. W., and De Mazière, M.: An intercomparison of total column-averaged nitrous oxide between ground-based FTIR TCCON and NDACC measurements at seven sites and comparisons with the GEOS-Chem model, *Atmospheric Measurement Techniques*, 12, 1393–1408, <https://doi.org/10.5194/amt-12-1393-2019>, publisher: Copernicus GmbH, 2019.