

REVIEW #1

More information is needed on SMR in Section 2, for instance Are the 2 side bands folded over on each other, or are they separate? (Could be part of more detailed explanation of Fig. 3). Section 2.1 has been updated with the following: “The output signal of the mixer is a function of the sum and difference of the frequencies of the input signals, therefore consisting of two sidebands. SMR is however run in single sideband mode so, of the two sidebands, only the one containing the signal of interest should be transmitted and the other one suppressed. This is achieved, before mixing, by accordingly setting the length of the arms of the Martin-Puplett interferometer.”

Is the scan continuous, or a step scan? It is a continuous scan with a constant speed of 0.75 km/s. This information has been added to Section 2.1.

Section 3.1: The method looks conceptually straightforward. Is there a reason the this has not been done before? The reason was a lack of personnel that would take care of this.

It looks as though the method described will work from 40 to 70 km, but what is done above that? Once the line has been identified via the spectra in the 40 – 60 km range, all the spectra in the scan are moved in bulk to the theoretical position. This is done by calculating the mean spectrum of all spectra in the scan, then the center frequency of the line is identified and the difference to the theoretical center is calculated. This same value is applied as a frequency correction for all spectra in the scan. After that a second correction, different for each spectrum, is applied, as described in Section 3.1 of the paper. We have clarified that point in Section 3.1.

Is the monthly a priori also divided into latitude bins? Yes, it is divided into 10° latitude bins. This information has been added to the text.

What is the “measurement response”? Is this the same as the Degrees of Freedom of the Signal (DFS)? The following was added to the text: “, where the measurement response is a measure of the degree to which the result may be contaminated by the a priori. It is defined as the sum over the row of the averaging kernel matrix (Rodgers, 2000)”. Also, the measurement response profile was added to Fig.5.

Section 3.2: Does the retrieval depend on temperature? If so, what temperature is used? What are its Biases? Yes, the retrievals depend on temperature. ERA-interim reanalysis data (Dee et al., 2001) is used up to 60 km and the Mass Spectrometer Incoherent Scatter model (version NRLMSISE-00; Picone et al., 2002) is used from 70 km upwards. Between 60 – 70 km, a spline interpolation of the two is applied. Information added to the text.

P. 7, Line 12 (regarding Fig. 5)- isn't it more that the concentrations above 90 km are influenced by the signals there? Yes, concentrations between 90 – 100 km are influenced by the signals there. In the text it is written that they are *also* influenced by higher altitudes measurements, as can be seen by the shape of the averaging kernels.

P. 9, l. 25: Please relate the vertical motions associated with the SAO with these signals. We have added the following explanation to the manuscript: “Although the SAO-induced variations in zonal mean vertical wind are too small to be measured directly, the accumulated effects can be seen in the distribution of the long-lived constituents like CO (Hamilton, 2015). Sinking of air, or at least anomalously weak rising motion, leads to the observed increase in the CO mixing ratios around the equinoxes.”

l. 28: clarify which is “this latitude band”. This has been changed to “high latitudes”.

P. 10, l. 1: Isn’t it more accurate to say SSW’s occur almost entirely in the N.H. We have changed “occur mainly in the N.H.” to “occur almost entirely in the N.H.”

P. 12: l.2: What was the criterion for sufficient coincidences? Coincidences are considered sufficient if they reach at least the number of 10. This information was added to the text.

l. 11: If this is absolute difference, please denote with abs. value brackets. Why use absolute values- aren’t you interested in whether the difference is positive or negative? If not, why? Absolute difference just means the subtraction $x_{\text{SMR}} - x_{\text{comp}}$ (without any normalization, as it is done for the relative difference), which of course can be negative, as it can be seen in the figures showing the comparisons. Absolute difference doesn’t mean “the absolute value of the difference” as the reviewer suggests.

ll. 17-18: Wording is unclear- is the average difference being calculated, or the median, or some combination? Is $\Delta(z)$ the mean of $|x_{\text{SMR}} - x_{\text{comp}}|(z)$? Where does the median come in? We changed the text to “the median difference $\Delta(z)$... is calculated.”

The SEM expression is the standard for calculating the standard deviation of the mean. True, it is the same as for the standard deviation of the mean, only that here $\Delta(z)$ is the median, not the mean.

P. 13, l. 12: Why were the 2 different periods with different MIPAS spectral resolution lumped together? Does the MIPAS spectral resolution not make a difference? The reviewer is right. The comparison has been re-done to consider the two periods separately and the paper has been updated accordingly. Figures 9 and 14 have been updated to show comparison with FR-NOM and OR-NOM separately. The following has been added to the text: “Regarding FR-NOM mode, with the above mentioned coincidence criteria we found 6088 coincident measurements with SMR over the period July 2002 - March 2004. The profiles and differences between 50 and 70 km altitude, averaged over the whole time period and over the whole globe, are shown in Figure 9a. The median relative difference between SMR and MIPAS has a value of +8% at 50 km which decreases to reach -15% at 60 km, and eventually remains constant between 60 - 70 km altitude. No bias characterising a particular latitude or season over time has been identified for this specific comparison, since FR-NOM mode has been operational for only a short period.

Comparing with OR-NOM mode, 88902 coincident measurements have been found over the period January 2005 - April 2012. Figure 9b shows that the median relative difference between SMR and MIPAS remains positive and almost constant with altitude, staying below +15%. The main contribution to this difference comes from latitudes between -25° and +50°. In particular during northern summer, around +25°, where it reaches values above +40%

between 65 and 70 km. Everywhere else relative difference values remain within $\pm 20\%$ (see Figure A1).”

The difference plots in Fig. 9a stop at ~ 70 km. What is plotted in the right panel above that—is it all a priori? Nothing is plotted above 70 km in the right panel, as well as in the center panel. In fact, differences between the two instruments are plotted in these panels and, since MIPAS nominal data extend only up to 70 km, there can be no difference shown above that altitude. In case the reviewer meant the left panel, note that what is plotted above 70 km is SMR CO concentration which is retrieved up to 100 km. So no, it is not all a priori.

P. 14, Fig. 9: What is the explanation, or theory for, the negative peak in the comparisons ~ 85 km? Since this occurs for all global comparisons, it would seem that it is a feature of the SMR measurements and/or analysis. On the same subject, there are very large and fluctuating differences as functions of season and latitude. Do the authors have any explanation for these variations? As explained more extensively in the dedicated points, SMR-MLS differences are consistent with a known bias of MLS (positive systematic error of 20–50%). Regarding concentration differences with other instruments we have checked that they are not due to differences in vertical resolution. This has been done by smoothing SMR profiles with a Gaussian filter to account for differences in vertical resolutions; however, this smoothing didn’t lead to significant changes in the profile. Also, we isolated MIPAS daytime profiles (which have a vertical resolution comparable to SMR) but the same relative differences with SMR were observed as in the globally averaged profiles. Moreover, it is unlikely that SMR-MIPAS relative differences are due to a misestimation of non-LTE effects from MIPAS, since systematic errors due to non-LTE were estimated to be less than 5% (in Funke et al., 2007, doi:10.1029/2006JD007933). This being said, the causes for the observed concentration differences, both globally and for different latitudes and seasons, are unknown. This information has been added to the Section “Conclusion” of the paper.

Note that the large number of comparisons make the SEM’s very small and perhaps not useful. It is true that the SEM is sometimes not visible, but the fact that it is very small is a useful information. Moreover, there are some cases where it is not negligible and, for the sake of consistency, we should indicate it in all comparisons.

P. 16, 13 MLS: Since MLS is another microwave instrument, it is surprising that its data are more different than that from MIPAS or ACE-FTS. The authors should comment on why this might be the case. Is it instrumental, or arising from the data analysis? It is a known bias of MLS, already referred to in the text in Section 5.3. MLS CO, as reported e.g. in Errera et al.(2019) and Livesey et al.(2018), is known to be affected by a positive systematic error of 20–50% throughout the mesosphere. This explains the observed SMR-MLS differences. Moreover, although SMR and MLS are both microwave instruments, they are not using the same spectral line, so there may be a variety of spectroscopic errors which would affect one instrument and not the other. However, CO is a species that radio astronomers have been studying for many decades; we rather suspect that its spectroscopy is better known than that of most other molecules. When MLS validation has been performed in Pumphrey et al. (2007), CO spectroscopy revealed itself to cause a 10% effect at worst; less serious than a variety of other systematic errors.

P. 17, ll. 4-5: Again, what is measurement response? Why a criterion of 0.75? For definition see point above. The value of 0.75 has been used for a long time. The reason is that, using a higher value as a threshold would result in the filtering out of a great part of the data.

Were OSO data only available for 2002-2007? The description seems to indicate an instrument improvement, especially after 2014. Most of the OSO data after 2014 has not been processed yet and the corresponding L2 data is therefore not available. The instrument improvement has been described just to shortly point out that OSO does not operate with a single sideband mixer anymore.

P. 19, Fig. 14: The difference of the comparisons between MLS and the other instruments is striking, as noted above. Also, the consistent decrease from higher to lower than the comparisons with altitude. As mentioned above, MLS CO is known to be characterized by a positive systematic error of 20–50% throughout the mesosphere.

Appendix: Latitudinal and seasonal differences in the biases at high altitudes- who is right? The causes for the observed concentration differences are unknown.

Looking at Figs. A2 and A3, it strikes this reviewer that the strong differences with MIPAS occur at the summer mesopause, the coldest region of the atmosphere. However, this difference does not show up in the ACE-FTS comparisons. This suggests that the MIPAS CO mixing effects might be too high, which could result from their temperatures being too low. At this level MIPAS temperatures must take non-LTE effects into account. These are notoriously difficult, and a negative error could lead to CO amounts too high.

1) non-LTE modelling of MIPAS CO has been extensively validated in Funke et al., 2007 (doi:10.1029/2006JD007933) and systematic errors due to non-LTE were estimated to be less than 5%. This information has been added to Section 5.1.

2) MIPAS CO (at least during daytime and hence polar summer) is rather independent on kinetic temperature, since CO daytime vibrational temperatures are dictated by solar excitation. A mapping of T errors is thus very unlikely, except for indirect effects via hydrostatics.

3) Further, the MIPAS kinetic temperature product used in the CO retrievals has been extensively validated in Garcia-Comas et al, 2014 (<https://www.atmos-meas-tech.net/7/3633/2014/>) and differences to correlative measurements were found to be less than 4K in the polar summer mesopause region.

4) the polar summer mesosphere, where the most striking differences occur, is characterised by very large vertical gradients in the CO distribution. In consequence, differences in vertical resolution between both instruments are likely important, there. While the vertical resolution of mesospheric MIPAS daytime CO is around 5 km (similar to that of SMR), nighttime vertical resolution is worse (>10 km) because of the lower sensitivity caused by smaller nighttime non-LTE emissions (no solar excitation). Therefore, it could be possible, that the SMR-MIPAS comparison is affected by the lower MIPAS nighttime resolution. This has been checked by restricting the comparison to MIPAS daytime profiles, but the resulting difference profiles are very similar to the ones considering data altogether. This suggests that differences in vertical resolution are not the cause of the observed biases. This information has been added to Section 5.1.

Technical Comments:

Page 1, Line 7: Much of the level 1. . . (The not needed) [OK](#)

l. 8: front end [OK](#)

l. 17: comparison instruments [OK](#)

P2, l. 19, 20- Suggest wording- “. . .obtained overall agreement over a two order of magnitude agreement” [Not applied. We think the current wording is appropriate.](#)

l. 22: front end [OK](#)

P3, l. 19 downward) scans [Not applied. We would like to avoid using the word “scan” in the definition of scan.](#)

P4, l. 7 front end [OK](#)

P. 10, Fig. 7- tick marks on abscissa can't be seen, also Fig. 8. [OK](#)

P. 15, l. 13: Does SCISAT-1 need to be explained and capitalized? [The name of the satellite is written in lowercase on the ESA website so we don't think it needs to be written in capital letters.](#)

l. 24: monotonically, not monotonously [OK](#)

P. 16, l. 17: better English would be . . .the MLS bias described is. . . [OK](#)

REVIEW #2

Major

There seems to be two major obstacles to overcome during periods when the PLL was malfunctioning: frequency shifting and line broadening. The approaches taken to overcome these obstacles seem reasonable, but more information must be provided in the paper in order to assess whether this is the case.

1. The paper is not entirely clear about this, but my impression is that from 2001-2003, the PLL was malfunctioning but only a frequency correction was needed. In 2003-2004, the PLL was working and no corrections were applied. Then from 2004 onward, the PLL was malfunctioning but both the frequency and broadening corrections were needed. My naive assumption was that the broadening was a result of the frequency shift occurring during the time of a single spectrum, but this seems inconsistent with how the corrections were applied. In any event, the paper needs to have more information to better clarify the malfunctions of the PLL during the pre 2003 and post 2004 time periods. [Section 3.3 has been updated to better clarify that we think two different kinds of PLL malfunctioning affected the instrument during these two periods: one before 8 October 2003 which had effect only at longer time scales and caused only frequency shifts; and another one after 8 October 2004 which have effects both within the integration time and at longer time scales, causing both line broadenings and frequency shifts.](#)

2. For the frequency correction, a "basic" correction applied to all spectra in a scan is derived using differences in vertical gradients of brightness temperature (T_b) between O3 and CO. This is a novel idea and it seems to work well. However, Fig 3 compares T_b profiles for

global averages, and one might expect to see differences according to latitude or season since vertical gradients in O₃ and CO mixing ratios do change with season and latitude. Whether seasonal or latitudinal variations matter for this "gradient threshold" method is something that needs to be demonstrated. There should also be bars or shading to show the sigma in Fig 3. A panel which represents the fitted values of the slopes together with uncertainties has been added to Figure 3 and is described in Section 3.1. In this panel it can be seen how the difference between the two slopes is bigger than any natural variability. We therefore confirm that the threshold we are using is valid for whatever season and latitude.

3. A second, single-altitude frequency correction seems to be applied using Gaussian fits, but it is not discussed in sufficient detail at the top of p. 6. Why is a Gaussian thought to be the appropriate lineshape (later, thermal broadening is mentioned but what about pressure broadening at the bottom of a scan)? Was the central frequency and half-width fitted independently using some least-squares method? If so, how do the fitted widths compare with those expected? Was this second stage of frequency correction required for both 2001-2003 and post 2004 periods, or only post 2004 as for the broadening correction? The Gaussian fits which are discussed here are not to be confused with the ones performed during inversions to retrieve concentration profiles. Here we are using Gaussian fits exclusively as a method to estimate the center frequency of the line so to assess what is the frequency correction that needs to be applied to the spectrum. The fitted width, in this case, is of no interest. This is done, prior to the inversion process, for all scans. Another method to obtain the same information could have been to locate the position of the maximum peak of the line. But the use of one method or the other didn't present significant differences in the estimated centrum. In fact, even where the line is not Gaussian, the central part can still reasonably be fitted with one and the obtained center will be correct.

4. Figures 2 and 4 should have color bar legends to show the approximate range of altitudes covered. The paper has been updated with legends reporting the colours corresponding to some of the altitudes. Not all altitudes are indicated in the legend for the sake of readability.

5. It is important to show how the frequency shift and broadening varied with time, to give an idea of the magnitudes and temporal variation of both corrections. For example, monthly means of both quantities such as in Fig 7, used for the total number scans, would give the reader this kind of information. As described in Section 3.3, the broadening correction applied was one and the same for the whole period after 8 October 2004. The sigma value chosen for the new response function is the best compromise over this period, since it is the one giving the lowest overall area differences. Thus, the broadening due to the PLL malfunction was assumed to have the same entity over the whole period after 8 October 2004. The period before instead was not affected by any broadening. Regarding frequency shift, it appeared to be random and not follow any trend with time. We also observed no dependency on the satellite temperature. Therefore, we chose not to include any time series reporting the temporal variations of the corrections. This information has been added in Section 3.1.

6. Related to number 5 above, in the dataset there should be quality or error flags (or raw information such as mean scan frequency shift and broadening correction) so that users can further filter or weight these data for scientific study. The paper does not discuss any error estimates in the dataset. ARTS retrieval algorithms are based on the Optimal Estimation Method (OEM), thus statistical errors come from there (information added to the paper in Section 3.2). Moreover, in Section 3.1 we already explain how we discard spectra were CO

line is not present. For all the other spectra, the PLL-generated frequency shifts and line broadening have been corrected, so there is no need of data flags.

Minor

1. abstract line 7 "The much of the..." -> "Much of the..." OK
2. p. 2 l. 3 "Schumann-Runge *bands* and continuum..." OK
3. p. 2. l. 27 "This is the first data being part of the..." is confusing Changed to "they are part of the..."
4. caption of Fig 3 "...and in the right place." -> "...and at the expected frequency." OK

5. p. 5 l. 13 "...a value higher than -0.0045 K/m..." Since all slopes are negative, this is confusing. I think what is meant is a slope with a magnitude less than 0.0045 K/m. Clearly, O3 has a larger magnitude in the vertical gradient of T_b, and CO a much smaller vertical gradient in T_b. Referring to slopes in Fig 3 is confusing because CO has a steeper slope, but a smaller vertical gradient. A panel which represents the fitted values of the slopes together with uncertainties has now been added to Figure 3. With the addition of this panel the text is not confusing anymore and it has been left unchanged.

6. p. 6 l. 7 define ARTS at first use OK

7. p. 7 l. 10 "Despite *the fact that* spectra above 40 km tangent altitude are considered..." OK

8. p. 8 l. 6 Is the given sigma FWHM or HWHM? Neither of the two. Sigma is the standard deviation of the Gaussian curve. The relation between sigma and FWHM is given by: $FWHM = 2 \cdot \sqrt{2 \cdot \ln(2)} \cdot \sigma$. We have not modified the text since this simply corresponds to the standard definition of a Gaussian function.

9. p. 10 l. 1 High CO vmr due to enhanced descent at the stratopause following an SSW was clearly demonstrated by Manney et al (Atmos. Chem. Phys., 9, 4775–4795, 2009). Reference added.

10. p. 12, l. 5-10 Not enough attention is given to possible biases due to differences in vertical resolution. There can be large vertical gradients in CO that depend on altitude and season. Between 70km and 90 km, where some biases are shown, CO increases by an order of magnitude over 20 km and differences in vertical averaging could be a factor. The mean averaging kernel widths between the various instruments should be given. For example, it is possible that ACE has at least a factor of 2 higher vertical resolution than SMR. Regarding comparisons with MIPAS, the polar summer mesosphere, where there most striking differences occur, is indeed characterised by very large vertical gradients in the CO distribution. In consequence, differences in vertical resolution between both instruments are likely important, there. While the vertical resolution of mesospheric MIPAS daytime CO is around 5 km (similar to that of SMR), nighttime vertical resolution is worse (>10 km) because of the lower sensitivity caused by smaller nighttime non-LTE emissions (no solar excitation). Therefore, it could be possible, that the SMR-MIPAS comparison is affected by the lower MIPAS nighttime resolution. This has been checked by restricting the comparison to MIPAS daytime profiles, but the resulting difference profiles are very similar to the ones obtained considering data altogether. This information has been added to Section 5.1.

Moreover, we smoothed SMR and comparison instruments profiles using Gaussian filters with various FWHMs. This has been done in order to account for differences in vertical resolution, but the test didn't show any improvement in concentration differences. This suggests that differences in vertical resolution are not the cause of the observed biases. This information has been added to the Section "Conclusion" of the paper.

Recovery and validation of Odin/SMR long term measurements of mesospheric carbon monoxide

Francesco Grieco¹, Kristell Pérot¹, Donal Murtagh¹, Patrick Eriksson¹, Peter Forkman¹, Bengt Rydberg², Bernd Funke³, Kaley A. Walker⁴, and Hugh C. Pumphrey⁵

¹Department of Space, Earth and Environment, Chalmers University of Technology, Gothenburg, 412 96, Sweden

²Molflow, Gråbo, 443 40, Sweden

³Instituto de Astrofísica de Andalucía, CSIC, Granada, Spain

⁴Department of Physics, University of Toronto, Toronto, M5S 1A7, ON, Canada

⁵School of GeoSciences, University of Edinburgh, Edinburgh, EH9 3FF, UK

Correspondence: Francesco Grieco (francesco.grieco@chalmers.se)

Abstract.

The Sub-Millimetre Radiometer (SMR) on board the Odin satellite performs limb sounding measurements of the middle atmosphere to detect molecular emission from different species. Carbon monoxide (CO) is an important tracer of atmospheric dynamics at these altitudes, due to its long photochemical lifetime and high vertical concentration gradient. In this study, we have successfully recovered over 18 years of SMR observations, providing the only dataset to date being so extended in time and stretching out to the polar regions, with regards to satellite-measured mesospheric CO. This new dataset is part of the Odin/SMR version 3.0 level 2 data. ~~The much~~ Much of the level 1 dataset - except the October 2003 to October 2004 period - was affected by a malfunctioning of the Phase-Lock Loop (PLL) in the ~~frontend~~ front end used for CO observations. Because of this technical issue, the CO line could be shifted away from its normal frequency location causing the retrieval to fail or leading to an incorrect estimation of the CO concentration. An algorithm was developed to locate the CO line and shift it to its correct location. Nevertheless, another artifact causing an underestimation of the concentration, i.e. a line broadening, stemmed from the PLL malfunctioning. This was accounted for by using a broader response function. The application of these corrections resulted in the recovery of a large amount of data that was previously being flagged as problematic and therefore not processed. A validation study has been carried out, showing how SMR CO volume mixing ratios are in general in good accordance with the other instruments considered in the study. Overall, the agreement is very good between 60 and 80 km altitude, with relative differences close to zero. A positive bias at low altitudes (50 - 60 km) up to +20% and a negative bias up to -20% at high altitudes (80 - 100 km) were found with respect to the ~~compared~~ comparison instruments.

1 Introduction

Of the carbon monoxide (CO) produced at surface level by anthropogenic sources (e.g. industrial activities, biomass burning, transport and heating) as well as by oceans and biogenic sources, very little is transported upwards to the stratosphere, as it is chemically destroyed by reacting with the hydroxyl radical (OH). This causes a sharp gradient in CO concentration from the

troposphere to the lower stratosphere (Zander et al., 1981). The CO which can be observed in the middle atmosphere is mainly produced via two processes: methane (CH_4) oxidation and CO_2 photolysis, the latter being the dominant one (Minschwaner et al., 2010). In particular, high altitudes are characterised by strong fluxes of radiation in the Schumann-Runge bands and continuum and Lyman- α wavelengths which are strongly absorbed by CO_2 , therefore CO_2 photolysis becomes more significant with height in the upper mesosphere and lower thermosphere, providing a major source of CO and resulting in a large vertical gradient in its concentration. CO from high altitudes is transported downwards in the winter hemisphere polar night region due to advection and vertical eddy mixing (Solomon et al., 1985). In the mesosphere and lower thermosphere, the photochemical lifetime of CO increases with altitude from a minimum of approximately one week up to several hundreds of years. This lifetime is greater than the zonal transport time scale and of the same order of magnitude as the meridional and vertical transport time scales in the stratosphere up to the middle mesosphere, while it is significantly greater than all transport times scales in the upper mesosphere (Brasseur and Solomon, 2005). Because of its strong horizontal and vertical concentration gradients and to its long lifetime, CO is commonly used as a tracer of middle atmospheric dynamics (e.g., Lee et al., 2011; de Zafra and Muscari, 2004).

Measurements of CO in the mesosphere have been carried out since the end of the 70's using ground-based instruments (Clancy et al., 1982) and later on with satellites such as UARS/ISAMS (Allen et al., 1999). More recent satellite measurements are the ones performed in the microwave band with Aura/MLS (e.g., Froidevaux et al., 2006), and in the infrared with Envisat/MIPAS (e.g., Funke et al., 2009) and Scisat-1/ACE-FTS (e.g., Clerbaux et al., 2008). The first retrieval results from CO measurements obtained with the Sub-Millimetre Radiometer (SMR) on board Odin were presented by Dupuy et al. (2004). They observed seasonal variations of CO concentration related to global circulation and chemical processes as predicted from the Whole Atmosphere Community Climate Model (WACCM), to which they compared their results obtaining an overall agreement within two orders of magnitude. Also, good agreement has been obtained comparing with observations from UARS/ISAMS.

There are not many studies about Odin/SMR measurements of CO, although such measurements have been performed from August 2001 until today. This is due to a malfunctioning of the Phase-Lock Loop (PLL) in the ~~frontend~~front end used for CO observations, which caused the majority of CO data to present artifacts which made them unusable for retrievals. The issue is illustrated in Section 2 together with a description of Odin/SMR. In Section 3 we explain how we implemented a correction for the artifacts caused by the PLL malfunctioning and other issues which arose during the retrieval process. As a result, retrieval products from over 18 years of Odin/SMR CO observations, between 50 km and 100 km altitudes, are now available and presented in Section 4. ~~This is the first data being~~They are part of the Odin/SMR v3.0 L2 dataset. Finally, for validation purposes, in Section 5 we compare Odin/SMR profiles with those from Envisat/MIPAS, Scisat-1/ACE-FTS and Aura/MLS, being the satellite instruments operating at the same time as SMR and observing similar altitudes. We also present comparisons with ground-based measurements from Onsala Space Observatory.

2 Odin/SMR CO measurements

2.1 The sub-mm radiometer

The Odin satellite, a Swedish-led project in collaboration with Canada, France and Finland, was launched on 20 February 2001 into a 600 km sun-synchronous orbit with inclination 97.77° and 18:00 hrs ascending node. Its observation time was shared between astronomical and atmospheric observations until 2007, when the astronomical part of Odin's mission was concluded. After that, the instruments on board of Odin have been used exclusively for limb sounding of the atmosphere. These instruments are OSIRIS (Optical Spectrograph and InfraRed Imaging System) and SMR. In this article, we are using the latter. The four sub-mm receivers in SMR can be tuned to cover frequencies between 486 - 504 GHz and 541 - 581 GHz, thus to observe emission due to rotational transitions for species such as O_3 , H_2O , CO, NO, ClO, N_2O , HNO_3 in the stratosphere and mesosphere (e.g., Frisk et al., 2003). There is also a mm receiver to observe the 118 GHz O_2 transition. The cold sky and a hot load are repeatedly observed for calibration purposes. ~~Through a Dicke switch, the signal from a calibrator and~~ (see Figure 1). ~~A Dicke switch allows to quickly change the source of the signal between the main beam signal are deviated towards different directions, and further and the calibrators. The signal is thereafter split according to polarisation, before being collected by different receivers (see Figure 1). Every signal is thereafter and injected into the mixer through a Martin-Puplett interferometer.~~ Here the source signal is converted to longer wavelengths by combining it with a local oscillator (LO) signal ~~through a mixer, also injected through the Martin-Puplett interferometer.~~ The frequency of the LO is fine tuned with the use of a PLL. The output signal of the mixer is a function of the sum and difference of the frequencies of the input signals, therefore consisting of two sidebands. SMR is however run in single sideband mode so, of the two sidebands, only the one containing the signal of interest should be transmitted and the other one suppressed. This is achieved, before mixing, by accordingly setting the length of the arms of the Martin-Puplett interferometer. The resulting signal can then be amplified and routed to the autocorrelator spectrometers (AC1 and AC2, as indicated in Figure 1).

Measurements are performed during both upward and downward vertical scanning with a vertical sampling of ~ 6 km in the mesosphere. ~~The scanning is continuous, with a constant speed of 0.75 km/s.~~ Odin/SMR L1 data are organised in scans, each of them consisting of a group of spectra collected during a single upward (or downward) scanning. Each spectrum corresponds to a single mean tangent altitude in the range 7 - 72 km (stratospheric scans), 7 - 110 km (strato-mesospheric scans) or 60 - 110 km (mesospheric scans) (Dupuy et al., 2004).

2.2 SMR CO operational modes

Odin/SMR measures CO, with a 3 - ~~4-7~~ km vertical resolution, from the thermal emission line corresponding to the $J = 5 \rightarrow 4$ rotational transition at 576.268 GHz. In Figure 2a we show an illustrative scan of spectra obtained in normal conditions where both a CO line and a O_3 line are measured. The SMR receivers can be adjusted to different configurations called frequency modes (FM), each of which corresponds a certain observed frequency band and a scheduled observation time. There are three FMs that cover bands in which the above mentioned CO transition is observed. Their characteristics are summarised in Table 1. All these measurements are carried out by using the B1 ~~frontend-front end~~ (the set of components thus denoted in Figure 1),

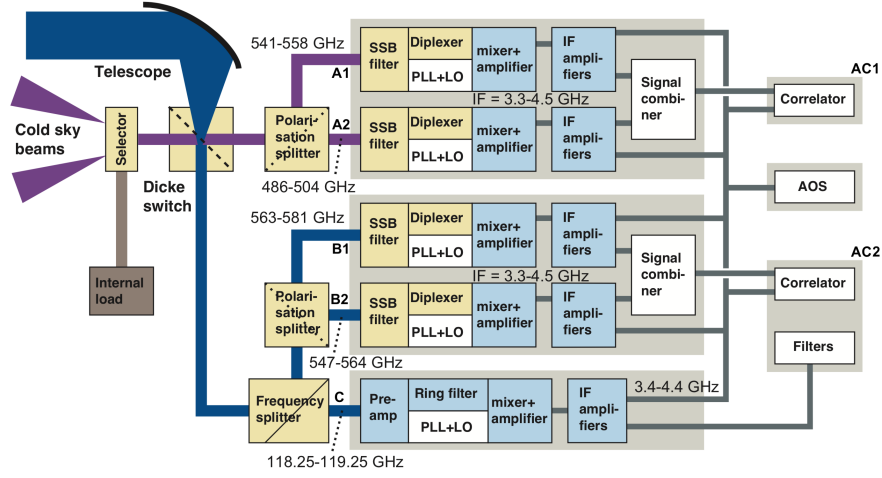


Figure 1. Block diagram of the Odin radiometer. From Frisk et al. (2003).

Table 1. FMs for observing CO. From Rydberg et al. (2017).

Frontend	Spectrometer	LO Freq. [GHz]	Freq. Range [GHz]	Species	FM
572 B1	AC2	572.762	576.062 - 576.862	CO, O ₃	14
		572.964	576.254 - 576.654 577.069 - 577.469	CO, O ₃ HO ² , ¹⁸ O ₃	22
	AC1	572.762	576.062 - 576.862	CO, O ₃	24

whose PLL element was working correctly for a year only, between 8 October 2003 and 8 October 2004. The measurements made during the rest of the Odin operational time, from 2001 until today, have all been affected by a malfunctioning of this PLL, leading to a shift of the LO frequency from its nominal value (Rydberg et al., 2017). Consequently each scan presents a different frequency shift. In extreme cases like the one shown in Figure 2b, the frequency shift causes the CO line to fall outside the observed bandwidth and thus the data are unrecoverable.

3 Recovery and retrieval

We explain here how we developed a frequency correction which was applied to the spectra before the retrieval process, leading to the recovery of a great part of the dataset. This is followed by descriptions of the set-up for the v3.0 CO retrievals and of the method used to estimate line broadening generated by the PLL failure.

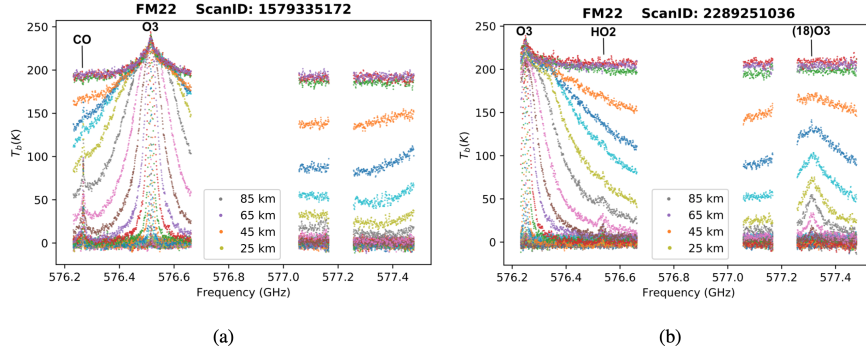


Figure 2. Spectra for altitudes ranging from 7 km to 110 km for two scans taken as examples. The bigger gap in the spectra around 576.8 GHz is due to the particular way the bands are arranged to cover the desired frequencies for FM22, while the smaller gap at 577.2 GHz is due to instrumental failure relative to one sub-band. In (a) both CO and O₃ lines are present and in-at the right place expected frequency. In (b) the frequency shift is so big that only the O₃ can be observed (shifted) while the CO line is shifted outside of the band and therefore not observed. Colors corresponds to different tangent altitudes within the reported scan. Not all altitudes are indicated in the legend for the sake of readability.

3.1 Basic frequency correction

In order to correct the frequency shift, as a first step, the centre frequency of the CO line in the scan's average spectrum has been compared with the theoretical centre frequency. The resulting difference is then applied to the LO frequency. To do that, the correction algorithm first needs to distinguish the CO line from the O₃ line. In Figure 3a it is shown how peak brightness temperatures of CO and O₃ lines vary with altitude. Here it can be noticed how in the 40–60 km altitude range the that the CO and O₃ slopes are most different between 40 and 60 km, allowing us to discern between the two species. Averaging In Figure 3b, averaging of observations during the PLL working period show that, in this altitude range, a -0.0004 ± 0.001 K/m slope corresponds to the CO line, while the value -0.009 ± 0.003 K/m identifies the O₃ line, where the uncertainties on slopes correspond to 3σ . Thus, the correction algorithm associates peak T_b gradients with a value higher than -0.0045 K/m to the CO line. The LO frequency can then be corrected for the whole altitude range under consideration in this study, as explained above.

If no line or only the O₃ line is found, the scan is not considered for further processing.

Despite the application of this first correction to the data, they still present artifacts that need to be corrected. In fact, as can be seen in Figure 4a, all spectra in a single scan, each corresponding to a different altitude, present a different frequency shift. This suggests that the PLL malfunctioning affects the observation at a time scale smaller than the scanning time scale (~ 2 minutes). Thus applying the same correction to each spectrum in a scan is not sufficient. To solve this problem, we modified the pre-correction algorithm by considering each single altitude spectrum and estimating the observed center frequency of each CO line by fitting them with a Gaussian. We then compared these values to the theoretical center frequency and therefore applied a different shift to each spectrum. An example of result obtained from this correction is shown in Figure 4b. Including this altitude dependent correction resulted in a significantly increased amount of data that could be recovered, i.e. from which

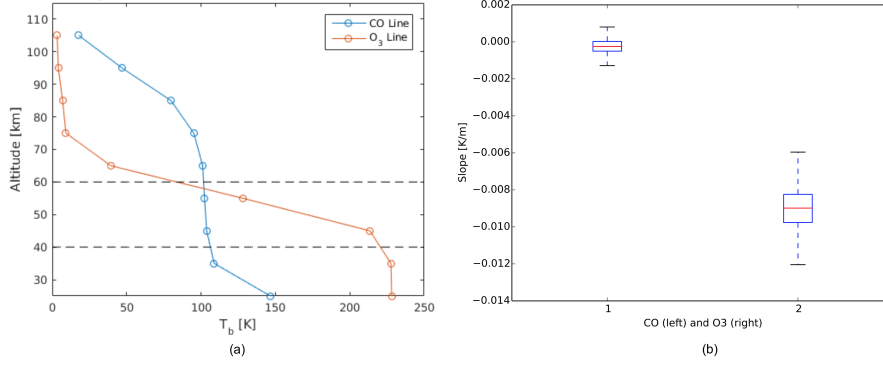


Figure 3. Panel (a): Peak brightness temperatures of CO and O₃ at different altitudes. Average considering all observations performed during the PLL working period, i.e. 14224 scans. At altitudes between 40 km and 60 km, the CO line curve presents a steeper slope. Panel (b): Linear fit of slopes of peak brightness temperatures between 40 - 60 km altitude for CO and O₃. The shown uncertainties on slopes correspond to 3σ .

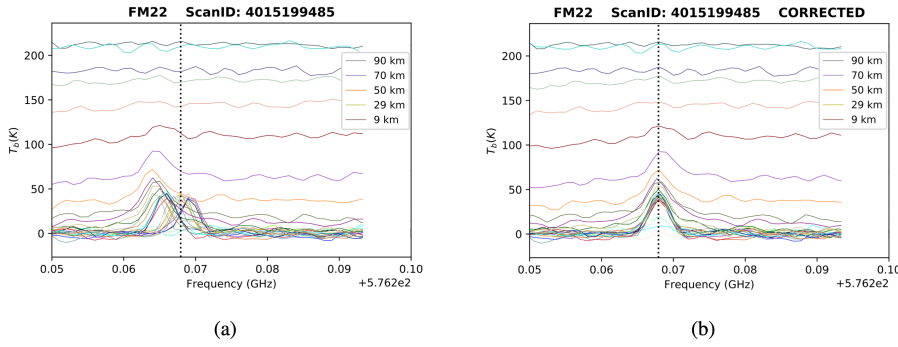


Figure 4. Spectra measured on 2 February 2009 presenting altitude dependent frequency shifts before (a) and after (b) altitude dependent correction. Colors corresponds to different tangent altitudes within the reported scan. Not all altitudes are indicated in the legend for the sake of readability.

CO concentration profiles could be derived (as shown in Figure 7 in Section 4). The frequency shifts, of which we presented the corrections in this Section, appear to be random and not follow any trend with time. No dependency on satellite temperature was observed either.

5 3.2 Retrieval set-up

The Atmospheric Radiative Transfer Simulator (ARTS) retrieval algorithms are based on the Optimal Estimation Method (Rodgers, 2000) from which statistical errors on the retrieved quantities originate. A new version of ~~the~~ ARTS operational processing system has been developed for SMR retrievals. The data produced by this system will be denoted as version 3.x.

The new retrieval system will be described in detail in a forthcoming publication, but a summary is provided below. The CO data discussed in this work have been assigned version number 3.0.

The new processing system is based on a MySQL database and is capable of distributing the calculations over multiple clusters. The actual retrievals are based on ~~the Atmospheric Radiative Transfer Simulator (ARTS, Buehler et al. (2018))~~ [ARTS \(Buehler et al., 2018\)](#). For the moment ARTS version 2.3.564 is applied. Besides atmospheric radiative transfer, the forward simulations consider the sensor's antenna response, double-sideband characteristics and spectrometer frequency response, using the approach of Eriksson et al. (2006). ARTS provides the needed weighting functions (mainly using analytical expressions), while the inversion of optimal estimation type is made in Matlab by code taken from Eriksson et al. (2005).

The L2 data provided contains a characterisation of the retrieval, following Rodgers (2000). As described in Baron et al. (2002), when retrieving multiple quantities in parallel, the "smoothing error" includes terms that describe the interference between the quantities. The retrieval error reported in SMR v3.0 data incorporates this cross-quantity interference, but excludes the classical smoothing error internal to the quantity. For example, the error reported for CO considers that the retrieval of instrumental parameters is non-ideal, but excludes the direct smoothing of the CO profile due to limitations in vertical resolution. In this way, the reported error matches the standard one that would be obtained if the instrumental parameters would instead be treated as forward model uncertainties (though strictly true only for a totally linear inversion problem).

Specifics for the retrievals presented in this paper include that only spectra recorded at tangent altitudes between 40 and 100 km are considered, to form a "mesospheric retrieval mode". The spectroscopic data for the frequency range of concern is solely taken from HITRAN 2012 (Rothman et al., 2013) (but the system allows to incorporate data from other sources). The set of variables retrieved is specified for each mode separately. In this case, the CO profile and off-sets for pointing, frequency and brightness temperature zero level (baseline) are retrieved.

The a priori dataset previously used for Odin/SMR CO retrievals was found outdated and inaccurate. As a consequence, for the new inversions, a new CO climatology was formed, based on MIPAS zonal means averaged over the years separately for each month. For this purpose, we are using the product V5R_CO_521, which at the moment is the most recent data version from MIPAS middle atmosphere observation mode (Garcia et al., 2014) (see Section 5.1). [The monthly a priori is divided into 10° latitude bins. As temperature model for the retrieval, ERA-interim reanalysis data \(Dee et al., 2011\) is used up to 60 km and the Mass Spectrometer Incoherent Scatter model \(version NRLMSISE-00; Picone et al., 2002\) is used from 70 km upwards. Between 60 - 70 km, a spline interpolation of the two is applied.](#)

In Figure 5 is shown the retrieval for an exemplary scan. Despite [the fact that](#) spectra above 40 km tangent altitude are considered in the inversion, only measurements above 50 km are reliable since at lower altitudes it is not possible to discern the CO line from the noise. Moreover, data with a measurement response lower than 0.75 are discarded. ~~This is the case of the~~, [where the measurement response is a measure of the degree to which the result may be contaminated by the a priori. It is defined as the sum over the row of the averaging kernel matrix \(Rodgers, 2000\). Values lower than 0.75 occur for the](#) retrieved profile above 100 km which is dominated by the a priori and is here shown out of completeness, together with the averaging kernel extending to higher altitudes, to display how retrieved concentrations between 90 and 100 km are also influenced by higher altitudes.

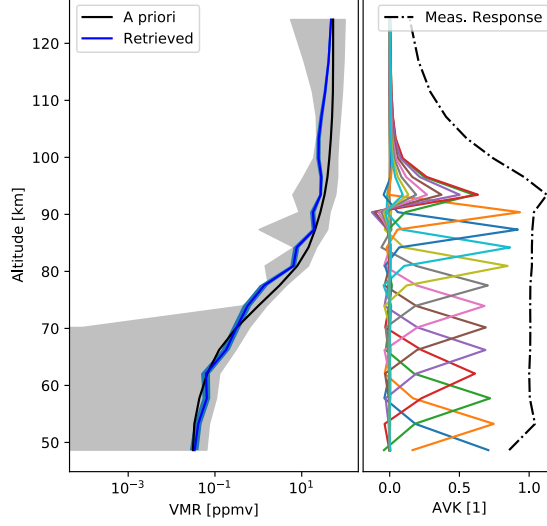


Figure 5. Example retrieval referring to ScanID 1579335172 from FM 22. Left: Retrieved concentration profile and error (in blue) and a priori including uncertainties (in grey). Right: Averaging ~~kernel~~ kernels plotted in a different colour for each altitude (not indicated) and measurement response.

3.3 Broadening correction

The fits of CO line performed during the inversion process - which are Gaussian shaped due to Doppler broadening - have significantly different amplitude and width compared to the observed line, causing an underestimation of the concentration. This suggests that the PLL malfunctioning even has an effect within the integration time, causing the observed line to be broadened with respect to what it would be in normal instrumental conditions. To take into account this broadening issue during the inversion, the original response function, a Gaussian with $\sigma = 0.8$ MHz, has been replaced with a Gaussian with a higher σ value. The response function $w_{ch}^i(\nu)$ of a spectrometer's channel i represents how the power of a signal $I(\nu)$ is weighted:

$$y_i = \int_0^\infty I(\nu) w_{ch}^i(\nu) d\nu \quad \text{with} \quad \int_0^\infty w_{ch}^i(\nu) d\nu = 1 \quad (1)$$

where y_i is the final, calibrated antenna temperature of the channel i (Eriksson et al., 2006).

In Figure 6a is represented the relative difference between the Gaussian area beneath the observed CO line and the Gaussian area beneath the fit, for a scan in the period when the PLL was working. Here there is no need for correction, and the lower values of the relative area difference are obtained with the original response function. As an example, in Figure 6b is plotted

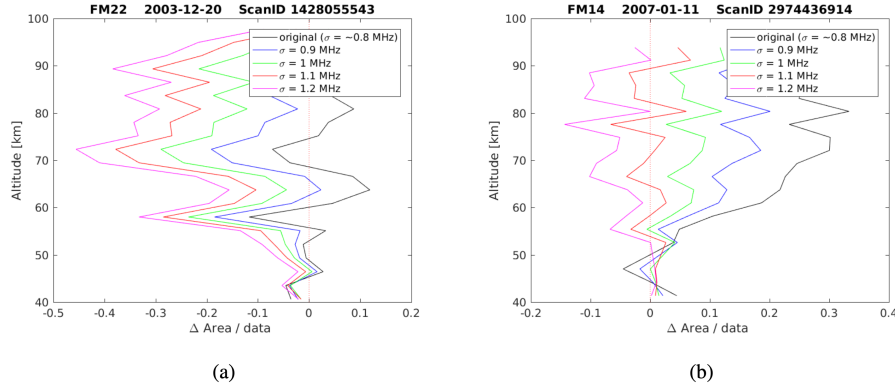


Figure 6. Relative difference between the Gaussian area beneath the observed CO line and the Gaussian area beneath the fit, obtained with different response functions for a scan in the PLL working period (a) and non-working period (b). The best response functions for the two cases are, respectively, the original one (black line) and the one with $\sigma = 1.1$ MHz (red line).

the relative area difference for one scan in the PLL malfunctioning period ~~after 8 October 2004~~. During this period, the most suitable response function is a Gaussian with $\sigma = 1.1$ MHz. This is indeed the one resulting in the lowest overall relative area differences, therefore this same response function has been used in the whole period after 8 October 2004, aforementioned.

Note that the original response function is still the one giving the best results during the period before 8 October 2003, ~~probably because the PLL malfunctioning in the very beginning of the satellite operation time, 2003. Consequently, the new response function has been applied only to inversions after 8 October 2004. This suggests that the PLL malfunctioning before 8 October 2003 is of different nature than the one after 8 October 2004. Consequently, the new response function has been applied only to inversions after 8 October 2004, 2004, with the former causing only frequency shifts and the latter causing both line broadenings and frequency shifts, i.e. having effects both within the integration time and at longer time scales.~~

4 The new dataset

The recovered dataset is part of the Odin/SMR v3.0 L2 data. As mentioned in Section 3, the new frequency correction algorithm also helped to recover many data that were previously being erroneously flagged, so that there is a very significant increase in the amount of CO level 2 data that are now available. This can be seen in Figure 7, showing the temporal distribution of the L1 and L2 data sets before and after the application of the correction algorithm, for the three CO observation modes together. The results are very satisfying, with 63% of the data being successfully processed (compared to 8% before the correction) until September 2017. The remaining data are discarded due to other quality flags. After this date, the frequency shift is causing the CO line of almost all the spectra to be outside of the observed band, resulting in the loss of most of the CO data during this time period, hence the almost total absence of L2 data from there on. The very high number of measurements performed with CO FMs during July 2002, July 2003 and August 2004 corresponds to special scheduling of the observation time, set to

monitor dynamics in the northern summer mesosphere associated with the study of noctilucent clouds (Karlsson et al., 2004).

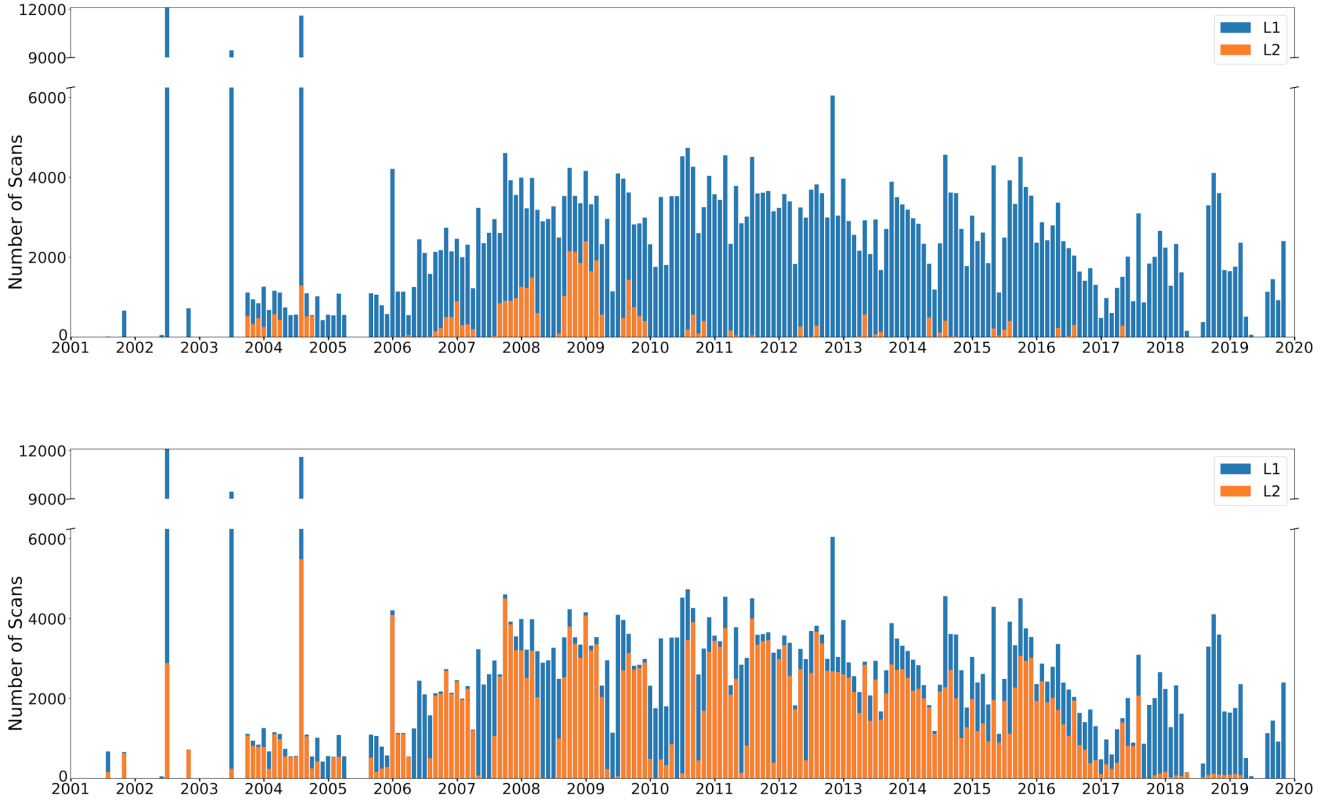


Figure 7. Number of L1 and L2 scans by month for FM14, FM22 and FM24 altogether, before (top) and after (bottom) the application of the correction algorithm. The ticks on the x-axis correspond to January 1st for each year. This figure gives an overview of how much data could be recovered.

The time series shown in Figure 8 gives an overview of the new CO data set, extended to cover the whole globe until the polar regions. They consist of monthly zonal means of CO volume mixing ratios over five different latitude bands. The white bands correspond to months during which the number of scans in the given latitude band is lower than 10. No concentrations are shown for some months in the early years of the mission, despite the presence of sufficient L2 data, because of plotting interpolation with adjacent months with no data (see Figure 7). No significant difference between the various FMs is observed (not shown). CO volume mixing ratios show noticeable variations with altitude, latitude and season, as well as longer-term variations. We observe a sharp increase with height, due to the photodissociation of CO_2 at high altitudes, as explained in Section 1. In these plots, it is also possible to notice the temporal variation of CO in the mesosphere throughout the years. In the tropics, maxima appear at equinoxes and minima at solstices. This is explained by the Semiannual Oscillation (SAO)

signal, which dominates at low latitudes (Lee et al., 2018). Although the SAO-induced variations in zonal mean vertical wind are too small to be measured directly, the accumulated effects can be seen in the distribution of the long-lived constituents like CO (Hamilton, 2005). Sinking of air, or at least anomalously weak rising motion, leads to the observed increase in the CO mixing ratios around the equinoxes. At high latitudes, the seasonal variations are mainly characterised by the downward transport of CO rich air in winter, from the upper mesosphere to the stratopause, induced by the meridional circulation. The CO volume mixing ratio values measured in summer are lower, due to upward circulation. The SAO signal is noticeable ~~in~~ this latitude band at high latitudes too, explaining the secondary minima that are visible in the middle of the winter (Lee et al., 2018). The seasonal variations at mid latitudes are similar, though the effect of the meridional circulation is significantly less pronounced than at high latitudes. Moreover, at high latitudes in the northern hemisphere, a particularly strong secondary peak in CO concentration appears during several of the winters. This is observed in particular in 2006, 2009, 2013 and 2019. As seen in Figure 8, unusually high volume mixing ratio values were measured in the mid and lower mesosphere, in those years, in late winter. Such a pattern is generated by Sudden Stratospheric Warming (SSW) events followed by an elevated stratopause event. SSWs occur mainly almost entirely in the northern hemisphere, during wintertime, and consist of warmings of the stratosphere at high latitudes by several tens of Kelvins, occurring within a few days. This is caused by planetary waves disturbing the polar vortex (e.g., Charlton and Polvani, 2007), thus reducing the descent of CO rich - air from higher altitudes. After such an event, the vortex recovers, and the stratopause sometimes reforms at higher altitudes than normal (Vignon and Mitchell, 2015), as it has already been observed by Odin/SMR (Pérot et al., 2014). In such a case, the downward transported air will come from higher altitudes (Orsolini et al., 2017), where CO is more abundant, resulting in the above mentioned higher concentration peak in late winter (Manney et al., 2009). Finally, it can be seen that, in all latitude bands, concentrations are in general higher during the period 2012-2016, in accordance with corresponding more intense solar activity. The described temporal variations are consistent with observations from other satellite instruments, such as MLS (Lee et al., 2018) and MIPAS (Funke et al., 2009; Garcia et al., 2014).

5 Comparison with other instruments

In this section we compare the SMR v3.0 CO data set with data from other limb-sounding satellite-borne instruments, namely MIPAS, ACE-FTS and MLS, and with ground-based measurements made by a radiometer located at Onsala Space Observatory (OSO). Our goal is to assess the quality of the new data set described above. With regards to satellite measurements, the comparison is performed between measurements that occurred within a maximum temporal separation of 24 hours and a maximum spatial separation of 500 km. It is possible to use such broad coincidence criteria because of CO's long chemical lifetime in the mesosphere (Minschwaner et al., 2010). CO in the mesosphere and lower thermosphere (MLT) is affected by tidal mixing (Garcia et al., 2014) which might contribute to the biases observed in comparisons with such a broad temporal coincidence criterium. To investigate if this is the case, we also carried out comparisons with 3h and 6h coincidence criteria. When not characterised by the absence of sufficient coincidences (i.e. less than 10), these comparisons do not present biases

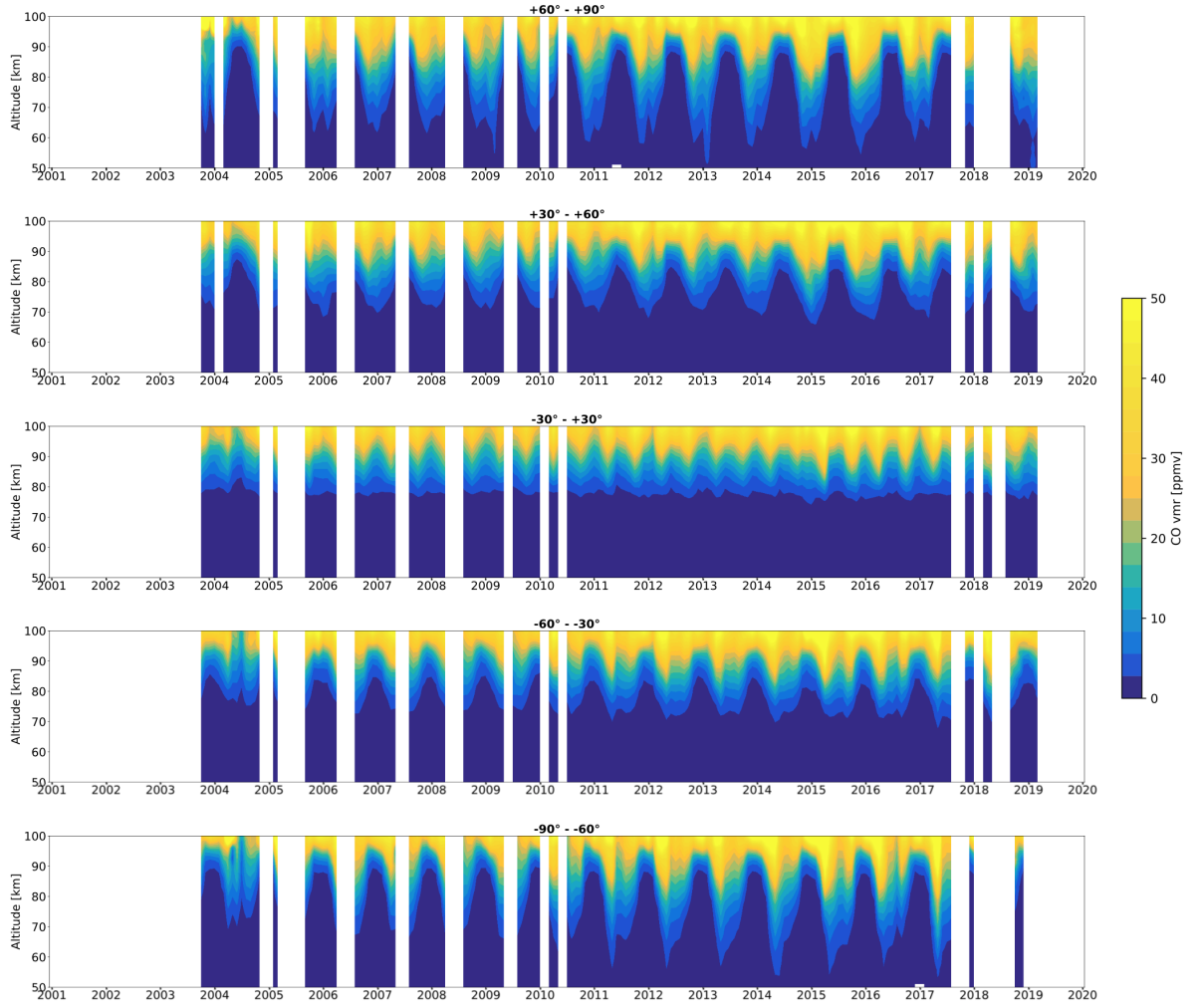


Figure 8. Time series of CO volume mixing ratios measured by SMR for different latitude bands. The white bands indicate periods during which the number of scans in the given latitude band is lower than 10. The ticks on the x-axis correspond to the beginning of each year.

which are significantly different from the 24h ones. No plot is shown for the 3h and 6h comparisons. All the validation plots shown hereafter refer to 24h coincidences.

CO vertical concentration profiles do not feature particular structures which would justify taking into account the satellite instruments' different vertical resolutions, also considering that such differences are not as marked as the ones between SMR and OSO (coincidence criteria and smoothing process for the SMR-OSO comparison are presented in Section 5.4). The comparison between space-borne instruments has therefore been performed simply by linearly interpolating each coincident profile

over a common ~~4 km~~ 1 km altitude grid, ranging from 50 to 100 km. Given a couple of coincident measurements denoted with i , the absolute difference of the two at altitude z will be given by:

$$\delta_{abs,i}(z) = x_{SMR} - x_{comp} \quad (2)$$

and the relative difference:

$$5 \quad \delta_{rel,i}(z) = \frac{x_{SMR} - x_{comp}}{(x_{SMR} + x_{comp})/2}, \quad (3)$$

where x_{SMR} and x_{comp} are respectively the CO mixing ratios measured from SMR and the comparison instrument at altitude z for the coincidence i . The relative difference has as its denominator the mean of the two concentrations. This is done because, being both satellite measurements, they can both be affected by large uncertainties and none of the two is preferable as a reference (Randall et al., 2003). To minimise the weight of outliers, the ~~average~~ median difference $\Delta(z)$ over all the $N(z)$ coincidences at altitude z is calculated ~~as a median~~. The dispersion of the results is represented by the standard deviation of the median, calculated as follows:

$$SEM(z) = \frac{1}{\sqrt{N(z)}} \sqrt{\frac{1}{N(z)-1} \sum_{i=1}^{N(z)} (\delta_i(z) - \Delta(z))^2}. \quad (4)$$

This is valid for both absolute and relative difference. The average and standard error of the concentration profiles are calculated using the same method. For the sake of clarity, in the following subsections we only show vertical profiles averaged over all the found coincidences, regardless of time or location. However, figures showing the relative differences between SMR and the other limb sounders as a function of altitude and latitude, for each season, has been included in the appendix and will be regularly referred throughout the text.

5.1 MIPAS

The Michelson Interferometer for Passive Atmospheric Sounding (MIPAS) is a mid-infrared spectrometer which was launched on board Envisat on 1 March 2002 and was operating until April 2012, when an unexpected loss of contact with the satellite occurred. The satellite travelled at 800 km altitude on a sun-synchronous orbit with a 98.55° inclination and a 22:00 hrs ascending node. The concentration profiles that we use for comparison in this study are retrieved from measurements of $^{12}\text{C}^{16}\text{O}$ roto-vibrational emissions in the $v = 1 \rightarrow 0$ band around $4.7\mu\text{m}$, using the retrieving processor developed at the Institute of Meteorology and Climate Research (IMK) in Karlsruhe and the Instituto de Astrofísica de Andalucía (IAA) in Granada (e.g., Funke et al., 2009; Sheese et al., 2016). The forward model used for CO retrieval takes into account non-LTE ~~effets~~ effects. We consider data from the most recent MIPAS datasets, as specified in Table 2 (Garcia et al., 2014, 2016). All recommendations about the quality filtering of the data have been followed (Kiefer and Lossow, 2017).

Table 2. Characteristics of the MIPAS CO datasets used for comparison. Vertical resolutions refer to the observations in the altitude range 50 - 100 km considered in this study.

Observation Mode	Altitude Range	Vertical Resolution	Spectral Resolution Mode	Time Period	Version
Nominal (NOM)	10 - 70 km	5 - 15 km	Full Resolution (FR)	July 2002 → March 2004	V5H_CO_20
		6 - 16 km	Optimized Resolution (OR)	January 2005 → April 2012	V5R_CO_220
Middle Atmosphere (MA)	20 - 100 km	5 - 15 km			V5R_CO_521
Upper Atmosphere (UA)	42 - 150 km	5 - 12 km			V5R_CO_621

5.1.1 Nominal mode

There are two observation modes called "Nominal": one in use during the period before the interferometer in MIPAS started malfunctioning, when the instrument was being used in full spectral resolution (FR mission); and one in use after the instrument was recovered and brought back to function with a reduced spectral resolution (OR mission) (Oelhaf, 2008). ~~Both-~~

- 5 ~~Regarding FR-NOM and OR-NOM datasets are considered here altogether. With mode, with~~ the above mentioned coincidence criteria we found ~~94989-6088~~ coincident measurements with SMR over the ~~period~~ July 2002 - ~~April-2012-period-during-which MIPAS-was-operating-March 2004~~. The profiles and differences between 50 and 70 km altitude, averaged over the whole time period and over the whole globe, are shown in Figure 9a. ~~SMR-difference-The median relative difference between SMR and MIPAS has a value of +8% at 50 km which decreases to reach -15% at 60 km, and eventually remains constant between 60 -~~
- 10 ~~70 km altitude. No bias characterising a particular latitude or season over time has been identified for this specific comparison, since FR-NOM mode has been operational for only a short period.~~
- ~~Comparing with OR-NOM mode, 88902 coincident measurements have been found over the period January 2005 - April 2012. Figure 9b shows that the median relative difference between SMR and MIPAS remains positive and almost constant with altitude, staying below +15%. The main contribution to this difference comes from latitudes between -25° and +50°. In~~
- 15 ~~particular during northern summer, around +25°, where it reaches values above +40 % between 65 and 70 km. Everywhere else relative difference values remain within ±20% (see Figure A1). To check if the observed differences are due to differences in vertical resolutions between the two instruments, we compared SMR and MIPAS profiles separately for daytime and nighttime observations. In fact, MIPAS vertical resolutions are similar to SMR's for daytime observations while are equal to more than 10 km for nighttime observations, due to lower sensitivity caused by smaller non-LTE emissions (Funke et al., 2009). The average~~
- 20 ~~comparison between daytime observations did not show differences that deviate significantly from the ones observed in Figure 9 (not shown). This suggests that the observed concentration differences are not due to differences in vertical resolution.~~

5.1.2 Middle atmosphere mode

Another observation mode in use during the OR mission is the middle atmosphere mode, covering a larger altitude range. Its average comparison with SMR over 17003 coincidences is shown in Figure 9bc. Both data sets are in general in good agreement

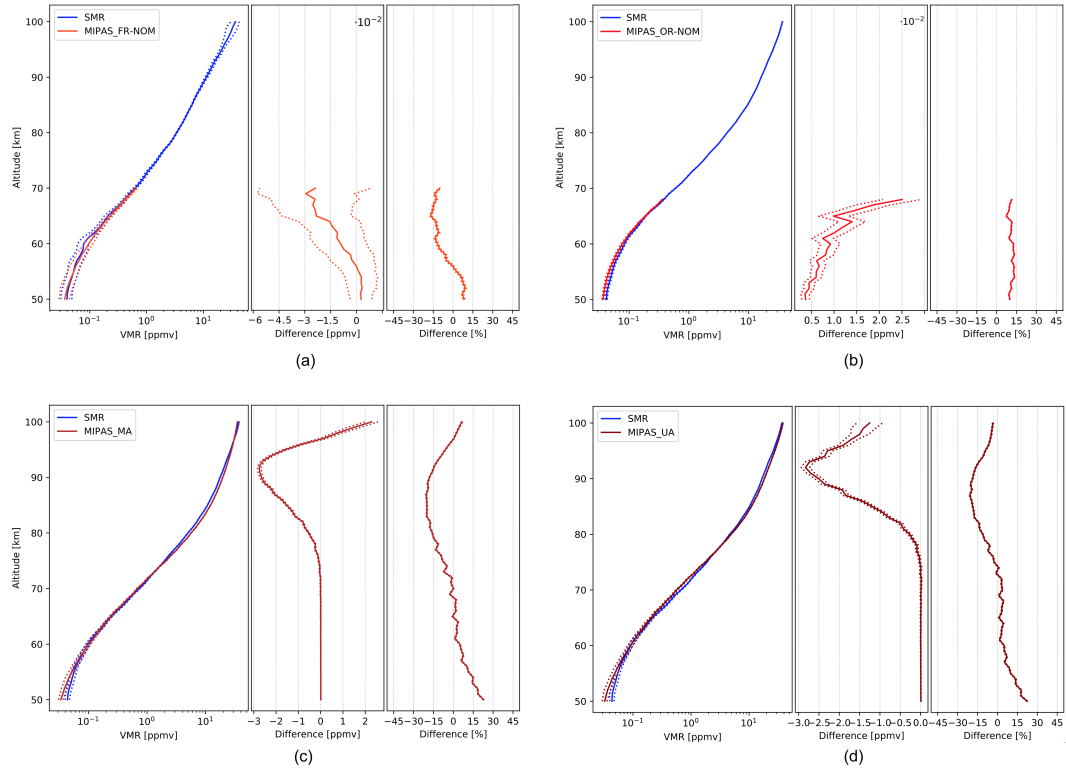


Figure 9. Comparison of SMR CO concentrations with the ones from MIPAS [nominal](#) [FR-nominal](#) (a), [OR-nominal](#) (b), middle atmosphere (c) and upper atmosphere (d) modes. The data plotted are global averages over the whole time periods indicated in Table 2. Left panels: volume mixing ratios, expressed in ppmv. Center panels: absolute differences, expressed in ppmv. Right panels: relative differences, expressed in percentage. The dashed lines represent the standard deviation of the median which, in some cases, is smaller than the thickness of the profile line, causing the dashed line not to be distinguishable.

with each other. It can be seen that the average difference always stays between -20 % and +20%. In particular, SMR presents a positive difference of around +20% at 50 km which decreases with altitude until it becomes null at 60 km. The difference keeps being negligible up until 70 km, then it becomes negative and keeps decreasing until it reaches a minimum of -20% around 85 km after which it goes back up to +5% at 100 km. [The observed difference around 85 km is not to be attributed to MIPAS misestimation of non-LTE effects; in fact non-LTE modelling of MIPAS CO has been validated in Funke et al. \(2007\) and systematic errors due to non-LTE were estimated to be less than 5%.](#) When looking at latitudes separately, the relative differences are mainly within $\pm 20\%$. The most extreme differences are observed around the northern spring equinox. Here we have very negative differences at the equator between 70 and 80 km altitude reaching peaks of -60%. -50% differences can be observed during local summer in both hemispheres at high latitudes between 80 and 90 km. Also peaks of positive difference around +60% are reached at 50 km altitude at -75° and $+50^\circ$ (see Figure A2). [Considerations about daytime and nighttime comparisons carried out separately are the same as in Section 5.1.1.](#)

5.1.3 Upper atmosphere mode

Figure 9e-d shows the comparison with OR upper atmosphere mode, averaged over 19084 coincidences. SMR difference in comparison with this mode is very similar to the one described in [Seet-Section 5.1.2](#). A small dissimilarity is given at higher altitudes where the difference approaches zero but stays negative. [Moreover, as described in Section 5.1.2, observed differences around 85 km are thought not to be caused by MIPAS misestimation of non-LTE effects.](#) At all latitudes and during all seasons, relative difference values are small and generally within $\pm 20\%$. Peaks of almost -50% are reached during the northern spring at the equator between 70 and 80 km and during local summer in both hemispheres at high latitudes between 80 and 90 km (see Figure A3). [Considerations about daytime and nighttime comparisons carried out separately are the same as in Section 5.1.1.](#)

5.2 ACE-FTS

The Fourier Transform Spectrometer (FTS) is an instrument which is part of the Canadian-led Atmospheric Chemistry Experiment (ACE) on board Scisat-1, launched on 12 August 2003 (and still operating) into a 74° inclination orbit at 650 km altitude (Bernath et al., 2005). ACE-FTS performs solar occultation measurements in the mid-infrared. In particular, CO concentrations are retrieved from the roto-vibrational absorption bands around $4.7\mu\text{m}$ ($v = 1 \rightarrow 0$) and $2.3\mu\text{m}$ ($v = 2 \rightarrow 0$). CO retrievals are performed between 5 and 105 km altitude, with a 3-4 km vertical resolution (Boone et al., 2005, 2013). In this study we use ACE-FTS v3.6 dataset for comparison. The data has been quality filtered according to the guidelines from the instrument team (Sheese et al., 2015).

A total of 12925 coincident CO measurements between ACE-FTS and SMR were found between February 2004 (first available ACE-FTS CO measurements) and March 2019. The number of coincidences found is lower compared with any of the MIPAS modes, despite the longer period considered. This is due to the fact that ACE-FTS is a solar occultation instrument, thus scanning the limb only twice per orbit, while both SMR and MIPAS are measuring almost continuously. The spatial and temporal average comparison with ACE-FTS is shown in Figure 10. The relative difference decreases almost [monotonously monotonically](#) with altitude, from a maximum of +15% at 50 km to a minimum of almost -15% at the highest altitudes. In particular, the difference decreases more quickly between 50 and 60 km from 15% to near 0%. SMR and ACE-FTS are in extremely good agreement with each other between 60 and 75 km, with a relative difference close to zero. The difference then slowly decreases with altitude up to about -15% at 90 km, and finally increases almost imperceptibly between 90 and 100 km. Looking at seasons separately, the relative differences present generally low values, almost always within $\pm 20\%$. Values only increase to more than +40% between 60 and 70 km during northern autumn around $+25^\circ$, and during southern winter at the equator and around -25° . The latter difference is similar to the one observed from ACE with respect to MIPAS (Sheese et al., 2016). Moreover, strong negative differences of about -40% are observed at the equator at 80 km altitude during northern spring, and also at $+25^\circ$ during northern autumn between 80 and 90 km (see Figure A4). Note that the majority of ACE-FTS

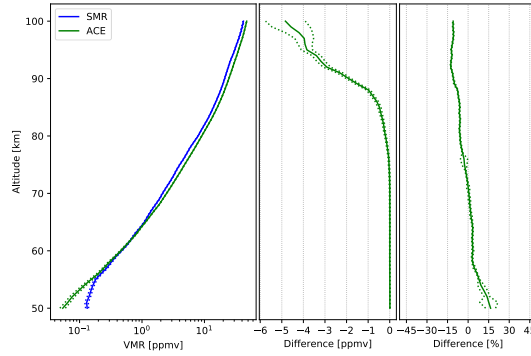


Figure 10. Comparison of SMR CO concentrations with the ones from ACE-FTS retrievals. The data plotted are global averages over the whole time between February 2004 and March 2019. Panels characteristics are the same as in Figure 9.

measurements occur at latitudes higher than 60° and that the above mentioned differences at low latitudes are probably related to averaging on a smaller number of coincidences.

5.3 MLS

The Microwave Limb Sounder (MLS) is an instrument on board the Aura satellite, still operating since its launch on 15 July 2004. Aura/MLS was launched on a 98° sun-synchronous orbit with 13:45 hrs ascending node at an altitude of 705 km. It performs measurements between 118 GHz and 2.5 THz with a 1.5 - 3 km vertical resolution (Schoeberl et al., 2006; Waters et al., 2004). In this study we use MLS CO concentrations from the v4 dataset, retrieved from the $J = 2 \rightarrow 1$ rotational transition emission line at 230.5 GHz, which are considered to be reliable between 0.0046 hPa and 215 hPa. Also, the suggested quality filtering has been performed (Livesey et al., 2018). A previous comparison between SMR and MLS CO retrievals was presented in Barret et al. (2006), based on older versions of the datasets. A validation study of the MLS CO v2 dataset has been performed by Pumphrey et al. (2007) and the differences between v2 and v4 datasets are documented in Livesey et al. (2018).

Because MLS uses the same observation technique as SMR, and because it has been functioning since 2004, a great number of coincident measurements could be used for this validation study. Indeed, 227820 coincidences were found between the beginning of MLS mission and March 2019. The overall average comparison of these measurements with the ones from SMR is shown in Figure 11. A negative difference of SMR with respect to MLS characterizes all of the altitude range. The relative differences are oscillating between -40% and -10% and are characterised by a significant variability. This is due to MLS CO profiles being rather jagged, as reported in Errera et al. (2019) where the ~~described-MLS-bias~~ MLS bias described is in accordance with what we obtain in this study. Also, the MLS-ACE bias which is reported in Sheese et al. (2016) - although referred to older versions of the two datasets - is consistent with the one we measure. The difference we measure is varying significantly with latitudes and seasons: between -40° and $+40^\circ$ and below 70 km, there are peaks of -150% and -80% during

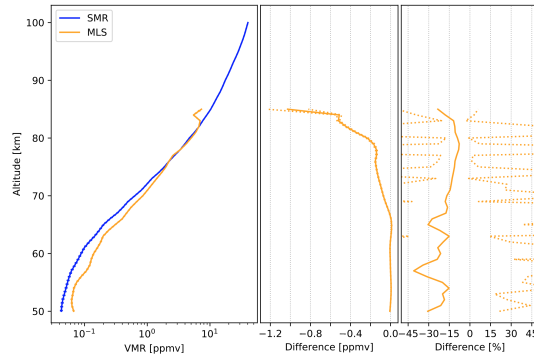


Figure 11. Comparison of SMR CO concentrations with the ones from MLS observations. The data plotted are global averages over the whole time between July 2004 and March 2019. Panels characteristics are the same as in Figure 9.

northern spring and autumn, respectively; while marked negative differences are registered during local summertime in both hemispheres, getting more pronounced toward high latitudes, especially around 50 km and 70 km altitude with peaks of -120%, as shown in Figure A5 (this is also observed in the ACE–MLS comparison in Sheese et al. (2016)).

5.4 OSO

- 5 Odin/SMR CO data are compared with data measured by a ground-based remote-sensing instrument at Onsala Space Observatory, OSO (57.4° N, 11.9° E), during 2002–2007. The recommended quality filtering has been followed.

The OSO instrument is a frequency-switched microwave radiometer for observations of the CO 1 → 0 transition at 115.27 GHz. During 2002–2007, a cooled Schottky single sideband mixer was used as the first stage. In 2014 the instrument was modified to a double sideband system with a low noise amplifier, at ambient temperature, as the first stage. During 2002–2007 a spec-

- 10 trometer with 20 MHz bandwidth and a resolution of 25 kHz was used. The Optimal Estimation Method together with the forward model ARTS, the Atmospheric Radiative Transfer Simulator (Eriksson et al., 2011), has been used to retrieve vertical CO profiles from the measured spectra. The OSO instrument, the calibration and retrieval methods are described in Forkman et al. (2012, 2016).

- 15 2002–2007 daily averages of CO profiles are retrieved from the OSO spectra. Only the 24h average CO spectra with a signal to noise ratio larger than 2 are used in the retrievals. SMR data are regarded coincident to OSO if the differences in latitude and longitude do not exceed $\pm 5^\circ$ and $\pm 10^\circ$ respectively and if the SMR data are taken within the OSO time average periods mentioned above. For the period 2002–2007, there are 89 coincident SMR profiles, see Figure 12.

Figure 13 shows the mean vertical CO profiles for the coincident SMR and OSO data sets. The limb sounding SMR has a much higher vertical resolution than the upward-looking OSO instrument. To compensate for this difference, the SMR profiles,

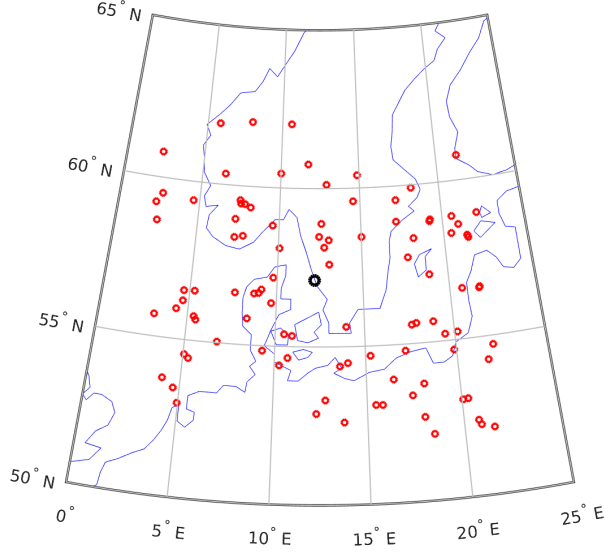


Figure 12. Coincidences Odin/SMR–OSO. Red circles show the 89 coincidences during 2002–2007 and the black circle shows the OSO site.

x_{sat} , were convoluted with the averaging kernels, A , of the OSO instrument (Rodgers, 2000).

$$x_s = x_a + A(x_{sat} - x_a) \quad (5)$$

where x_a is the OSO apriori and x_s the smoothed SMR profile. The measurement response of SMR is very close to 1 in the 50–100 km range. The altitude range in Figure 13 is more narrow since it is given by the range where the measurement response of the OSO instrument is > 0.75 .

OSO data from 2002–2007 have been compared to data from the satellite instruments ACE-FTS, MIPAS and MLS in a previous study (Forkman et al., 2012). As seen in Figure 13, during this period, the difference between SMR data and the coincident averaged data from OSO was found to be of 30% at 50 km, decreasing with altitude to reach -10% at 60 km and remaining constant up until 90 km altitude. The standard error of the mean, $\sigma_{\bar{x}}$, for the average difference between SMR and OSO, is about 5 % above 55 km, see Figure 13.

6 Conclusions

Before the application of the corrections described in this study, almost the whole Odin/SMR CO dataset was unusable due to line shifts and broadening of instrumental origin, due to the Phase-Lock Loop malfunctioning. Line displacement resulted in the failure of inversions or inaccurate retrievals, while the instrumental broadening caused underestimation of the concentration values. We estimated and corrected the different impacts of the PLL malfunctioning on SMR CO measurements. Line shifts were addressed by developing a correction algorithm which allowed the CO lines to be re-positioned to their theoretical center, also considering different frequency shifts for each tangent altitude within a scan. This resulted in the recovery of a great part of

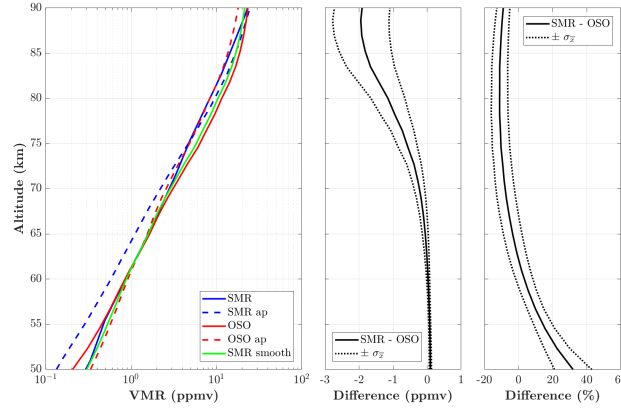


Figure 13. SMR–OSO comparison 2002–2007. Left: blue solid/dashed are the average SMR retrieved/apriori profiles, red solid/dashed are the average OSO retrieved/apriori profiles and green is the smoothed SMR profile. Middle and right: the average difference between the smoothed SMR and OSO profiles, solid, together with the standard error of the mean, $\sigma_{\bar{x}}$, dotted, are shown in ppmv and %, respectively.

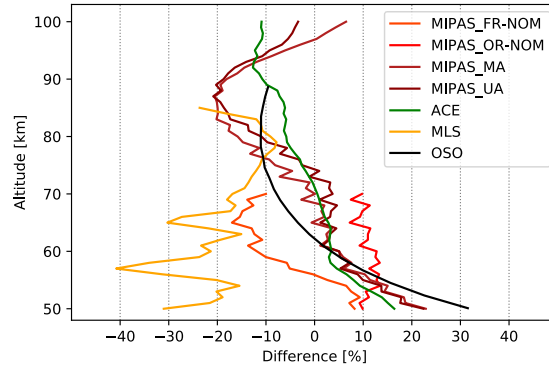


Figure 14. Summary of relative differences between SMR CO concentrations and the ones measured by all other instruments considered in this study. For the sake of clarity, errors are not shown.

the dataset. Line broadening was taken into account using a broader response function for the retrievals. That led to a new, good quality, Odin/SMR v3.0 mesospheric CO dataset, covering more than 18 years of observations. Time series of the retrieved volume mixing ratios reveal variations consistent with known mesospheric dynamical patterns, such as the annual cycle of the meridional circulation, SAO, SSWs, as well as a clear 11-year solar cycle signal. The validation study shows, on average, a good agreement with both the ground-based radiometer OSO and the three satellite-borne instruments (MIPAS, ACE-FTS, MLS) considered for comparison (see Figure 14). In particular, between 60 and 80 km SMR agrees very well with almost all instruments, presenting relative differences close to zero. Comparisons with MIPAS, ACE-FTS and OSO show a positive bias of SMR of up to +20% at low altitudes (50 - 60 km) and a negative bias of up to -20% at high altitudes (80 - 100 km).

Something different is found with regards to MLS - i.e. negative difference at all altitudes, ranging from -40% to -10% - which is in accordance with the stated MLS bias (Errera et al., 2019). To investigate if the reason for the observed differences is to be attributed to differences in vertical resolutions, in the case where SMR has better vertical resolution than the comparison instrument, we tried to smooth SMR profiles using Gaussian filters with FWHMs equal to the comparison instrument's vertical resolution; vice versa in the opposite case. This test didn't show any improvement in concentration differences (not shown), suggesting that differences in vertical resolution are not the cause of the observed bias. Another evidence for this, regarding comparisons with MIPAS modes, is given by the fact that comparisons of daytime measurements (which are characterised by similar vertical resolutions for SMR and MIPAS) do not show differences in concentration that vary from what is observed when comparing measurements all together. Moreover, systematic errors from non-LTE modelling in MIPAS measurements amount to less than 5% (Funke et al., 2007) and are therefore not thought to be the cause for the observed SMR-MIPAS differences at high altitudes. Thus, the causes for the observed concentration differences between SMR and the comparison instruments, both globally and for different latitudes and seasons, are unknown at the moment and require further investigation.

Given its unique extension in time and geographical coverage, this new mesospheric CO dataset provides a valuable tool for further studies of mesosphere dynamics.

15 Appendix A

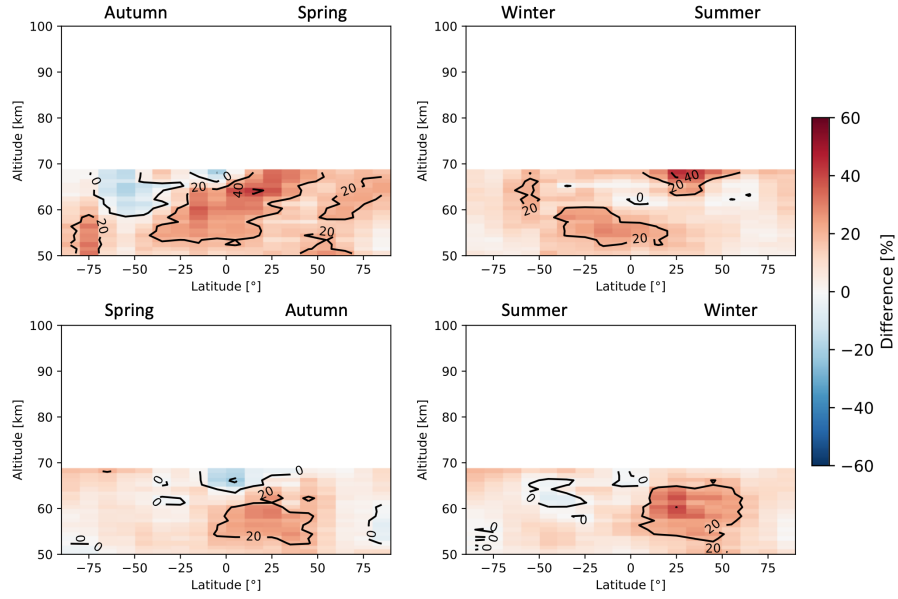


Figure A1. Seasonal zonal means of SMR-MIPAS Nominal-OR-NOM relative differences averaged over the time period indicated in Table 2. The seasons are intended as the time between solstice and equinox.

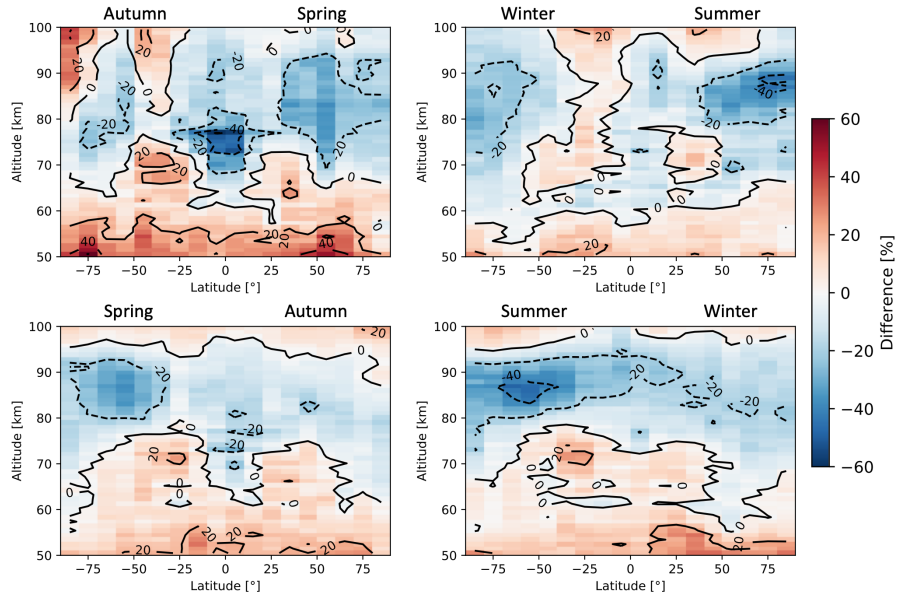


Figure A2. Seasonal zonal means of SMR-MIPAS Middle Atmosphere relative differences averaged over the time period indicated in Table 2. The seasons are intended as the time between solstice and equinox.

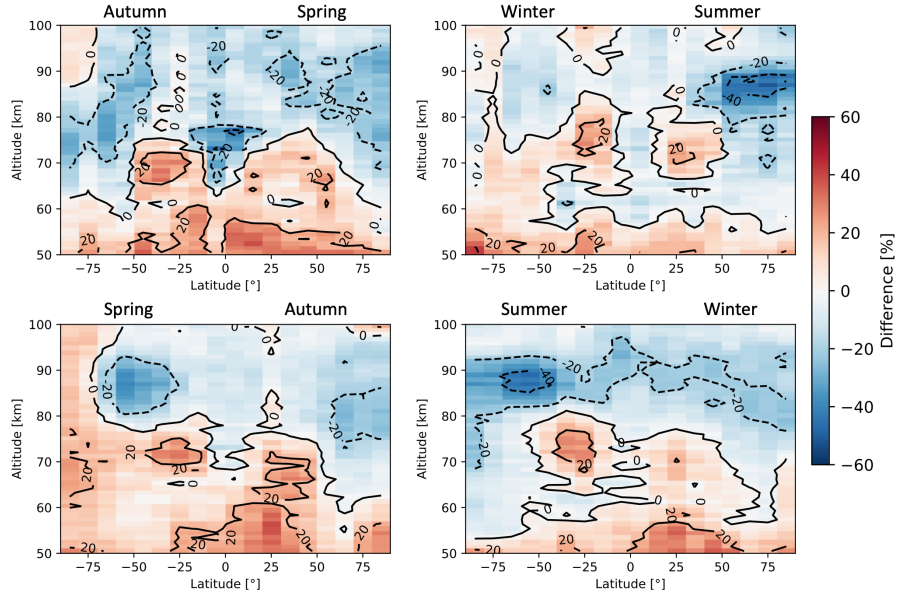


Figure A3. Seasonal zonal means of SMR–MIPAS Upper Atmosphere relative differences averaged over the time period indicated in Table 2. The seasons are intended as the time between solstice and equinox.

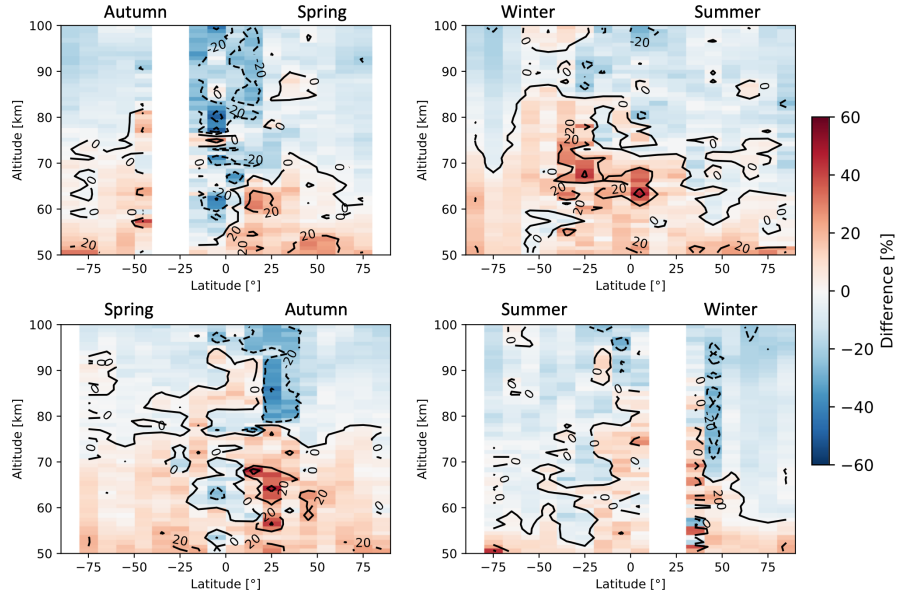


Figure A4. Seasonal zonal means of SMR–ACE relative differences averaged over the whole time between February 2004 and March 2019. The seasons are intended as the time between solstice and equinox.

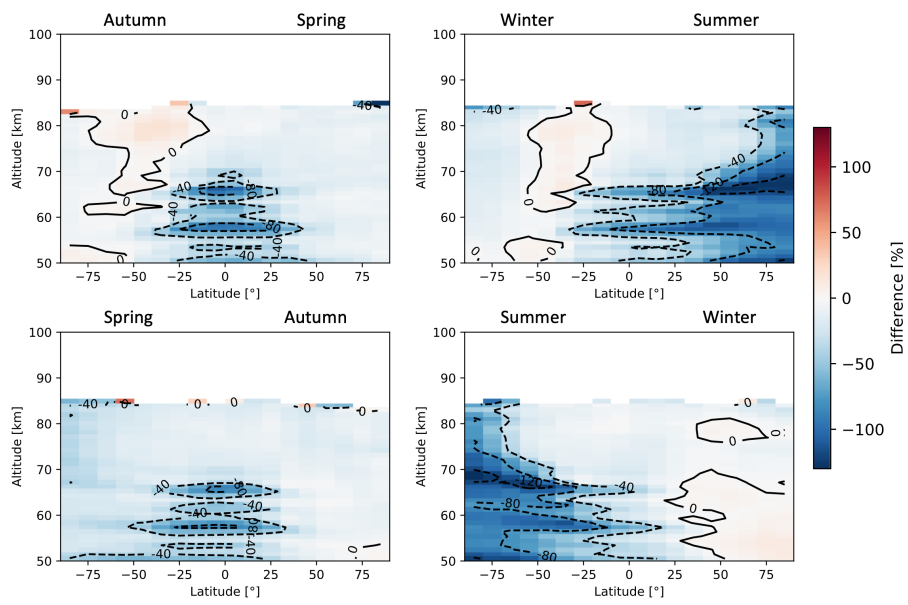


Figure A5. Seasonal zonal means of SMR–MLS relative differences averaged over the whole time between July 2004 and March 2019. The seasons are intended as the time between solstice and equinox. Note the different colour scale compared to the previous figures.

Data availability. Odin/SMR v3.0 L2 data are publicly accessible at <http://odin.rss.chalmers.se/level2>; MIPAS IMK/IAA L2 data (both NOM and MA/UA) can be downloaded upon registration at <http://www.imk-asf.kit.edu/english/308.php>; ACE-FTS L2 data are available upon request at <https://database.scisat.ca/l2signup.php>; MLS L2 data are available at <https://doi.org/10.5067/Aura/MLS/DATA2005> (Schwartz et al., 2015).

- 5 *Author contributions.* FG has developed the correction algorithm, processed the CO data, made most of the plots and written most of the text. KP, DM and PE have initiated the project and supported FG throughout it. KP has contributed to the writing of the text. PF has performed the comparison study between SMR and the OSO radiometer and has written the corresponding section. BR has helped with the processing of the data. All co-authors have contributed to the interpretation of the results and have proofread the text.

Competing interests. The authors declare that they have no conflict of interest.

- 10 *Acknowledgements.* The Chalmers team acknowledges support from the Swedish National Space Agency (Dnr 88/14 and 72/17). Odin is a Swedish-led satellite mission, which is also part of the European Space Agency's (ESA) third party mission programme. The reprocessing of the SMR data was supported by ESA (MesosphEO and Odin/SMR reprocessing projects). The authors would like to thank Julia Ringsby for her contribution to the correction algorithm. BF acknowledges support by the Spanish MCINN (ESP2017-87143-R) and EC FEDER funds. The Atmospheric Chemistry Experiment is a Canadian-led mission mainly supported by the Canadian Space Agency. MLS research
15 in Edinburgh was funded by NERC. [The authors also thank the two anonymous reviewers for their helpful comments.](#)

References

- Allen D.R., Stanford J.L., López-Valverde M.A., Nakamura N., Lary D.J., Douglass A.R., Cerniglia M.C., Remedios J.J., and Taylor F.W.: Observations of Middle Atmosphere CO from the UARS ISAMS during the Early Northern Winter 1991/92, *J. Atmos. Sci.*, 56, 563-583, doi:10.1175/1520-0469(1999)056%3C0563:OOMACF%3E2.0.CO;2, 1999.
- 5 Baron Ph., Ricaud Ph., de la Noë J., Eriksson J.E.P., Merino F., Ridal M., and Murtagh D.P.: Studies for the Odin sub-millimetre radiometer. II. Retrieval methodologies, *Canadian Journal of Physics*, 80, 341-356, doi:10.1139/p01-150, 2002.
- Barret B., Ricaud P., Santee M.L., Attié J.-L., Urban J., Le Flochmoën E., Berthet G., Murtagh D., Eriksson P., Jones A., de La Noë J., Dupuy E., Froidevaux L., Livesey N.J., Waters J.W., and Filipiak M.J.: Intercomparisons of trace gases profiles from the Odin/SMR and Aura/MLS limb sounders, *J. Geophys. Res.*, 111, D21302, doi:10.1029/2006JD007305, 2006.
- 10 Bernath P.F., McElroy C.T., Abrams M.C., Boone C.D., Butler M., Camy-Peyret C., Carleer M., Clerbaux C., Coheur P.-F., Colin R., DeCola P., DeMazière M., Drummond J.R., Dufour D., Evans W.F.J., Fast H., Fussen D., Gilbert K., Jennings D.E., Llewellyn E.J., Lowe R.P., Mahieu E., McConnell J.C., McHugh M., McLeod S.D., Michaud R., Midwinter C., Nassar R., Nichitiu F., Nowlan C., Rinsland C.P., Rochon Y.J., Rowlands N., Semeniuk K., Simon P., Skelton R., Sloan J.J., Soucy M.-A., Strong K., Tremblay P., Turnbull D., Walker K.A., Walkty I., Wardle D.A., Wehrle V., Zander R., and Zou J.: Atmospheric Chemistry Experiment (ACE): Mission overview, *Geophys. Res. Lett.*, 32, L15S01, doi:10.1029/2005GL022386, 2005.
- 15 Boone C.D., Nassar R., Walker K.A., Rochon Y., McLeod S.D., Rinsland C.P., and Bernath P.F.: Retrievals for the Atmospheric Chemistry Experiment Fourier-Transform Spectrometer, *Appl. Opt.*, 44, 7218– 7231, doi:10.1364/AO.44.007218, 2005.
- Boone, C. D., K. A. Walker, and P. F. Bernath: Version 3 Retrievals for the Atmospheric Chemistry Experiment Fourier Transform Spectrometer (ACE-FTS), *The Atmospheric Chemistry Experiment ACE at 10: A Solar Occultation Anthology*, pp. 103– 127 A. Deepak Publishing, Hampton, Va., 2013.
- 20 Brasseur G.P. and Solomon S.: *Aeronomy of the Middle Atmosphere*, Springer, The Netherlands, 3rd edition, 2005.
- Buehler S. A., Mendrok J., Eriksson P., Perrin A., Larsson R., and Lemke O.: ARTS, the Atmospheric Radiative Transfer Simulator – version 2.2, the planetary toolbox edition, *Geosci. Model Dev.*, 11, 1537–1556, doi:10.5194/gmd-11-1537-2018, 2018.
- Clancy R.T., Muhleman D.O., and Berge G.L.: Microwave spectra of terrestrial mesospheric CO, *J. Geophys. Res.*: 87, 5009, doi:10.1029/JC087iC07p05009, 1982.
- 25 Charlton A.J. and Polvani L.M.: A New Look at Stratospheric Sudden Warmings. Part I: Climatology and Modeling Benchmarks, *J. Climate*, 20, 449–469, doi:10.1175/JCLI3996.1, 2007.
- Clerbaux C., George M., Turquety S., Walker K.A., Barret B., Bernath P., Boone C., Borsdorff T., Cammas J.P., Catoire V., Coffey M., Coheur P.-F., Deeter M., De Mazière M., Drummond J., Duchatelet P., Dupuy E., de Zafra R., Eddounia F., Edwards D. P., Emmons L., Funke B., Gille J., Griffith D.W.T., Hannigan J., Hase F., Höpfner M., Jones N., Kagawa A., Kasai Y., Kramer I., Le Flochmoën E., Livesey N.J., López-Puertas M., Luo M., Mahieu E., Murtagh D., Nédélec P., Pazmino A., Pumphrey H., Ricaud P., Rinsland C.P., Robert C., Schneider M., Senten C., Stiller G., Strandberg A., Strong K., Sussmann R., Thouret V., Urban J., and Wiacek A.: CO measurements from the ACE-FTS satellite instrument: data analysis and validation using ground-based, airborne and spaceborne observations, *Atmos. Chem. Phys.*, 8, 2569–2594, doi:10.5194/acpd-7-15277-2007, 2008.
- 30 de Zafra R.L. and Muscari G.: CO as an important high-altitude tracer of dynamics in the polar stratosphere and mesosphere, *J. Geophys. Res.*, 109, D06105, doi:10.1029/2004JD005102, 2004.

- Dee, D.P., Uppala S.M., Simmons A.J., Berrisford P., Poli P., Kobayashi S., Andrae U., Balmaseda M.A., Balsamo G., Bauer P., Bechtold P., Beljaars A.C.M., van de Berg L., Bidlot J., Bormann N., Delsol C., Dragani R., Fuentes M., Geer A.J., Haimberger L., Healy S.B., Hersbach H., Hólm E.V., Isaksen I., Kållberg P., Köhler M., Matricardi M., McNally A.P., Monge-Sanz B.M., Morcrette J.-J., Park B.-K., Peubey C., de Rosnay P., Tavolato C., Thépaut J.-N. and Vitart F.: The ERA-Interim reanalysis: configuration and performance of the data assimilation system, *Q.J.R. Meteorol. Soc.*, 137: 553-597, doi:10.1002/qj.828, 2011.
- 5 Dupuy É., Urban J., Ricaud P., Le Flochmoën É., Lautié N., Murtagh D., De La Noë J., El Amraoui L., Eriksson P., Forkman P., Frisk U., Jégou F., Jiménez C., and Olberg M.: Strato-mesospheric measurements of carbon monoxide with the Odin Sub-Millimetre Radiometer: Retrieval and first results, *Geophys Res Lett*: 31, L20101, doi:10.1029/2004GL020558, 2004.
- Eriksson P., Jiménez C., and Buehler S.A.: Qpack, a general tool for instrument simulation and retrieval work, *J. Quant. Spectrosc. Ra.*, 91, 1, 47-64, doi:10.1016/j.jqsrt.2004.05.050, 2005.
- 10 Eriksson P., Ekström M., Melsheimer C., and Buehler S.: Efficient forward modelling by matrix representation of sensor responses, *International Journal of Remote Sensing*, 27, 9, 1793-1808, doi:10.1080/01431160500447254, 2006.
- Eriksson P., Buehler S.A., Davis C.P., Emde C., and Lemke O.: ARTS, the atmospheric radiative transfer simulator, Version 2, *J. Quant. Spectrosc. Ra.*, Vol. 112, p. 1551-1558, doi:10.1016/j.jqsrt.2011.03.001, 2011.
- 15 Errera Q., Chabrillat S., Christophe Y., Deboscher J., Hubert D., Lahoz W., Santee M.L., Shiotani M., Skachko S., von Clarmann T., and Walker K.: Technical note: Reanalysis of Aura MLS chemical observations, *Atmos. Chem. Phys.*, 19, 13647–13679, doi:10.5194/acp-19-13647-2019, 2019.
- Forkman P., Christensen O.M., Eriksson P., Urban J., and Funke B.: Six years of mesospheric CO estimated from ground-based frequency-switched microwave radiometry at 57° N compared with satellite instruments, *Atmos. Meas. Tech.*, 5, 2827–2841, doi:10.5194/amt-5-2827-2012, 2012.
- 20 Forkman P., Christensen O.M., Eriksson P., Billade B., Vassilev V., and Shulga V.M.: A compact receiver system for simultaneous measurements of mesospheric CO and O₃, *Geosci. Instrum. Method. Data Syst.*, 5, 27–44, doi:10.5194/gi-5-27-2016, 2016.
- Frisk U., Hagström M., Ala-Laurinaho J., Andersson S., Berges J.-C., Chabaud J.-P., Dahlgren M., Emrich A., Florén H.-G., Florin G., Fredrixon M., Gaier T., Haas R., Hirvonen T., Hjalmarsson Å., Jakobsson B., Jukkala P., Kildal P. S., Kollberg E., Lassing J., Lecacheux A., Lehtikoinen P., Lehto A., Mallat J., Marty C., Michet D., Narbonne J., Nexon M., Olberg M., Olofsson A. O. H., Olofsson G., Origné A., Petersson M., Piironen P., Pons R., Pouliquen D., Ristorcelli I., Rosolen C., Rouaix G., Räisänen A. V., Serra G., Sjöberg F., Stenmark L., Torchinsky S., Tuovinen J., Ullberg C., Vinterhav E., Wadefalk N., Zirath H., Zimmermann P., and Zimmermann R.: The Odin satellite I. Radiometer design and test, *A&A*, 402(3):L27-L34, doi:10.1051/0004-6361:20030335, 2003.
- 30 Froidevaux L., Livesey N.J., Read W.G., Jiang Y.B., Jimenez C., Filipiak M.J., Schwartz M.J., Santee M.L., Pumphrey H.C., Jiang J.H., Wu D.L., Manney G.L., Drouin B.J., Waters J.W., Fetzer E.J., Bernath P.F., Boone C.D., Walker K.A., Jucks K.W., Toon G.C., Margitan J.J., Sen B., Webster C.R., Christensen L.E., Elkins J.W., Atlas E., Lueb R.A., and Hendershot R.: Early validation analyses of atmospheric profiles from EOS MLS on the aura Satellite, *IEEE Trans. Geosci. Rem. Sens.*, 44, 1106 – 1121, doi:10.1109/TGRS.2006.864366, 2006.
- 35 Funke B., López-Puertas M., Bermejo-Pantaleón D., von Clarmann T., Stiller G.P., Höpfner M., Grabowski U., and Kaufmann M.: Analysis of nonlocal thermodynamic equilibrium CO 4.7 μ m fundamental, isotopic, and hot band emissions measured by the Michelson Interferometer for Passive Atmospheric Sounding on Envisat, *J. Geophys. Res.*, 112, D11305, doi:10.1029/2006JD007933, 2007.
- Funke B., López-Puertas M., Garcia-Comas M., Stiller G., von Clarmann T., Höpfner M., Glatthor N., Grabowski U., Kellmann S., and Linden A.: Carbon monoxide distributions from the upper troposphere to the mesosphere inferred from 4.7 μ m non-local thermal equilibrium emissions measured by MIPAS on Envisat, *Atmos. Chem. Phys.*, 9, 2387-2411, doi:10.5194/acp-9-2387-2009, 2009.

- Garcia R.R., López-Puertas M., Funke B., Marsh D.R., Kinnison D.E., Smith A.K., and González-Galindo F.: On the distribution of CO₂ and CO in the mesosphere and lower thermosphere, *J. Geophys. Res. Atmos.*, 119, 5700–5718, doi:10.1002/2013JD021208, 2014.
- Garcia R.R., López-Puertas M., Funke B., Kinnison D.E., Marsh D.R., and Qian L.: On the secular trend of CO_x and CO₂ in the lower thermosphere, *J. Geophys. Res. Atmos.*, 121, 3634–3644, doi:10.1002/2015JD024553, 2016.
- 5 [Hamilton K., MIDDLE ATMOSPHERE I Semiannual Oscillation, Encyclopedia of Atmospheric Sciences \(Second Edition\), Academic Press, pp. 26-29, doi:10.1016/B978-0-12-382225-3.00233-4, 2015.](#)
- Karlsson B., Gumbel J., Stegman J., Lautier N., Murtagh D., and Odin Team: Studies of Noctilucent Clouds by the Odin Satellite, in: Proceedings of the 35th COSPAR Scientific Assembly, 35, 1921, 2004.
- MIPAS-IMK/IAA L2 Data ReadMe, FMI-TN-MesosphEO-WP4-003, Version 1.0, 2017.
- 10 Lee J.N., Wu D.L., Manney G.L., Schwartz M.J., Lambert A., Livesey N.J., Minschwaner K.R., Pumphrey H.C., and Read W.G.: Aura Microwave Limb Sounder observations of the polar middle atmosphere: Dynamics and transport of CO and H₂O, *J. Geophys. Res.*, 116, D05110, doi:10.1029/2010JD014608, 2011.
- Lee J.N., Wu D.L., Ruzmaikin A., and Fontenla J.: Solar cycle variations in mesospheric carbon monoxide, *Journal of Atmospheric and Solar-Terrestrial Physics*, 170, 21-34, doi:10.1016/j.jastp.2018.02.001, 2018.
- 15 Livesey N.J., Read W.G., Wagner P.A., Froidevaux L., Lambert A., Manney G.L., Millán Valle L.F., Pumphrey H.C., Santee M.L., Schwartz M.J., Wang S., Fuller R.A., Jarnot R.F., Knosp B.W., Martinez E., and Lay R.R.: Earth Observing System (EOS) Aura Microwave Limb Sounder (MLS) Version 4.2x Level 2 data quality and description document, Tech. Rep. D-33509 Rev. D, JPL, 2018.
- [Manney G.L., Harwood R.S., MacKenzie I.A., Minschwaner K., Allen D.R., Santee M.L., Walker K.A., Hegglin M.I., Lambert A., Pumphrey H.C., Bernath P.F., Boone C.D., Schwartz M.J., Livesey N.J., Daffer W.H., and Fuller R.A.: Satellite observations and modeling of transport in the upper troposphere through the lower mesosphere during the 2006 major stratospheric sudden warming, Atmos. Chem. Phys., 9, 4775–4795, doi:10.5194/acp-9-4775-2009, 2009.](#)
- 20 Minschwaner K., Manney G. L., Livesey N. J., Pumphrey H. C., Pickett H. M., Froidevaux L., Lambert A., Schwartz M. J., Bernath P. F., Walker K. A., and Boone C. D.: The photochemistry of carbon monoxide in the stratosphere and mesosphere evaluated from observations by the MicrowaveLimb Sounder on the Aura satellite, *J. Geophys. Res.*, 115, D13303, doi:10.1029/2009JD012654, 2010.
- 25 Oelhaf H.: MIPAS Mission Plan, Issue 4, Version 3, ESA Technical Note ENVI-SPPA-EOPG-TN-07-0073, 2008.
- Orsolini Y.J., Limpasuvan V., Pérot K., Espy P., Hibbins R., Lossow S., Raaholt Larsson K., and Murtagh D.: Modelling the descent of nitric oxide during the elevated stratopause event of January 2013, *Journal of Atmospheric and Solar-Terrestrial Physics*, 155, 50-61, doi:10.1016/j.jastp.2017.01.006, 2017.
- Pérot K., Urban J., and Murtagh D.: Unusually strong nitric oxide descent in the Arctic middle atmosphere in early 2013 as observed by Odin/SMR, *Atmos. Chem. Phys.*, 14, 8009-8015, doi:10.5194/acpd-14-3563-2014, 2014.
- 30 [Picone J.M., Hedin A.E., Drob D.P., and Aikin A.C.: NRLMSISE-00 empirical model of the atmosphere: Statistical comparisons and scientific issues, J. Geophys. Res., 107\(A12\), 1468, doi:10.1029/2002JA009430, 2002.](#)
- Pumphrey H.C., Filipiak M.J., Livesey N.J., Schwartz M.J., Boone C., Walker K.A., Bernath P., Ricaud P., Barret B., Clerbaux C., Jarnot R.F., Manney G.L., and Waters J.W., Validation of middle-atmosphere carbon monoxide retrievals from the Microwave Limb Sounder on Aura, *J. Geophys. Res.*, 112, D24S38, doi:10.1029/2007JD008723, 2007.
- 35 Randall C.E., Rusch D.W., Bevilacqua R.M., Hoppel K.W., Lumpe J.D., Shettle E., Thompson E., Deaver L., Zawodny J., Kyrö E., Johnson B., Kelder H., Dorokhov V.M., König-Langlo G., and Gil M.: Validation of POAM III ozone: comparison with ozonesonde and satellite data, *J. Geophys. Res.*, 108(D12), 4367, doi:10.1029/2002JD002944, 2003.

- Rodgers C.D.: Inverse methods for atmospheric sounding: Theory and practise, World Scientific Publishing, 1st edition, 2000.
- Rothman L.S., Gordon I.E., Babikov Y., Barbe A., Chris Benner D., Bernath P.F., Birk M., Bizzocchi L., Boudon V., Brown L.R., Campargue A., Chance K., Cohen E.A., Coudert L.H., Devi V.M., Drouin B.J., Fayt A., Flaud J.-M., Gamache R.R., Harrison J.J., Hartmann J.-M., Hill C., Hodges J.T., Jacquemart D., Jolly A., Lamouroux J., Le Roy R.J., Li G., Long D.A., Lyulin O.M., Mackie C.J., Massie S.T., Mikhailenko S., Müller H.S.P., Naumenko O.V., Nikitin A.V., Orphal J., Perevalov V., Perrin A., Polovtseva E.R., Richard C., Smith M.A.H., Starikova E., Sung K., Tashkun S., Tennyson J., Toon G.C., Tyuterev V.I., and Wagner G.: The HITRAN2012 molecular spectroscopic database, *J. Quant. Spectrosc. Ra.*, 130, 4-50, doi:10.1016/j.jqsrt.2013.07.002, 2013.
- Rydberg B., Eriksson P., Kiviranta J., Ringsby J., Skyman A., and Murtagh D.: Odin/SMR Algorithm Theoretical Basis Document: Level 1 Processing, Technical Report, Chalmers University of Technology, Department of Space, Earth and Environment, <http://odin.rss.chalmers.se/#documents>, 2017.
- Schoeberl M.R., Douglass A.R., Hilsenrath E., Bhartia P.K., Beer R., Waters J.W., Gunson M.R., Froidevaux L., Gille J.C., Barnett J.J., Levelt P.F., and DeCola P.: Overview of the EOS Aura Mission, *IEEE Trans. Geosci. Remote Sensing*, 44(5):1066 - 1074, doi:10.1109/TGRS.2005.861950, 2006.
- Schwartz M., Pumphrey H., Livesey N., and Read W.: MLS/Aura Level 2 Carbon Monoxide (CO) Mixing Ratio V004, Green- belt, MD, USA, Goddard Earth Sciences Data and Information Services Center (GES DISC), available at: <https://doi.org/10.5067/Aura/MLS/DATA2005>, 2015.
- Sheese P.E., Boone C.D., and Walker K.A.: Detecting physically unrealistic outliers in ACE-FTS atmospheric measurements, *Atmos. Meas. Tech.*, 8, 741–750, doi:10.5194/amt-8-741-2015, 2015.
- Sheese P.E., Walker K.A., Boone C.D., Bernath P.F., Froidevaux L., Funke B., Raspollini P., and von Clarmann T.: ACE-FTS ozone, water vapour, nitrous oxide, nitric acid, and carbon monoxide profile intercomparisons with MIPAS and MLS, *J. Quant. Spectrosc. Ra.*, 186, 63–80, doi:10.1016/j.jqsrt.2016.06.026, 2016.
- Solomon S., Garcia R.R., Olivero J.J., Bevilacqua R.M., Schwartz P.R., Clancy R.T., and Muhleman D.O.: Photochemistry and Transport of Carbon Monoxide in the Middle Atmosphere, *J. Atmos. Sci.*, 42, 10721083, doi:10.1175/1520-0469(1985)042%3C1072:PATOCM%3E2.0.CO;2, 1985.
- Vignon E. and Mitchell D.M.: The stratopause evolution during different types of sudden stratospheric warming event, *Clim Dyn*, 44: 3323, doi:10.1007/s00382-014-2292-4, 2015.
- Waters J.W., Froidevaux L., Jarnot R.F., Read W.G., Pickett H.M., Harwood R.S., Cofield R.E., Filipiak M.J., Flower D.A., Livesey N.J., Manney G.L., Pumphrey H.C., Santee M.L., Siegel P.H., and Wu D.L.: An Overview of the EOS MLS Experiment, version 2.0, JPL D-15745, ATBD-MLS-01, 2004.
- Zander R., Leclercq H., and Kaplan L.D.: Concentration of carbon monoxide in the upper stratosphere, *Geophys Res Lett*: 8, 365, doi:10.1029/GL008i004p0036, 1981.