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Validation of ACE-FTS version 3.5 NO_y species profiles using correlative satellite measurements

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Abstract. The ACE-FTS (Atmospheric Chemistry Experiment – Fourier Transform Spectrometer) instrument on the Canadian SCISAT satellite, which has been in operation for over 12 years, has the capability of deriving stratospheric profiles of many of the NO_y (N + NO + NO₂ + NO₃ + 2 × N₂O₅ + HNO₃ + HNO₄ + ClONO₂ + BrONO₂) species. Version 2.2 of ACE-FTS NO, NO₂, HNO₃, N₂O₅, and ClONO₂ has previously been validated, and this study compares the most recent version (v3.5) of these five ACE-FTS prod-

ucts to spatially and temporally coincident measurements from other satellite instruments – GOMOS, HALOE, MAESTRO, MIPAS, MLS, OSIRIS, POAM III, SAGE III, SCIAMACHY, SMILES, and SMR. For each ACE-FTS measurement, a photochemical box model was used to simulate the diurnal variations of the NO_y species and the ACE-FTS measurements were scaled to the local times of the coincident measurements. The comparisons for all five species show good agreement with correlative satellite measurements. For

NO in the altitude range of 25–50 km, ACE-FTS typically agrees with correlative data to within –10 %. Instrument-averaged mean relative differences are approximately –10 % at 30–40 km for NO₂, within ±7 % at 8–30 km for HNO₃, better than –7 % at 21–34 km for local morning N₂O₅, and better than –8 % at 21–34 km for ClONO₂. Where possible, the variations in the mean differences due to changes in the comparison local time and latitude are also discussed.

1 Introduction

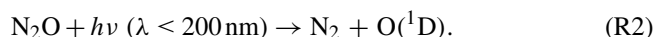
Currently, the only way to get global observational coverage of the Earth's atmosphere is with satellite-based observations. In addition, no single instrument can give us the full picture. Several instruments are needed in order to give us full global, vertical, and temporal coverage. Understanding biases between instruments is thus critical to understanding the true state of the atmosphere.

NO_y is the complete set of reactive nitrogen species. Its concentration is calculated as [N] + [NO] + [NO₂] + [NO₃] + 2 × [N₂O₅] + [HNO₃] + [HNO₄] + [ClONO₂] + [BrONO₂]. The abundances of NO_y as well as the partitioning and interactions of its components are important to understand because they play a significant role in ozone chemistry. The main source of the NO_y species in the stratosphere is through oxidation of N₂O. NO and NO₂ can also descend from the lower thermosphere, where they are mainly produced via energetic particle precipitation, into the upper stratosphere during the polar winter (Randall et al., 1998, 2007, 2009; Funke et al., 2005a). A detailed description of stratospheric NO_y photochemistry is given, for example, by Brasseur and Solomon (2005) and a summary for the species validated in this study is given below.

The main source of NO in the stratosphere is through dissociation of N₂O via reactions with excited O(¹D) atoms,



and the majority of stratospheric N₂O originates from surface soil and ocean emissions. The predominant destruction mechanism of stratospheric N₂O is photolysis, producing O(¹D) in the process:



NO is also produced through reactions of O₂ with atomic nitrogen, which can be produced by dissociation of N₂ by cosmic rays. Cosmic rays can be a nontrivial source of NO in the polar regions.

Stratospheric NO₂ is produced through the reaction of NO with O₃,

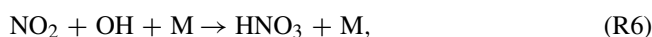


as well as with ClO, BrO, HO₂, and CH₃O₂. NO₂ is destroyed through reactions with atomic oxygen and through

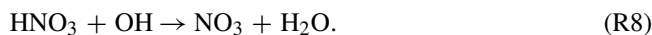
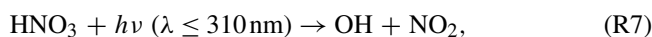
photolysis – both processes produce NO:



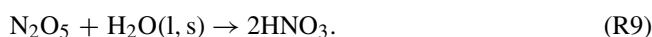
The main source of HNO₃ is through the three-body reaction,



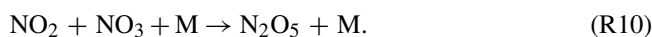
where M is an air molecule. The main sinks are through photolysis and through destruction via reactions with OH:



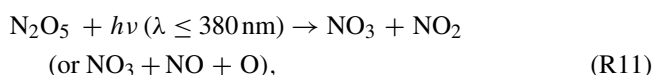
HNO₃ is also produced on the surface of ice (H₂O) particles, water droplets, nitric acid ice, and sulfate aerosols through the heterogeneous reaction,



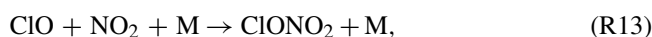
N₂O₅ is produced mainly at night, when there is an abundance of NO₃, through the three-body reaction:



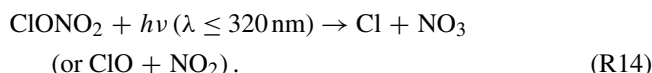
The main sinks of N₂O₅ are through photolysis and through collisions:



The main source of ClONO₂ is through the three-body reaction,



and the main sink is through photolysis,



Concentrations of these NO_y species can have large diurnal variations because the reactions governing their production and destruction depend on sunlight. To account for diurnal variations, calculations made using the “Pratmo” photochemical box model (McLinden et al., 2002) are used to scale local times between the two instruments. This model was used by Kerzenmacher et al. (2008) in the validation of version 2.2 of the ACE-FTS (Atmospheric Chemistry Experiment – Fourier Transform Spectrometer) NO and NO₂ data.

ACE-FTS level 2 version 3.5 data are used in comparison to coincident level 2 data from the satellite instruments Global Ozone Monitoring by Occultation of Stars (GOMOS), Halogen Occultation Experiment (HALOE), Measurements of Aerosol Extinction in the Stratosphere and Troposphere Retrieved by Occultation (MAESTRO), Michelson

Table 1. Details of the instruments and the NO_y data sets used in the comparisons.

Instrument data set	Observation method	Comparison period	Comparison coverage	Vertical resolution (km)				
				NO	NO ₂	HNO ₃	N ₂ O ₅	ClONO ₂
ACE-FTS v3.5	Solar occultation	2004–2013	85° S–87° N	3–4	3–4	3–4	3–4	3–4
GOMOS v6.01	Stellar occultation	2004–2012	60° S–60° N	–	2	–	–	–
HALOE v19	Solar occultation	2004–2005	23–68° S 48–67° N	4–7	2	–	–	–
MAESTRO v1.2	Solar occultation	2004–2010	85° S–87° N	–	1–2	–	–	–
MIPAS ESA v6	Limb emission	2005–2012	86° S–89° N	–	3–5	2–5	3–5	2–8
MIPAS IMK-IAA v5R (L1)	Limb emission	2005–2012	86° S–89° N	4–6	3–5	2–8	2–6	2–8
MLS v3.3	Limb emission	2004–2013	82° S–82° N	–	–	2–4	–	–
OSIRIS v3	Limb scatter	2004–2013	83° S–82° N	–	2.5	–	–	–
POAM III v4	Solar occultation	2004–2005	63–86° S 56–70° N	–	1–3	–	–	–
SAGE III v4	Solar occultation	2004–2005	38–60° S 50–81° N	–	1–4	–	–	–
SCIAMACHY v3.1	Limb scatter	2004–2012	81° S–84° N	–	3–10	–	–	–
SMILES v3.0	Limb emission	2009–2010	66° S–66° N	–	–	5–9	–	–
SMR v2.0	Limb emission	2004–2009	84° S–85° N	–	–	1.5–3	–	–

Table 2. Reported retrieval uncertainties for the data sets used in this study. The listed ACE-FTS values represent mean statistical fitting errors (Boone et al., 2005). The values given are in the altitude range of 20–60 km for NO, 20–40 km for NO₂, 15–30 km for HNO₃, 20–40 km for N₂O₅, and 17–38 km for ClONO₂. The MIPAS IMK-IAA uncertainties were obtained from the respective validation studies discussed in Sect. 2.2.2. Note that these are the uncertainties reported as “systematic” and “random” uncertainties and are not all necessarily at the same confidence level.

Instrument data set	NO		NO ₂		HNO ₃		N ₂ O ₅		ClONO ₂		Reference
	Sys. (%)	Rand. (%)	Sys. (%)	Rand. (%)	Sys. (%)	Rand. (%)	Sys. (%)	Rand. (%)	Sys. (%)	Rand. (%)	
ACE-FTS v3.5	15–80		2–5		<5		7–50		6–29		
GOMOS v6.01	–	–	1–5	10–25	–	–	–	–	–	–	Verronen et al. (2009)
HALOE v19	10–60 ^a		5–30 ^a		–	–	–	–	–	–	Gordley et al. (1996)
MAESTRO v1.2	–	–	5–10	<5	–	–	–	–	–	–	Kar et al. (2007)
MIPAS ESA v6	–	–	<9	<15	5–10	2–10	11–42 ^b		6–60 ^b		Raspollini et al. (2006)
MIPAS IMK-IAA v5R (L1)	5–40 ^a		10–40 ^a		5–15	2–6	10–45	5–30	7–32 ^a		See caption
MLS v3.3	–	–	–	–	5–10	<10	–	–	–	–	Livesey et al. (2013)
OSIRIS v3	–	–	11–31	6–15	–	–	–	–	–	–	Brohede et al. (2007a)
POAM III v4	–	–	0–12	2–7	–	–	–	–	–	–	Randall et al. (2002)
SAGE III v4	–	–	Not given	15	–	–	–	–	–	–	SAGE (2002b)
SCIAMACHY v3.1	–	–	<14	<15	–	–	–	–	–	–	Bauer et al. (2012)
SMILES v3.0	–	–	–	–	Not given	15–80	–	–	–	–	Kikuchi et al. (2010)
SMR v2.0	–	–	–	–	<10	10–15	–	–	–	–	Urban et al. (2009)

^a Values represent the total uncertainty. ^b Values calculated from the error profiles given in the respective level 2 data set.

Interferometer for Passive Atmospheric Sounding (MIPAS), Microwave Limb Sounder (MLS), Optical Spectrograph and InfraRed Imaging System (OSIRIS), Polar Ozone and Aerosol Measurement (POAM) III, Stratospheric Aerosol and Gas Experiment (SAGE) III, SCanning Imaging Absorption SpectroMeter for Atmospheric CHartography (SCIAMACHY), Superconducting Submillimeter-Wave Limb-Emission Sounder (SMILES), and Sub-Millimetre Radiometer (SMR). These instruments and their data sets are described in the next section and their key details and ran-

dom/systematic uncertainties are outlined in Tables 1 and 2, respectively. Section 2 outlines the instruments and the data sets used in this study. Section 3 describes the methodology as well as the Prato photochemical box model. The results of the comparisons with ACE-FTS, with and without the use of the photochemical box model, are detailed in Sect. 4. A summary and discussion of the results is given in Sect. 5.

2 Instrumentation

2.1 Instruments on SCISAT

2.1.1 ACE-FTS

The ACE-FTS instrument (Bernath et al., 2005) is a solar occultation, high-resolution (0.02 cm^{-1}) spectrometer operating between 750 and 4400 cm^{-1} . It was launched in August 2003 into a high-inclination orbit of 74° near an altitude of 650 km, and ACE-FTS has been providing volume mixing ratio (VMR) profiles of over 30 atmospheric trace gases and of over 20 isotopologue species since February 2004. During either sunset or sunrise, ACE-FTS makes a measurement approximately every 2 s between ~ 5 and 150 km with a vertical sampling between ~ 2 and 6 km, depending on the orbital geometry. The vertical extent of the instrument field of view is ~ 3 –4 km at the tangent point.

The trace species VMR retrieval, as described by Boone et al. (2005), is a nonlinear, least-squares, global-fitting technique that fits the observed spectra in given spectral microwindows (dependent on the retrieved species) to forward modelled spectra. Modelled spectra use line strengths and widths from HITRAN 2004 (Rothman et al., 2005) (with various updates, as detailed by Boone et al., 2013) and use the derived temperature and pressure profiles determined by fitting CO₂ lines in the observed spectra. The main updates in v3.5 (compared to v2.2) are improved sets of microwindows for the majority of species, along with an increase in the number of interfering species in their retrievals; improved temperature/pressure retrievals resulting in a reduction of profiles exhibiting unrealistic temperature oscillations; and the inclusion of trace species COCl₂, COClF, H₂CO, CH₃OH, and HCFC-141b and the exclusion of ClO.

The ACE-FTS v3.5 NO retrieval uses 39 microwindows between 1649.3 and 1977.6 cm^{-1} . The main interfering species within the NO microwindows is O₃, but spectral features of CO₂ and H₂O isotopologues and COF₂ are also present. The retrieval has a lower altitude limit of 6 km and an upper altitude limit of 107 km. ACE-FTS v2.2 NO was validated by Kerzenmacher et al. (2008), and there were two known issues with the v2.2 results (still present in the v3.5 NO results). At altitudes below ~ 20 km, NO VMRs suffer from a significant negative bias that causes many unphysical negative results. This is most likely due to strong diurnal variation along the line of sight that is not taken into account in the NO retrievals. Also, in polar winter around 35–50 km, where the NO VMR profile has a large vertical gradient, during times of increased downwelling, NO VMRs can exhibit large negative spikes. Kerzenmacher et al. (2008) found that, on average, ACE-FTS v2.2 NO agreed with coincident HALOE data on the order of 8 % within the altitude range of 22–64 km and exhibited a positive bias of ~ 10 % from 93 to 110 km, and that the uncertainties were too large

for statistically significant comparisons in the 64–93 km region.

The v3.5 NO₂ retrieval uses 40 microwindows between 1204.4 and 2950.9 cm^{-1} . The majority of microwindows added since v2.2 were chosen because of their information content with respect to the spectrally interfering isotopologues of CH₄ and H₂O. Between 7 and 20 km, CO₂ and OCS also significantly interfere with the NO₂ lines. The retrieval has a lower altitude limit of 7 km and an upper altitude limit of 52 km. ACE-FTS v2.2 NO₂ was validated by Kerzenmacher et al. (2008), who concluded that ACE-FTS NO₂ typically exhibited a ~ 15 % low bias with coincident satellite data near the peak (~ 35 km) and on average was within 20 % in the altitude range of approximately 20–40 km.

The v3.5 HNO₃ retrieval uses 41 microwindows between 865.5 and 1977.6 cm^{-1} . Interfering species include CCl₂F₂, H₂O, CO₂, OCS, and O₃. The retrieval has a lower altitude limit of 5 km and an upper altitude limit of 62 km. ACE-FTS v2.2 HNO₃ was validated by Wolff et al. (2008), who found that the ACE-FTS data and all coincident satellite data agreed to within 20 % in the altitude range of 18–35 km.

The v3.5 N₂O₅ retrieval, with altitude limits of 8 and 45 km, has only one spectral window, 30.0 cm^{-1} wide and centred at 1244.0 cm^{-1} . Interfering species include O₃ and isotopologues of H₂O, CO₂, CH₄, and N₂O. ACE-FTS v2.2 update N₂O₅ profiles (herein v2.2) were compared with MIPAS IMK-IAA N₂O₅ profiles by Wolff et al. (2008), who used climatological results from a chemical transport model to calculate diurnal scaling factors in order to match the local times of the two instruments. Without the use of diurnal scaling, Wolff et al. (2008) found that ACE-FTS v2.2 N₂O₅ typically exhibited a low bias on the order of 30–50 %, whereas with diurnal scaling ACE-FTS typically exhibited a ~ 10 –35 % low bias.

The v3.5 ClONO₂ retrieval uses five microwindows between 780.2 and 2672.7 cm^{-1} . Interfering species include N₂O, CH₄, O₃, HNO₃, and isotopologues of N₂O, CO₂, H₂O, and CH₄. The retrieval has a lower altitude limit of 10 km and an upper altitude limit of 41 km at high latitudes and 36 km near the equator. ACE-FTS v2.2 ClONO₂ was compared to co-located MIPAS IMK-IAA data by Wolff et al. (2008), who used diurnal scaling factors to match the local times of the two instruments. With the use of diurnal scaling, Wolff et al. (2008) showed that ACE-FTS v2.2 and MIPAS IMK-IAA ClONO₂ values typically differed by less than 1 % between 16 and 24 km. Above the peak (~ 25 km), ACE-FTS exhibited a positive bias with respect to MIPAS of up to 20 % near 33 km.

It should be noted that ACE-FTS also derives VMR profiles of HNO₄; however, because HNO₄ does not contribute substantially to the overall NO_y budget and due to a lack of multiple correlative satellite data sets with which to validate, it is not included in this study. All ACE-FTS data used in this study were screened for physically unrealistic outliers using the recommended quality flags version 1.1, as described by

Sheese et al. (2015). Any profile known to be affected by instrument or processing errors (flag values of 7) or any profile containing a data point determined to be an extreme outlier (flag value in the range of 4–6) was excluded from the analysis.

2.1.2 MAESTRO

The MAESTRO instrument (McElroy et al., 2007) consists of two spectrographs designed to cover the spectral range 210–1025 nm – with a 1.5–2.5 nm spectral resolution, which observe direct solar radiation occulted by the Earth's atmosphere. The MAESTRO solar occultation measurements are used to retrieve profiles of O₃, NO₂, H₂O, aerosol extinction, and other various atmospheric properties. The instrument has been providing measurements since February 2004. The NO₂ retrieval algorithm, described by McElroy et al. (2007), uses a two-step process. The spectral fitting of apparent optical depth spectra is used to derive slant column densities, assuming temperature-independent NO₂ and O₃ absorption cross sections from Burrows et al. (1998, 1999). Then an iterative Chahine inversion technique (Chahine, 1968) is used to retrieve NO₂ VMR profiles from the slant column values. The spectral fitting algorithm is performed over a spectral range of 420–750 nm, and NO₂ profiles are retrieved in an altitude range of ~5–52 km, with a vertical resolution on the order of 1–2 km.

Version 1.2 of the NO₂ data (used in this study) was validated by Kerzenmacher et al. (2008), who found that between 25 and 40 km, when comparing to correlative satellite measurements, diurnally scaled MAESTRO NO₂ tends to exhibit a bias within –20 and +10%. In the same altitude region, scaled MAESTRO NO₂ also tends to exhibit a high bias of 0–50% when compared to correlative ground- and balloon-based measurements. The poorer comparison with ground-based instruments was attributed to not accounting for diurnal variations along the MAESTRO line of sight in the NO₂ retrieval algorithm. It should be noted that this issue would similarly affect ACE-FTS NO₂ retrievals.

The ACE-FTS outlier detection method described by Sheese et al. (2015) was used to detect physically unrealistic outliers in the MAESTRO NO₂ data set. Any profile that was found to contain such an outlier was rejected prior to any comparisons. This method was ineffective at removing many of the outliers below 19 km. Therefore at altitude levels below 19 km, NO₂ VMR values greater than 3 ppb were screened out. At all altitude levels, any values with a corresponding fractional error of 1 or greater were also removed. Only data between February 2004 and September 2010 were used in the analysis.

2.2 Instruments on Envisat

In March 2002, the European Space Agency (ESA) launched the Envisat satellite (Fischer et al., 2008) into a polar, sun-

synchronous orbit near 800 km, with an ascending node of 22:00 LT (local time). On board the Envisat satellite were a number of atmospheric sounding instruments, including the limb sounders GOMOS, MIPAS, and SCIAMACHY, which are described in following sections. Ground control lost communication with the satellite in early April 2012, thus ending all observations from the Envisat instruments.

2.2.1 GOMOS

The GOMOS instrument (Kyrölä et al., 2004) on the Envisat satellite employed a grating spectrometer that observed the attenuation of stellar emission, from the ultraviolet (UV) to the near-infrared, through the limb of the Earth's atmosphere. The stellar occultation technique was employed to retrieve vertical profiles of nighttime O₃, NO₂, NO₃, H₂O, OClO, BrO, O₂, and aerosol extinction nominally between altitudes of 5 and 150 km, using three different bands within the spectral range of 248–954 nm. GOMOS was capable of obtaining hundreds of occultations each day with a vertical sampling typically between 0.4 and 1.7 km. GOMOS measurements span from March 2002 to April 2012.

The NO₂ retrieval algorithm is described by Kyrölä et al. (2010) and makes use of a Tikhonov-type regularization (Tikhonov, 1963), which leads to a retrieval vertical resolution of 4 km. Version 6 of the GOMOS NO₂ data set is used in this study. Version 5 of the NO₂ retrievals was validated by Verronen et al. (2009), who compared the GOMOS profiles to nighttime MIPAS ESA NO₂ data (described below). It was found that in the low to midlatitudes, between approximately 25 and 60 km, GOMOS NO₂ tended to exhibit a positive bias with respect to MIPAS on the order of 0–25%. In the high latitudes, the two data sets agreed within 35% at altitudes above ~45 km where nighttime NO₂ VMR was at a maximum. However, at lower altitudes (in the high latitude regions) the bias reached up to 65%, which was greater than the combined systematic errors. Since the ACE-FTS NO₂ profiles only extend up to 52 km, GOMOS comparisons have been limited to between 60° S and 60° N.

Only GOMOS profiles where the local solar zenith angle is greater than 97° at altitudes below 50 km and greater than 110° at altitudes below 100 km were used in the analysis. In order to eliminate the presence of extreme outliers, any GOMOS NO₂ profile that contained an absolute VMR value greater than 0.5 ppm in the altitude range of 0–52 km was also rejected; in the limited latitude region this rejected less than 1% of the GOMOS profiles.

2.2.2 MIPAS

The MIPAS instrument (Fischer and Oelhaf, 1996; Fischer et al., 2008) on the Envisat satellite was a limb-viewing Fourier transform spectrometer that observed atmospheric emissions. The spectrometer had five spectral bands in the range of 685–2410 cm⁻¹ and scanned the Earth's limb be-

tween altitudes from approximately 6 to 70 km in nominal mode and up to 170 km in special modes. The MIPAS vertical field of view was 3 km and the instrument had a vertical sampling that ranged from 1.5 to 5 km, depending on the altitude. Prior to 2005, MIPAS operated at its full spectral resolution of 0.025 cm⁻¹, with a sampling time of 4.5 s. In 2004, an anomaly occurred in the interferometer mirror slide mechanism and it was determined that the spectral resolution needed to be downgraded to 0.0625 cm⁻¹ with a consequent reduction of the sampling time to 1.8 s, exploited to allow for a finer vertical sampling. In order to avoid any discontinuities that may arise from switching the observation mode, only MIPAS measurements from the period of January 2005 to April 2012 were used in this study.

Two different MIPAS level 2 products, based on two different retrieval algorithms, were used in this study – the first is from the ESA and the second is the result of a collaboration between the Institute of Meteorology and Climate Research at the Karlsruhe Institute of Technology and the Instituto de Astrofísica de Andalucía (IMK-IAA). The ESA algorithm that produces version 6 of the level 2 retrievals (used in this study) is described by Raspollini et al. (2013). It is a least-squares, global-fitting technique, using the regularized Levenberg–Marquardt method (Hanke, 1997), which fits spectra in species-dependent microwindows to a forward model. A parameter setting has been chosen that leaves results largely independent from the initial guess profiles. The forward model assumes horizontal homogeneity and local thermodynamic equilibrium at all altitudes. An a posteriori regularization, using a self-adapting regularization constraint, is then applied to the retrieved profile (Ceccherini, 2005; Ceccherini et al., 2007).

The IMK-IAA algorithm is described by von Clarmann et al. (2009) and Funke et al. (2014), and the most recent version of the level 2 data (used in this study) is version 5. The IMK-IAA algorithm uses an iterative variant of Tikhonov regularization (Tikhonov, 1963) on species-dependent sets of microwindows. This inversion technique is implemented to constrain the shape of the resulting profile without pushing the values towards an a priori profile. The retrieval is performed on a 1 km grid, and the altitude-dependent strength of the smoothing constraint was chosen in order to optimize vertical resolution in the upper troposphere to lower mesosphere while still minimizing artificial oscillations in the retrieved profile. The NO and NO₂ retrievals are performed in log(VMR) space, and the forward model allows for horizontal variation in temperature. In the forward model, NO and NO₂ line-of-sight variations are considered and a line-of-sight NO_x gradient is retrieved concurrently. Further, the forward model can allow for deviations from local thermodynamic equilibrium (LTE), which mainly affects mesospheric retrievals, and LTE is assumed for all NO_y species except NO and NO₂. The NO_y microwindows are chosen, in part, in order to reduce non-LTE effects.

MIPAS IMK-IAA NO_x retrievals (only in the original resolution mode) were compared to HALOE measurements by Funke et al. (2005b). It was found that the two NO_x data sets typically agreed within 20 % between 25 and 50 km. Wetzel et al. (2007) found that, in the mid-stratospheric MIPAS ESA version 4.6 NO₂, diurnally scaled using data from a 1-D photochemical model agreed best with balloon-borne measurements, with biases typically better than 10 %. In similar comparisons with correlative satellite-based solar occultation measurements, the MIPAS ESA profiles typically agreed within 10–30 %.

Wang et al. (2007a, b) assessed the quality of the MIPAS IMK-IAA version 3 and MIPAS ESA version 4.6 HNO₃ data sets, respectively. Comparing MIPAS ESA HNO₃ with correlative data sets from ground-based and balloon-borne instruments, both Wang et al. (2007a, b) studies determined that relative differences were typically better than 10 %. In their comparisons with ACE-FTS v2.2 HNO₃, relative differences in the lower to mid-stratosphere were on the order of 5–15 %.

MIPAS IMK-IAA ClONO₂ profiles were validated by Höpfner et al. (2007), who showed that the MIPAS data set agreed well with correlative balloon and airborne data sets, typically to better than 10 %. Höpfner et al. (2007) also compared the MIPAS IMK-IAA profiles to ACE-FTS v2.2 ClONO₂ using diurnal correction factors obtained from a chemical transport model. The diurnally corrected MIPAS data and ACE-FTS typically agreed within 10 % at altitudes between 15 and 27 km. However, above 27 km, the ACE-FTS exhibited a ~ 20 % low bias with the diurnally corrected MIPAS data and a ~ 20 % high bias with the uncorrected data.

Neither the MIPAS ESA nor IMK-IAA N₂O₅ data set has been the focus of a MIPAS validation study, but MIPAS ESA N₂O₅ and ClONO₂ data were compared with the balloon-based MIPAS-B instrument by Wetzel et al. (2013). It was found that N₂O₅ concentrations typically agree within ±40 % and ClONO₂ concentrations typically within ±30 %. Also, the IMK-IAA N₂O₅ data set was used in the ACE-FTS v2.2 N₂O₅ validation study of Wolff et al. (2008), the results of which are summarized in Sect. 2.1.1.

All MIPAS vertical resolutions, listed in Table 1, were calculated as the full-width, half-maximum of the retrieval averaging kernels. MIPAS IMK-IAA data were used only where the corresponding averaging kernel diagonal values were greater than 0.03.

2.2.3 SCIAMACHY

The SCIAMACHY instrument (Burrows et al., 1995; Bovensmann et al., 1999) was an eight-channel grating spectrometer that observed the Earth's atmosphere in the wide spectral range of 240–2400 nm, using three different viewing geometries – limb viewing of scattered sunlight, solar occultation, and nadir viewing. The NO₂ data used in this study are the profiles retrieved from limb-viewing observa-

tions in the channel that observed in the spectral window of 394–620 nm (spectral channel 3). The instrument scanned the Earth's limb from the surface up to 100 km with a 2.5 km vertical field of view and a ~ 3 km vertical sampling. The NO₂ retrieval algorithm, detailed by Rozanov et al. (2005) and summarized by Bauer et al. (2012), uses limb-scattered radiances measured from 420 to 470 nm and solves the inverse problem using the DOAS technique and Tikhonov regularization (Tikhonov, 1963). In each profile, the spectra are normalized by the limb radiances nearest 43 km. The regularization matrix smooths the retrievals using an empirically determined height-dependent smoothing parameter, chosen in order to minimize physically unrealistic oscillations in profiles while maximizing vertical resolution. The retrieval makes use of a forward model that takes into account absorption by O₃ (simultaneously retrieved) and O₂–O₂ and uses pressure and temperature profiles from the European Centre for Medium-Range Weather Forecasts (ECMWF). The NO₂ and O₃ absorption cross sections were obtained from Bogumil et al. (1999). The algorithm retrieves NO₂ profiles between 10 and 40 km with a typical vertical resolution of 3–5 km, degrading to ~ 10 km at the upper and lower retrieval altitude limits.

This study used v3.1 of the SCIAMACHY level 2 NO₂ profiles, which was validated by Bauer et al. (2012). The NO₂ profiles were compared to correlative satellite measurements that were diurnally scaled to the SCIAMACHY local times. It was found that in the altitude range of 25–35 km SCIAMACHY NO₂ tends to exhibit a 2% low bias with respect to HALOE v19 profiles and tends to exhibit a 5% high bias with respect to ACE-FTS v2.2 profiles.

Only SCIAMACHY data below 40 km with a retrieval response greater than 0.8 were used in the analysis.

2.3 Solar occultation instruments

2.3.1 HALOE on the Upper Atmosphere Research Satellite (UARS)

The HALOE instrument (Russell et al., 1993), on the UARS, was a solar occultation instrument that provided observations of the Earth's limb between October 1991 and November 2005. The UARS precessing orbit allowed for HALOE measurements to observe all latitudes between 80° S and 80° N approximately every 36 days. Profiles of O₃, HCl, HF, CH₄, H₂O, NO, NO₂ concentrations, temperature, and aerosols were derived from observations within four radiometric channels and four radiometric/gas-filter correlation channels.

The HALOE NO measurements use a gas-filter correlation method with a spectral filter band pass near 1900 cm⁻¹ and are virtually insensitive to interfering absorbers. The NO₂ measurements are made using a broadband radiometric channel centred near 1600 cm⁻¹ and the effects of interfering species O₂, H₂O, and CH₄ are accounted for in

the retrieval. The interfering species N₂O is not accounted for, although the effect on NO₂ is very small. Retrievals of NO and NO₂ profiles use a modified onion peel approach and account for aerosol extinction and interfering attenuation. The NO retrievals have a vertical resolution of 4 km at altitudes below ~ 60 km (degrading to 7 km at higher altitudes), and the NO₂ retrievals have a vertical resolution of 2 km. The HALOE version 17 NO and NO₂ data were validated by Gordley et al. (1996). This study uses HALOE version 19 NO and NO₂, which have very small differences relative to v17 (James M. Russell III, Hampton University, personal communication, December 2015). Gordley et al. (1996) found that above 25 km HALOE v17 NO tended to agree with correlative satellite and balloon-based measurements within 15%, but with a maximum low bias reaching 35%. Also, above 25 km HALOE v17 NO₂ agreed with correlative satellite, balloon, and ground-based measurements to within 15%.

2.3.2 POAM III on SPOT 4

The POAM III instrument (Lucke et al., 1999) was a nine-channel photometer that viewed the Earth's limb in solar occultation. POAM III, on board the Satellite Pour l'Observation de la Terre (SPOT) 4 satellite, was launched in March 1998 into a sun-synchronous orbit with a descending node of 10:30 LT, at an altitude of ~ 830 km and a 98.7° inclination. Designed to measure atmospheric profiles of O₃, H₂O, NO₂, and aerosol extinction, POAM III observed the limb at tangent heights between cloud-top and 60 km in nine different narrow passbands in the near-UV to near-infrared spectral region, with a total spectral range from 354 to 1018 nm. POAM III started taking measurements in April 1998, and measurements stopped in December 2005 due to instrument failure.

NO₂ profiles were retrieved between 20 and 45 km from differential measurements in the 439.6 nm (NO₂ “on”) and 442.2 nm (NO₂ “off”) channels, both with a full-width, half-maximum passband of 2.1 nm. The vertical resolution of retrieved NO₂ was ~ 1.5 km from 25 to 35 km, increasing to nearly 3 km at 20 km and > 7 km at 45 km. The retrieval algorithm is described in detail by Lumpe et al. (2002). The algorithm inverts slant column densities to vertical profiles using the Newtonian optimal estimation technique (Rodgers, 2008) for all target species. The forward model assumes horizontal homogeneity. Randall et al. (2002) validated POAM III version 3.0 NO₂ measurements through comparisons with data from multiple instruments. They found no evidence for any systematic bias below 35 km; e.g. differences with respect to HALOE were within approximately ± 0.2 ppbv ($\sim 6\%$). Relative to HALOE, POAM III NO₂ mixing ratios were shown to be higher by up to 0.7 ppbv ($\sim 17\%$) from 35 to 42 km; about 5% of that bias was attributed to an error in HALOE retrievals, but no explanation for the remaining 12% was identified. Although the version 4 NO₂ data (used in

this study) have not been the focus of a validation study, it was used by Kerzenmacher et al. (2008) in comparison with ACE NO₂. It was shown that above 25 km, POAM III typically agreed within $\pm 6\%$ with respect to ACE-FTS v2.2 and within $\pm 8\%$ with respect to MAESTRO v1.2.

2.3.3 SAGE III on Meteor 3M

The SAGE III instrument (SAGE III ATBD Team, 2002a) was a solar and lunar occultation atmospheric sounder on board the Russian Meteor 3M satellite, which was launched in December 2001 and was operational until March 2006. Meteor 3M was launched into a 1020 km altitude, sun-synchronous orbit with a descending node of $\sim 09:00$ LT. In solar occultation mode, SAGE III was designed to retrieve vertical profiles of O₃, NO₂, H₂O, and aerosol extinction (plus NO₃ and OCIO in lunar mode) throughout the stratosphere from observations in the near-UV to near-infrared spectral region. The instrument consisted of a grating spectrometer that observed in the spectral range of 280–1040 nm and an InGaAs infrared detector that observed in a band pass between 1530 and 1560 nm.

The SAGE III NO₂ retrieval algorithm is detailed by SAGE III ATBD Team (2002b). The algorithm first uses a multiple linear regression technique to derive slant column densities for both O₃ and NO₂ simultaneously from calculated slant column optical depths. The O₃ and NO₂ region wavelength-dependent optical depths are derived from observations in two spectral channels spanning 433–450 and 563–622 nm. The NO₂ column densities are inverted into vertical density profiles (on a 0.5 km grid between 0 and 100 km with a vertical resolution of 1–2 km) using a modified Chahine technique (Chahine, 1968), assuming horizontal homogeneity.

There has not yet been a rigorous SAGE III NO₂ validation study. Kar et al. (2007) found that SAGE III NO₂ version 3 data (used in this study) typically exhibited a high bias (within ~ 10 – 15%) above 25 km with respect to v1.2 MAESTRO data. Similarly, Kerzenmacher et al. (2008) found that the SAGE III v3 data also tended to exhibit a high bias (typically within $\sim 10\%$) with respect to v2.2 ACE-FTS data. These results are consistent with Polyakov et al. (2005), who reported that their SAGE III NO₂ product, derived using the Newtonian iterative optimal estimation technique, was systematically lower than the SAGE III operational product.

2.4 MLS on Aura

The MLS instrument (Waters et al., 2006) aboard the Aura satellite observes atmospheric thermal emission in the Earth's limb. It was launched into a sun-synchronous orbit at an altitude of ~ 700 km and with an ascending node of 13:45 LT. The MLS consists of seven radiometers measuring in the spectral range of 118 GHz to 2.5 THz, and the spectra are used to retrieve atmospheric profiles of temper-

ature, geopotential height, and concentrations of over 15 atmospheric trace species and cloud ice on a pressure vertical grid.

HNO₃ retrieved from MLS is scientifically useful between pressure limits of 215 and 1.5 hPa. In the lower altitude range, at pressures of 22 hPa or greater, the 240 GHz radiometer measurements are used and result in a HNO₃ vertical resolution on the order of 3–4 km; at higher altitudes, at pressures of 15 hPa or less, the HNO₃ retrievals use measurements from the 190 GHz radiometer and have a vertical resolution of 4–6 km. In both pressure regimes, HNO₃ level 2 v3.3/3.4 profiles (Livesey et al., 2006, 2013) use a Newtonian optimal estimation technique (Rodgers, 2008), with a forward model that assumes horizontal homogeneity and uses absorption cross sections from the JPL Spectral Line Catalogue (Pickett et al., 1998).

Version 2.2 HNO₃ was validated by Santee et al. (2007), where the MLS data were compared to multiple data sets retrieved from ground-based, balloon-borne, aircraft, and satellite platforms. It was found that the MLS HNO₃ profiles were scientifically useful within the altitude range of approximately 10–40 km and that throughout the stratosphere MLS HNO₃ tended to exhibit a low bias on the order of 10–30%. That low bias was largely eliminated in version 3.3 (Livesey et al., 2013).

All MLS measurements with corresponding negative precision values, indicating poor retrieval response, have not been included in the analyses, nor have any profiles determined to contain cloud contamination. However, the adverse effects on MLS v3 HNO₃ due to clouds were substantially mitigated in the most recent version, v4.2 (Livesey et al., 2015). The altitude-dependent vertical resolution was assumed to be constant for all retrievals and was calculated as the full-width, half-maximum of the mean averaging kernels.

2.5 The Odin satellite

Odin is a Swedish/Canadian/Finnish/French satellite (Murtagh et al., 2002) that was launched in February 2001. It was launched into a sun-synchronous orbit at an altitude of ~ 600 km, with an ascending/descending node of 06:00/18:00 LT. Aboard the Odin satellite are two Earth observing instruments, OSIRIS (Llewellyn et al., 2004) and SMR (Frisk et al., 2003).

2.5.1 OSIRIS

The optical spectrograph of the OSIRIS instrument operates in the spectral range of 280–810 nm, with ~ 1 nm spectral resolution, and observes Rayleigh and Mie scattered sunlight in the Earth's limb between altitudes of ~ 7 and 110 km with a vertical field of view of approximately 1 km. The NO₂ retrievals, described by Haley and Brohede (2007), use the DOAS technique to calculate NO₂ slant columns. These are calculated in the spectral window of 435–451 nm and

between altitudes of 10 and 46 km, with the OSIRIS 46–60 km averaged radiances as the reference spectrum. The slant columns are then inverted into density profiles using the optimal estimation technique (Rodgers, 2008), using LIMBTRAN (Griffioen and Oikarinen, 2000) for the forward model. The NO₂ retrievals have a vertical resolution of approximately 2 km at all altitudes.

Version 3 of the data set (used in this study) was validated by Brohede et al. (2007a), who found that OSIRIS NO₂ typically agrees with correlative satellite and balloon-borne data sets within 20 % between 25 and 35 km for all seasons and latitudes. Between 35 and 45 km, the agreement was within 30 %, with smaller absolute systematic differences for comparisons in the high latitudes than for those nearer the equator.

2.5.2 SMR

SMR observes thermal emission in the Earth's limb using four tunable receivers in the spectral range of 486–581 GHz and a millimetre-wave receiver near 119 GHz. The HNO₃ profile retrieval algorithm (Urban et al., 2005) uses observations in a 1 GHz band centred at 544.6 GHz and is based on the Newtonian Levenberg–Marquardt optimal estimation technique (Rodgers, 2008). The forward model used is that of the MOLIERE-5 forward/inversion model (Urban et al., 2004). HNO₃ is retrieved at altitudes above 18 km, with vertical resolutions on the order of 2–3 km. As discussed by Urban et al. (2009), the SMR HNO₃ data exhibit a ~1–1.5 km vertical bias. Therefore, in this study the version 2.1 HNO₃ data were offset upwards by 1.5 km prior to any analysis.

Urban et al. (2009) showed that the SMR HNO₃ climatology exhibits reasonably good agreement with UARS/MLS climatology from measurements taken between 1991 and 1998. Wolff et al. (2008) showed that SMR HNO₃ profiles exhibit a ~20 % high bias with respect to ACE-FTS v2.2 HNO₃ at altitudes below 30 km, and exhibit systematic differences within ±20 % between 30 and 35 km.

Only profiles that had retrieval response values greater than 0.75 were used in the analysis. Due to a level 2 processing error that affected SMR data for May 2009 and onwards, only SMR data before May 2009 were used in this study.

2.6 SMILES on the Japanese Experiment Module (JEM) on the International Space Station (ISS)

The SMILES instrument (Kikuchi et al., 2010) was an atmospheric limb sounder that operated on ISS/JEM between October 2009 and April 2010. SMILES measured atmospheric thermal emissions in three bands within the spectral region of 624–650 GHz. The ISS orbits the Earth at an altitude of ~375 km with an inclination of 52°. In order to observe northern high latitudes, the SMILES line of sight was angled 45° from the ISS orbital plane, giving SMILES a nominal latitudinal coverage of 38° S to 66° N. The angle of the line of

sight was occasionally shifted to give a latitudinal coverage of 66° S to 38° N. SMILES scanned the Earth's limb between tangent heights of 10 and 60 km with a vertical resolution on the order of 3.5–4 km, and the local time coverage was such that it took 2 months to sample an entire diurnal cycle.

The SMILES operational retrieval algorithm, detailed by Takahashi et al. (2010), makes use of the optimal estimation technique combined with the Levenberg–Marquardt method, with a forward model that accounts for instrument attributes, single-ray temperature brightness, and absorption cross sections from the JPL Spectral Line Catalogue (Pickett et al., 1998). The resulting HNO₃ data, derived from observations in the two spectral bands covering 624.32–625.52 (band A) and 649.12–650.32 (band C) GHz, have a typical vertical resolution on the order of 5–9 km.

No studies focusing specifically on SMILES-derived HNO₃ have previously been published, mainly because the line parameters used in the forward model are theoretical, rather than laboratory, values. This study uses version 2.4 of the level 2 SMILES data from the operational processor. Only level 2 SMILES data derived from band C measurements were used in the analysis, as the HNO₃ retrievals from band A have been found to typically converge to a priori values (Makato Suzuki, personal communication, 30 October 2015). Only data with corresponding precision values greater than 0 (indicating reasonable measurement response values) were used in the analysis.

3 Methodology

In this section, when discussing comparisons between the ACE-FTS data set and the correlative data sets from other instruments, the term INST will be used to refer in general to one of the other instruments' data sets. Prior to analysis, all profiles (from every data set) have been linearly interpolated onto the ACE-FTS 1 km grid. In cases where an ACE-FTS profile was coincident with multiple profiles within an INST data set, only the profile measured closest in time to the ACE-FTS occultation was used.

In order to keep the level of vertical smoothing consistent between data sets, vertical resolution matching was carried out on coincident profiles where the INST vertical resolutions are finer than 3 km or coarser than 4 km (the range of the ACE-FTS vertical resolution). The profile with the finer vertical resolution, X_f , was smoothed by taking a weighted average of the profile at each altitude level. The weight used was a normalized Gaussian centred at the altitude level:

$$X_f^{\text{smooth}}(h) = \frac{\int X_f(z) G(h, z) dz}{\int G(h, z) dz}, \quad (1)$$

where h is the altitude on the ACE-FTS 1 km grid, z is altitude, and $G(h, z)$ is the normalized Gaussian distribution,

$$G(h, z) = \frac{2\sqrt{2\ln 2}}{v_s(h)\sqrt{2\pi}} \exp\left(-\frac{1}{2} \frac{(z-h)^2}{(v_s(h)/2\sqrt{2\ln 2})^2}\right). \quad (2)$$

v_s is the square root of the difference between the squared coarser vertical resolution of profile X_c and the squared vertical resolution of X_f (in order to avoid over-smoothing):

$$v_s(h) = \sqrt{v_c^2 - v_f^2}, \quad (3)$$

where v_c and v_f are the vertical resolutions of X_c and X_f , respectively, at altitude level h . In instances where the INST vertical resolution is finer than 3 km in one altitude region and coarser than 4 km in another region, the INST profiles are smoothed only in the altitude regions where $v_{\text{INST}}(h) < 3$ km, the ACE-FTS profiles are smoothed only in the altitude regions where $v_{\text{INST}}(h) > 4$ km, and neither is smoothed in the intermediate regions.

This method of using a Gaussian as an approximation of an averaging kernel is used in place of applying the averaging kernels directly because averaging kernels are not always available for all data sets. In fact, the ACE-FTS data sets do not include corresponding averaging kernels. One drawback of this approach is that any distortion of the profiles due to asymmetric averaging kernels (especially for retrievals performed in log(VMR) space) remains unaccounted for. However, as discussed in the Appendix, vertical smoothing in the altitude regions where the ACE-FTS retrievals have been validated typically only affect average relative differences on the order of 1 % or less.

For all of the species analyzed, three main diagnostics have been calculated at each altitude: correlation, mean relative difference, and standard deviation of relative differences. In all comparisons, differences are with respect to ACE-FTS v3.5 data. In the following definitions, X will represent ACE-FTS values at a given height, and Y will represent the corresponding INST values. The correlation coefficient, r , for comparisons between ACE-FTS and the other individual correlative data sets is determined at each height in the usual way:

$$r = \frac{1}{n-1} \sum_i^n \left(\frac{X_i - \bar{X}}{\sigma_X} \right) \left(\frac{Y_i - \bar{Y}}{\sigma_Y} \right), \quad (4)$$

where n is the number of co-located measurements and σ refers to the standard deviation over the co-located measurements. The means of the relative differences are calculated at each altitude as the mean of the absolute differences (relative to ACE-FTS) divided by the mean of both the ACE-FTS and INST values:

$$\text{rel diff} = 2 \frac{\sum_i^n X_i - Y_i}{\sum_i^n X_i + Y_i} \times 100 \%. \quad (5)$$

The overall mean is used as the denominator because the ACE-FTS retrievals, along with certain INST retrievals, allow for negative concentrations (which are included in the analysis so as to not bias the respective means); negative values can cause unrealistically large percent differences if the average of two compared values is near zero. The relative difference calculated as per Eq. (5) can also have unrealistically large values when the overall mean is near zero (if one of the ACE-FTS or INST averages is negative); however, this is much more unlikely than when using the standard calculation of the percent difference. Similarly, the standard deviation of the relative differences is calculated at each height as the standard deviation of the absolute differences (relative to ACE-FTS) divided by the overall mean of the ACE-FTS and INST values.

When comparing ACE-FTS data to multiple instruments it is desirable to calculate an overall average of each of the diagnostic values. A simple mean of the values is not useful, as it does not take into account the quality of the INST data sets used in the comparisons. Therefore, a weighted average is calculated, using the inverse of the squared standard error of the relative means ($\sigma_s^{-2} = (\sigma/\sqrt{n})^{-2}$) as the weight. Using σ_s assumes that all data sets exhibit similar natural variability. In certain regions, it is possible for comparisons to have unreasonable standard errors with data set values approximately equal to the a priori. Unfortunately, not all data sets include retrieval response and, therefore, at each height the weights are calculated as the INST inverse-squared standard error multiplied by the INST correlation coefficient, i.e.

$$W_{\text{INST}} = \frac{r_{\text{INST}}}{\sigma_s^2}. \quad (6)$$

For rare cases where there is anti-correlation between ACE-FTS and INST ($r_{\text{INST}} < 0$), the weights are set to zero. These weights are used to calculate the weighted-average ACE-FTS correlation coefficients, mean differences, and standard deviations of the relative differences.

All recommended status, quality, and convergence flags have been applied to all data sets where such flags have been made available (as described in Sect. 2).

Diurnal scaling

For each pair of coincident profiles, the ACE-FTS profile was scaled to the local time of the other instrument's profile. This was done by using a photochemical box model in order to determine altitude-dependent diurnal-scale factors for each ACE-FTS NO_y profile. Similar approaches have been used before in other studies, e.g. Bracher et al. (2005), Fussen et al. (2005), Wetzel et al. (2007), Brohede et al. (2007a), and Wolff et al. (2008).

The University of California Irvine photochemical box model (Prather, 1997; McLinden et al., 2002), also known as Pratmo, simulates the diurnal cycle of nitrogen and chlorine species, including NO, NO₂, HNO₃, N₂O₅, and ClONO₂. It

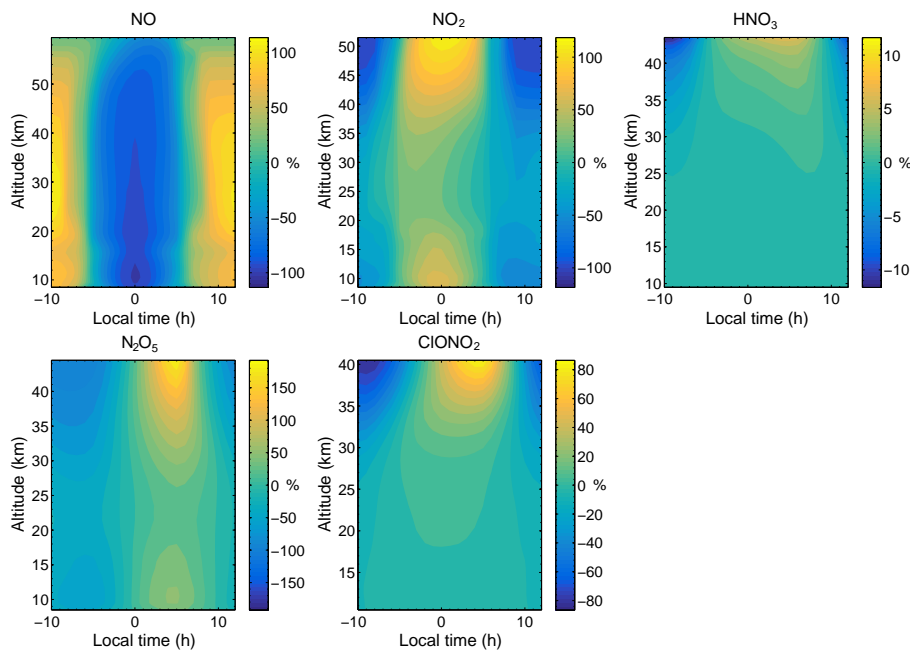


Figure 1. Diurnal variation in percent for NO, NO₂, HNO₃, N₂O₅, and ClONO₂ profiles, calculated by the Prtmo model for all ACE-FTS v3.5 data. At each altitude, variation is the percent deviation from the mean VMR at that altitude.

was used by Brohede et al. (2007b) in producing NO₂ climatologies from OSIRIS measurements, by Kerzenmacher et al. (2008) in the validation of ACE-FTS v2.2 NO₂ and by Bauer et al. (2012) in the validation of SCIAMACHY v3.1 NO₂. In the simulation of the diurnal cycle for an ACE-FTS profile, the model is constrained using the corresponding ACE-FTS temperature, pressure, and O₃ profiles. The model takes into account altitude, latitude, and day of year, using NO_y and N₂O climatologies from a 3-D chemical transport model (Olsen et al., 2001), Cl_y and Br_y climatologies (as described by Brohede et al., 2007a), and climatological SAGE II background aerosol data. All photochemical reaction rates were obtained from Sander et al. (2003). Updated reaction coefficients have more recently been suggested for Reactions (R6) and (R10) by Burkholder et al. (2015). Since HNO₃ does not have a significant diurnal variation, excluding the updated coefficients for Reaction (R6) is unlikely to affect the results of this study; however, excluding updates to the coefficients for Reaction (R10) may add additional uncertainty to the comparisons of N₂O₅ that use diurnal scaling. Latitude- and longitude-dependent albedo values from the Medium Resolution Imaging Spectrometer (MERIS) 412 nm albedo climatology (Popp et al., 2011) were also used as input into the model. The MERIS albedo climatology data were obtained from http://www.temis.nl/surface/meris_bsa.html. Figure 1 shows the mean altitude-dependent diurnal variations for NO, NO₂, HNO₃, N₂O₅, and ClONO₂, as calculated by Prtmo using all ACE-FTS v3.5 data. The variation values shown at a given altitude are the mean percent deviations from the mean concentration at that altitude.

The output of the Prtmo model for a given profile is the variation of the concentration of the given species on the given day of year (from midnight to midnight the next day). At each altitude, the diurnal-scale factor value, s_{diurnal} , is calculated as

$$s_{\text{diurnal}} = \frac{X_{\text{mod}}(\text{LT}_{\text{INST}})}{X_{\text{mod}}(\text{LT}_{\text{ACE}})}, \quad (7)$$

where X is the species concentration, LT is the local time, and the mod, ACE, and INST subscripts refer to the model, ACE-FTS, and the compared instrument values, respectively. The ACE-FTS concentration values can then be scaled to the compared instrument local time using

$$X_{\text{ACE}}(\text{LT}_{\text{INST}}) = X_{\text{ACE}}(\text{LT}_{\text{ACE}}) s_{\text{diurnal}}. \quad (8)$$

As discussed by Brohede et al. (2007b) and Kerzenmacher et al. (2008), for NO₂, the uncertainties due to the diurnal-scale factor profiles are typically less than 20 % in the lower and upper stratosphere and typically less than 10 % in the middle stratosphere. Uncertainties are expected to be of the same order or less for the other NO_y species. For a small fraction of ACE-FTS occultations, the photochemical model failed to produce results. Therefore, in the following section, comparisons between scaled and non-scaled results between ACE-FTS and each INST may not always contain exactly the same number of coincident pairs.

4 Results

4.1 Direct comparisons of ACE-FTS versions 2.2 and 3.5

Direct comparisons between v3.5 and v2.2 of the ACE-FTS NO_y species are shown in Fig. 2. From left to right in each panel, Fig. 2 shows the v3.5 and the v2.2 mean profiles, the correlation coefficient profiles, the mean of the relative differences (v3.5–v2.2 divided by the mean v2.2 profile), and the standard deviation of the relative differences. Figure 1a–e show results for NO, NO₂, HNO₃, N₂O₅, and ClONO₂.

For NO, it can be seen that up to 60 km the two versions are highly correlated, with a correlation coefficient of nearly 1 at most altitudes, dropping to 0.92 at the lowest altitude level. Between altitudes of 25 and 43 km, the relative differences are better than 2% with standard deviations less than 10%. At higher altitudes, up to 60 km, v3.5 NO concentrations are ~5% lower with standard deviations on the order of 30%. This difference can be considered an improvement, as Kerzenmacher et al. (2008) showed that near 60 km ACE-FTS v2.2 NO had a positive bias on the order of 10–15%. Below 22 km, the differences are much worse; however, this is in a region where the NO retrievals are often negative, and below 17 km the mean NO profile of both versions is negative and NO concentrations are over an order of magnitude smaller than above 22 km.

In the altitude region of 17–37 km, v2.2 and v3.5 NO₂ retrievals are very similar. The correlation coefficients are all near 1, relative differences are within 2% and standard deviations are better than 5%. From 37 to 47 km, v3.5 NO₂ reaches a maximum difference of –8% with a standard deviation of 15%. Above 37 km, where there is only a weak NO₂ signal, the standard deviations of the relative means and the correlation coefficients get worse, reaching 137% and 0.7, respectively. Below 17 km, where NO₂ VMR values are significantly lower, v3.5 exhibits lower VMRs than v2.2, with differences reaching –12%.

For HNO₃, correlation coefficients are greater than 0.95 at altitudes of 10 km and higher. Between 10 and 23 km, v3.5 HNO₃ tends to exhibit differences between –1 and 5% with standard deviations on the order of 4–14%. Between 23 and 37 km, v3.5 HNO₃ exhibits 4–8% higher VMRs with standard deviations of 4–13%. Below 10 km, where v3.5 HNO₃ VMR values are lower, the comparison results get much worse with decreasing altitude and at 6 km the correlation coefficient is 0.42, the mean of the relative differences is –53%, and the standard deviation of the relative differences is 130%.

The v3.5 N₂O₅ data exhibit a positive difference that is within 5% between 22 and 37 km and within 15% at all altitudes above 17 km. Above 20 km, correlation coefficients are better than 0.95 and the standard deviations of the relative means are between 15 and 44%. Below 20 km, the compar-

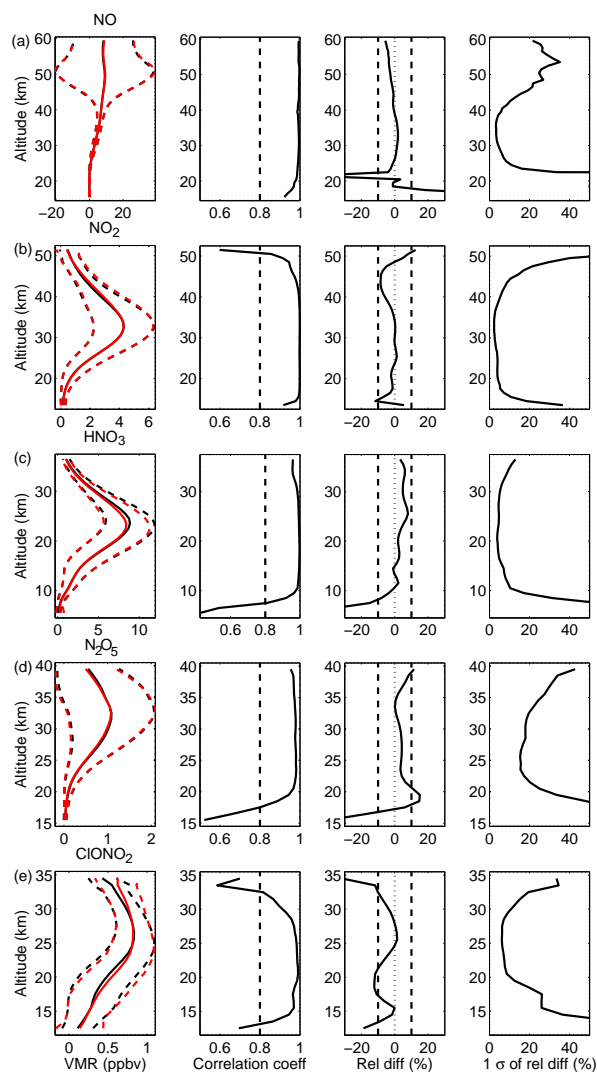


Figure 2. Direct comparisons of ACE-FTS v2.2 with ACE-FTS v3.5 (a) NO, (b) NO₂, (c) HNO₃, (d) N₂O₅, and (e) ClONO₂. From left to right the panels show the mean concentration profiles (red solid for v2.2, black solid for v3.5) with corresponding 1 σ (red dashed for v2.2, black dashed for v3.5) in parts per billion volume (ppbv), correlation coefficient profiles, the mean of the percent differences (v3.5–v2.2 divided by the mean v2.2 profile), and standard deviation of the percent differences. Dashed lines in the correlation (at 0.8) and relative difference plots (at –10, 0, and 10%) are provided for visual clarity.

ison results get worse with decreasing altitude, as the N₂O₅ concentration decreases.

ClONO₂ correlation coefficients are all greater than 0.95 in the altitude region of 15–29 km and greater than 0.8 between 13 and 32 km. Both the v3.5 and v2.2 ClONO₂ mean profiles peak between 26 and 27 km, but the v3.5 peak exhibits a positive difference of 1 \pm 6% and is vertically narrower, with v3.5 exhibiting lower VMRs with differences of 12 \pm 23% at 18 km and 11 \pm 27% at 33 km. The lower v3.5

Table 3. Optimized coincidence criteria and maximum number of coincident profiles.

Instrument data set	Maximum number of coincident profiles				
	NO (3 h, 500 km ^a) (3 h, 100 km ^b)	NO ₂ (4 h, 350 km)	HNO ₃ (6 h, 100 km)	N ₂ O ₅ (3 h, 100 km)	ClONO ₂ (4 h, 100 km)
GOMOS	–	124	–	–	–
HALOE	47	25	–	–	–
MAESTRO	–	17017	–	–	–
MIPAS ESA	–	3965	1022	375	488
MIPAS IMK-IAA	342	3142	854	332	172
MLS	–	–	3445	–	–
OSIRIS	–	1589	–	–	–
POAM III	–	205	–	–	–
SAGE III	–	517	–	–	–
SCIAMACHY	–	6983	–	–	–
SMILES	–	–	29	–	–
SMR	–	–	329	–	–

^a Coincidence criteria for comparisons with HALOE. ^b Coincidence criteria for comparisons with MIPAS IMK-IAA.

VMRs above the peak would improve on the v2.2 high bias of $\sim 20\%$ reported by Wolff et al. (2008); however the lower v3.5 VMRs below the peak would worsen the reported $\pm 1\%$ bias.

4.2 Satellite instrument comparisons

Throughout the discussion of the results, when it is remarked that there are “better” comparison results, what is meant (unless explicitly stated otherwise) is that the correlation coefficients are higher while the standard deviations of the relative differences are lower. Conversely, by “worse” comparison results, it is meant that the correlation coefficients are lower and that standard deviations of the relative differences are higher. When discussing the coincidence criteria for each species, the “optimal” criteria are those that allow for a significant number of coincident profiles (minimum number of 10), but loosening the criteria would generally worsen the comparison results and tightening the criteria would not significantly affect the comparison results. When the “bias” and the “standard deviation” between two data sets are mentioned, unless stated otherwise, these refer to the mean of the relative differences and the standard deviation of the relative differences, respectively. In the following figures, plots of relative differences include error bars that represent the standard error of the mean of the relative differences (shown every 5 km, error bars that are less than $\sim 1\%$ may not be visible); thick solid black lines represent the weighted-average profiles for comparisons that have been diurnally scaled, and the thick dashed black lines represent the weighted-average profiles for comparisons that have not been scaled. Table 3 gives the maximum number of coincident profiles between ACE-FTS and the respective instruments using the optimized coincidence criteria. It should be noted that the number of co-

incidences are typically not constant in altitude due to screening of the data sets using metrics (e.g. retrieval response, quality flags) that are not always constant in altitude.

4.2.1 Comparisons of NO

Note that only HALOE and MIPAS IMK-IAA are being compared with ACE-FTS (MIPAS ESA does not have an NO data product). Figure 3 shows the mean NO VMR profiles for coincident ACE-FTS and HALOE profiles and coincident ACE-FTS and MIPAS IMK-IAA profiles. Since ACE-FTS and HALOE are both solar occultation instruments and only overlapped between 2004 and 2005, there are not many coincident measurements. As such, the spatial coincidence criterion was kept somewhat lax, within 500 km, in order to ensure a statistically significant number of coincidences. It was found that a temporal coincidence criterion of within 3 h also led to a statistically significant number of coincidences (47 profiles). Comparisons with a less stringent criterion (greater than 3 h) led to a larger number of coincidences but significantly reduced the correlation and increased the standard deviation of the relative differences between the data sets. Using a tighter spatial criterion, e.g. within 350 km, also yields a significant number of coincidences but does not significantly improve the comparison results.

At all altitudes, with any temporal coincidence criterion, it was found that using the diurnal scaling factors did not greatly improve the HALOE comparison results. This is likely due to the fact that both ACE-FTS and HALOE are solar occultation instruments, and hence measurements at a common geographic location do not differ greatly in local time. Figure 4a shows the ACE-FTS and HALOE NO comparison results, with and without diurnal scaling. The two data sets are only strongly correlated in the altitude region of

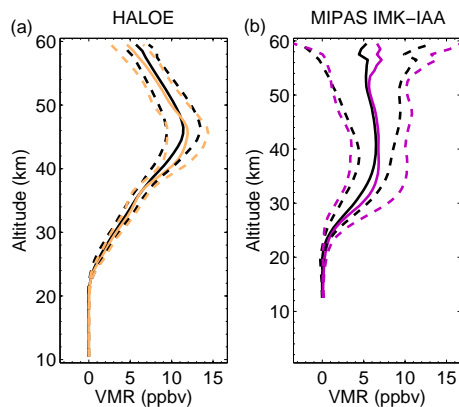


Figure 3. Mean NO profiles (in ppbv) for coincident diurnally scaled ACE-FTS (black) and HALOE (left panel) and MIPAS IMK-IAA (right panel) measurements and corresponding measurement standard deviations (dashed lines). Coincidence criteria for HALOE comparisons are within 3 h and 500 km and are within 3 h and 100 km for MIPAS IMK-IAA comparisons.

approximately 25–55 km. In this region, the relative difference shows that ACE-FTS NO tends to exhibit a low bias of less than 10 %, with standard deviations on the order of 10 % with respect to HALOE. Of the 47 coincident profiles 41 are local sunset occultations, and the remaining 6 are local sunrise occultations. Due to the lack of sunrise measurements, it was not possible to determine whether or not there is a significant bias between the sunrise and sunset (or similarly local morning and local evening) NO profiles.

It was found that the temporal and spatial coincidence criteria that optimized the comparison results for diurnally scaled ACE-FTS and MIPAS IMK-IAA NO profiles were within 3 h and within 100 km. Similar to the HALOE comparisons, using the diurnal scaling factors did not greatly improve the comparison results at most altitudes. However, for temporal differences larger than 3 h, using the diurnal scaling factors worsened comparison results at all altitudes.

Figure 4b shows the ACE-FTS and MIPAS IMK-IAA NO comparison results with and without diurnal scaling, using coincidence criteria of within 3 h and 100 km. Throughout the middle stratosphere, the diurnal scaling generally increased the correlation coefficients by ~ 0.05 and lowered the standard deviations by ~ 3 %. Relatively strong correlation is seen above 25 km, where ACE-FTS exhibits a negative bias within -10 and -22 % between 25 and 35 km and an approximate -5 % bias between 40 and 50 km. The lowest standard deviations are observed in the 30–50 km region, on the order of 35–50 %. The higher standard deviations (relative to comparisons with HALOE, Fig. 4a) reflect the higher variance within the MIPAS IMK-IAA NO data set. Below 25 km, the relative differences get more negative with decreasing altitude – more negative than -100 % below 21 km. An ACE-FTS NO low bias with respect to non-solar occultation instruments, on the order of ~ 10 –40 %, is expected in

this region due to not accounting for diurnal variations of NO along the line of sight (Brohede et al., 2007a).

Figure 5a shows NO comparison results for data separated by local time using all available MIPAS IMK-IAA data. It can be seen that there is an apparent significant local time bias in the ACE-FTS–MIPAS IMK-IAA comparison results. Between 19 and 52 km, the correlation coefficients are better for local evening (PM) comparisons than for local morning (AM) comparisons by up to 0.4, and at all altitudes the evening comparisons exhibit lower standard deviations by ~ 15 –50 %. This leads to improved relative differences when only using the evening data in the 25–34 and 52–60 km ranges. However, due to the orbital geometries and the MIPAS IMK-IAA retrieval sensitivity to NO, the only coincident PM data are during November–January in the Southern Hemisphere (SH) and May–July in the Northern Hemisphere (NH), hereafter referred to as “summer” months. Figure 5b shows NO comparison results between ACE-FTS and MIPAS IMK-IAA for data separated by local time and using only the summer months (both NH and SH). It can be seen that correcting for this seasonal bias greatly improves the AM comparison results, as there is less NO variation in the polar summer regions than in the winter. At most altitudes the summer PM comparisons still tend to exhibit better correlation than the AM, but the summer AM and PM standard deviation profiles are rather similar – values of ~ 100 % near 18 km, then decreasing with altitude to ~ 15 –20 % near 45 km, and from there increasing with altitude. Between 22 and 52 km, the summer AM and PM relative difference profiles are also quite similar. ACE-FTS exhibits a negative bias with respect to MIPAS IMK-IAA of approximately -100 to -10 % between 22 and 27 km. Above 27 km, up to ~ 50 km, ACE-FTS NO is typically systematically lower than MIPAS IMK-IAA by 0–10 %. Above 52 km, the summer PM results (correlation coefficients and standard deviations) are typically better than the AM; the PM relative differences are between 0 and $+7$ %, and the AM relative differences decrease with altitude from 0 to -32 % between 53 and 60 km.

4.2.2 Comparisons of NO₂

From Fig. 6 it is apparent that in the comparisons with all other NO₂ data sets, the diurnally scaled ACE-FTS profiles have a low bias near the NO₂ peak, ~ 33 km. It can be seen in Fig. 7a that using coincidence criteria of within 350 km and within 4 h without any diurnal scaling leads to relatively poor agreement between ACE-FTS and many instruments in the middle to upper stratosphere. Near the NO₂ peak, without diurnal scaling mean relative differences between ACE-FTS and INST data range from -38 to $+2$ %, with standard deviations that reach up to ~ 50 %. With diurnal scaling (Fig. 7b), near the NO₂ peak the ACE-FTS low bias is on the order of -20 to -5 %, with standard deviations on the order of 7–35 %. Between 30 and 40 km, the weighted-average mean relative differences are on the order of -10 %, and the standard deviations are on the order of 10–20 %.

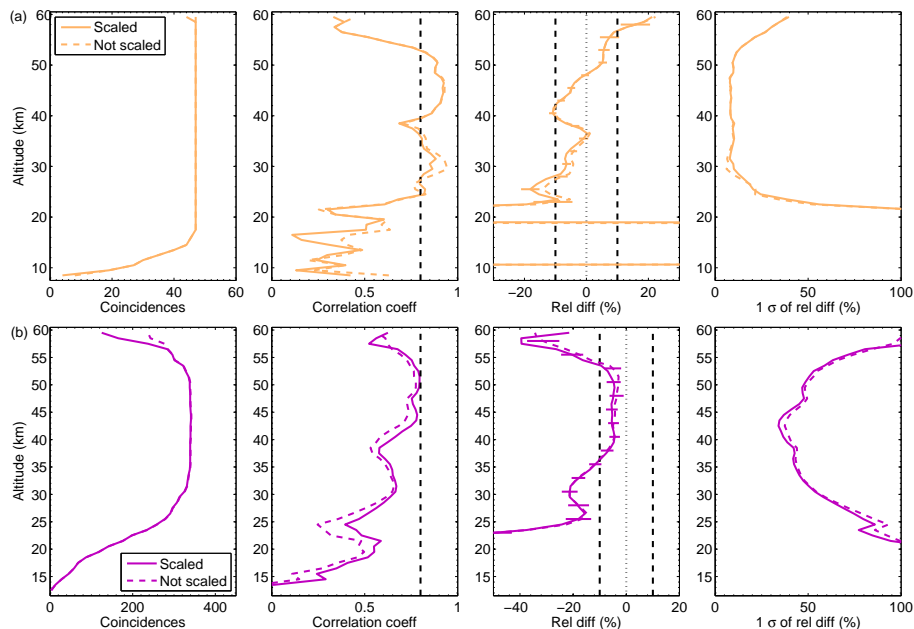


Figure 4. Comparisons of diurnally scaled (solid lines) and non-scaled (dashed lines) ACE-FTS NO profiles with (a) HALOE data within 3 h and 500 km, and (b) MIPAS IMK-IAA data within 3 h and 100 km. From left to right, the plots show number of coincident profiles, correlation coefficient profiles, mean relative difference profiles (ACE-FTS–INST) in percent, and standard deviation of relative difference profiles in percent. Error bars in the relative difference profiles represent the standard error of the mean.

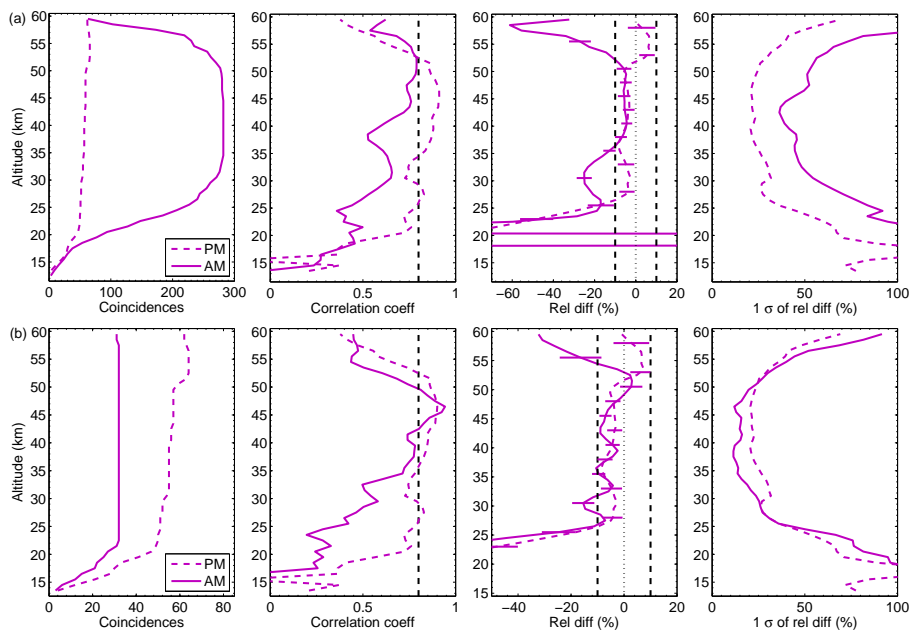


Figure 5. Comparisons of ACE-FTS NO profiles with MIPAS IMK-IAA data within 3 h and 100 km for local morning (solid lines) and evening (dashed lines) data. From left to right, the plots in each panel show number of coincident profiles, correlation coefficient profiles, mean relative difference profiles (ACE-FTS–MIPAS IMK-IAA) in percent, and standard deviation of relative difference profiles in percent. (a) All data; (b) only summer months (May–Jul in NH, Nov–Jan in SH) are included. Error bars in the relative difference profiles represent the standard error of the mean.

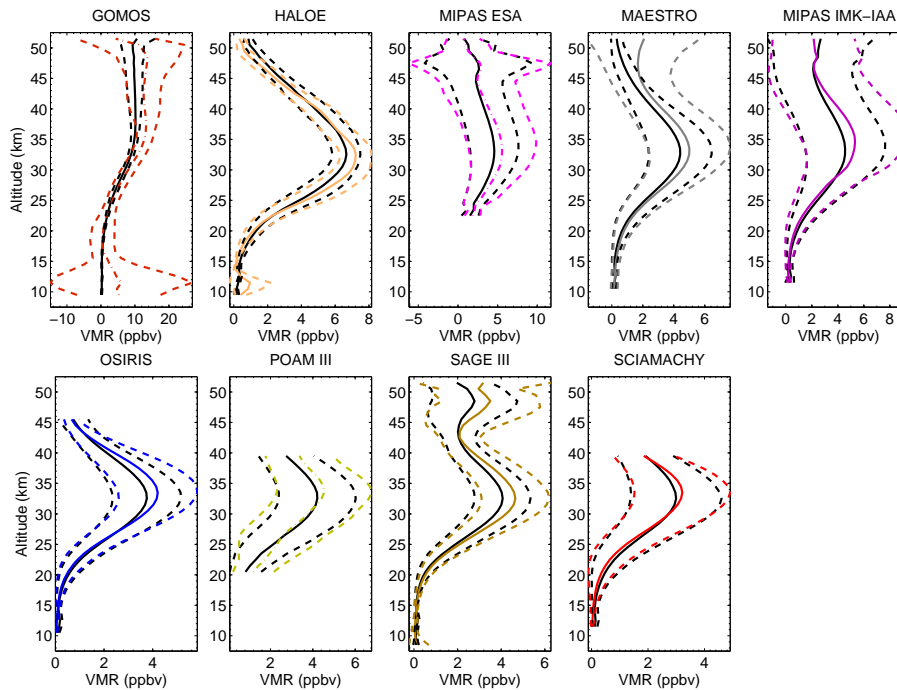


Figure 6. Mean NO₂ profiles (in ppbv) for coincident diurnally scaled ACE-FTS (black) and INST (coloured) measurements and corresponding measurement standard deviations (dashed lines). Coincidence criteria for all comparisons are within 4 h and 350 km.

with weighted-average standard deviations within 18–43 %. Within these altitudes, most comparisons typically yielded correlation coefficients that were greater than 0.8, the exception being GOMOS which measures at nighttime. The weighted-average correlation coefficients are better than 0.8 between 15 and 40 km and better than 0.9 between 17 and 35 km.

Below 25 km, an ACE-FTS NO₂ positive bias is expected with respect to instruments that do not use the solar occultation viewing geometry due to not accounting for diurnal variations in NO₂ along the line of sight in the forward model. In solar occultation viewing geometry, not accounting for this diurnal effect is expected to lead to a ~ 10 –40 % positive bias (Brohede et al., 2007a). It can be seen from Fig. 7b that, below 25 km, ACE-FTS does have a positive bias on the order of 5–40 % with respect to MIPAS IMK-IAA, OSIRIS, and SCIAMACHY. As well, below 22 km, ACE-FTS exhibits a positive bias with respect to HALOE, which is a solar occultation instrument but accounts for the diurnal effect in the NO₂ retrieval algorithm.

Diurnal scaling has less of an effect on comparisons with the solar occultation instruments (HALOE, POAM III, SAGE III) than on those with other viewing geometries, as there is less of a difference in measurement local times, and diurnal scaling has no effect on the ACE-FTS comparisons with MAESTRO as measurements are co-located (although they do have differing vertical and horizontal resolutions). In order to determine biases in the comparisons due to lo-

cal time or hemispheric coverage, comparisons were made in the 20–40 km region where the NO₂ peak is well sampled and the majority of instruments have sufficient coverage. For local time differences, GOMOS data have been excluded, as it only contains local evening data, and the solar occultation instruments have been excluded as they tend to only have a significant number of coincidences in either local morning or local evening. For hemispheric differences only HALOE data were excluded, as the vast majority of HALOE data are from the NH.

As can be seen in Fig. 8, at all altitudes within the 20–40 km range, the weighted-average results are generally better for the evening comparisons than for the morning comparisons. The weighted-average standard deviations are better by up to 18 % and the correlation coefficients are better by ~ 0.05 in the evening comparisons. The evening average relative differences are ~ 0 % near 20 km, reach -11 % near 35 km, and 6 % near 40 km. Whereas for morning results, compared to evening results, average relative differences are more negative above 35 km (reaches -13 %) and more positive below 30 km (up to $+40$ %). The better evening results are likely due to differences in the diurnal variation along the line of sight between sunrise and sunset observations. For sunrise (local morning) observations, ACE-FTS samples a region of the atmosphere that has yet to be sunlit long enough for NO₂ to be in equilibrium. For sunset (local evening), however, the entire sampled area should be relatively stable. As can be seen in Fig. 9, there were no major differ-

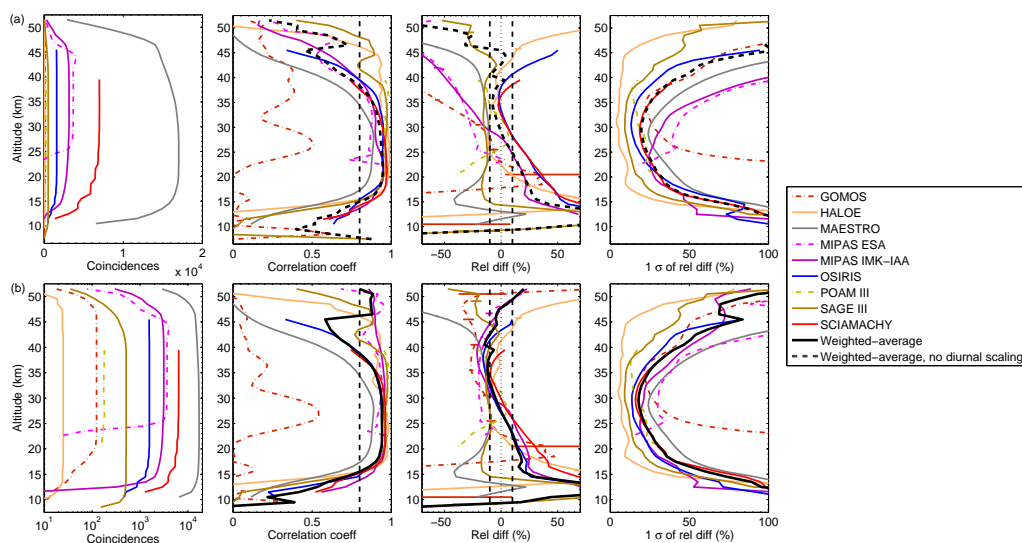


Figure 7. Comparisons of ACE-FTS NO₂ profiles with correlative data sets using coincidence criteria of within 4 h and 350 km. From left to right, the plots in each panel show number of coincident measurements, correlation profiles between ACE-FTS and INST, mean relative difference profiles (ACE-FTS–INST) in percent, and standard deviation of relative difference profiles in percent. **(a)** Comparisons without diurnal scaling; **(b)** comparisons with diurnal scaling. Error bars in the relative difference profiles represent the standard error of the mean (values less than ~ 1 % may not be visible).

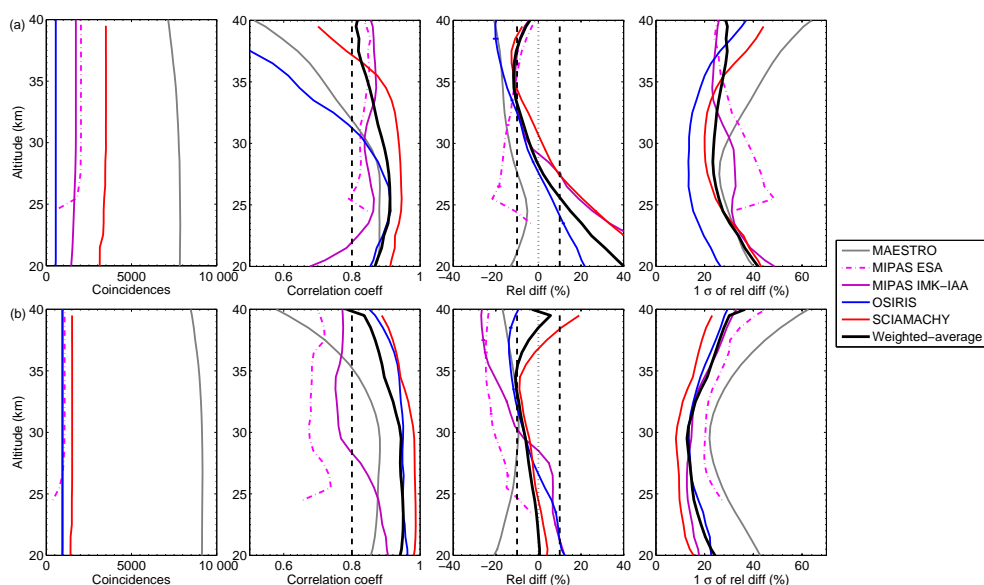


Figure 8. Comparisons of ACE-FTS NO₂ profiles with correlative data sets using coincidence criteria of within 4 h and 350 km: **(a)** comparisons for local morning and **(b)** local evening. Note that GOMOS and the solar occultation instruments have been excluded. Error bars in the relative difference profiles represent the standard error of the mean (values less than ~ 1 % may not be visible).

ences in the weighted-average results between the NH comparisons and the SH comparisons. The only significant difference in the weighted-average relative differences is below 25 km, with the SH exhibiting larger values by up to 7 %.

4.2.3 Comparisons of HNO₃

Due to the relatively weak diurnal variation of HNO₃ in the stratosphere, using the photochemical box model did not improve the HNO₃ comparison results at any altitude level. In addition, a lax temporal coincidence criterion of within 6 h was used, as tightening the criterion did not significantly im-

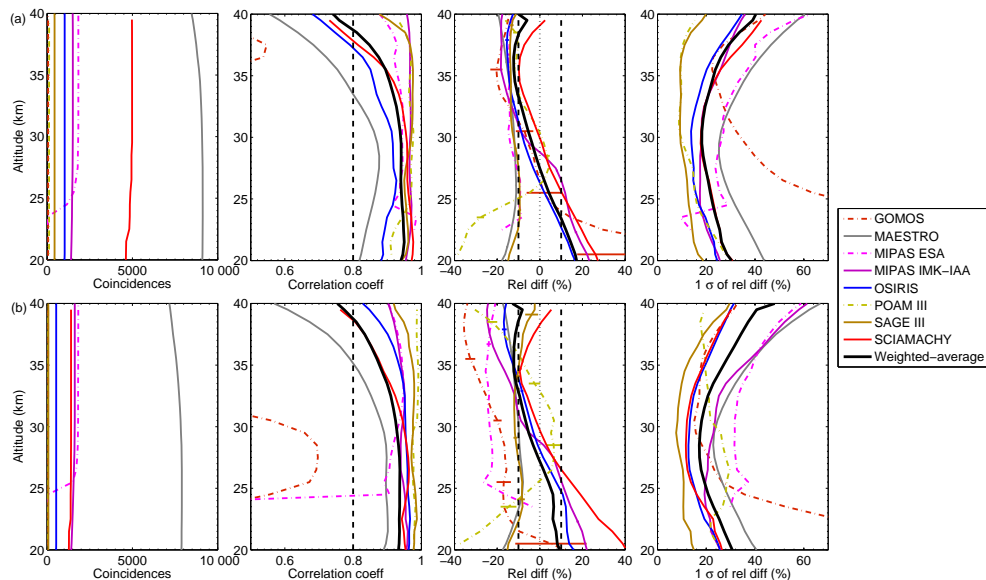


Figure 9. Comparisons of ACE-FTS NO₂ profiles with correlative data sets using coincidence criteria of within 4 h and 350 km: **(a)** northern hemispheric data and **(b)** southern hemispheric data. Note that HALOE has been excluded. Error bars in the relative difference profiles represent the standard error of the mean (values less than $\sim 1\%$ may not be visible).

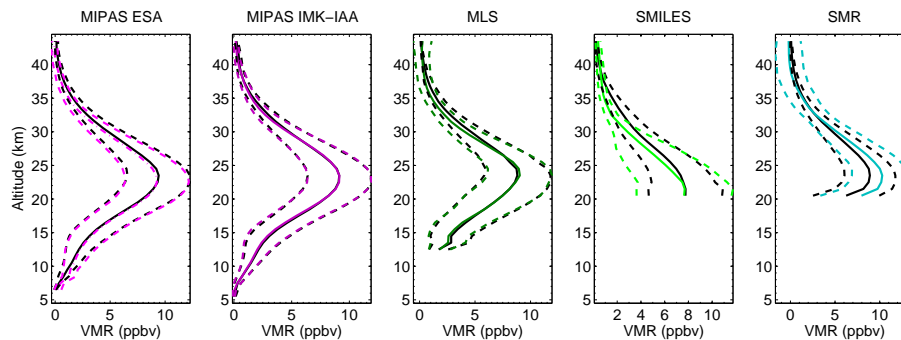


Figure 10. Mean HNO₃ profiles for coincident diurnally scaled ACE-FTS (black) and INST (coloured) measurements and corresponding measurement standard deviations (dashed lines). Coincidence criteria for all comparisons are within 6 h and 100 km.

prove comparison results. As such, it was possible to use a spatial coincidence criterion of within 100 km, which optimized the comparison results.

Figure 10 shows the mean coincident ACE-FTS and INST HNO₃ profiles along with the 1σ measurement variation. There is typically good agreement between ACE-FTS and the other instruments, and HNO₃ comparison results are shown in Fig. 11. Near the HNO₃ peak, ~ 20 – 25 km, there is excellent agreement, with weighted-average relative differences within -1% , correlation coefficients of ~ 0.97 , and standard deviations of $\sim 8\%$.

The weighted-average correlation coefficients are greater than 0.5 for altitudes of 7–40 km and greater than 0.9 for altitudes of 12–31 km. Between 9 and 38 km the weighted-average standard deviations are below 50%, reaching a minimum of 7% near 24 km. The weighted-average relative

differences are within $\pm 6\%$ between 9 and 29 km. Above 30 km, the average relative differences increase with altitude to 37% at 40 km; however, at that altitude only the MIPAS IMK-IAA comparisons exhibit standard deviations below 50%, and the ACE-FTS–MIPAS IMK-IAA relative difference at 40 km is on the order of 20%. Below 30 km, the ACE-FTS–MIPAS ESA relative differences are within $\pm 10\%$, and the ACE-FTS–MLS differences are typically on the order of -5 to 10%. Between 20 and 38 km ACE-FTS typically exhibits a high bias with respect to SMILES, which is on the order of 1% near 20 km and increases to 55% near 34 km. With respect to SMR, ACE-FTS exhibits a negative bias on the order of -9 to -13% between 25 and 30 km. This is an improvement from the $\sim 20\%$ low bias exhibited in the ACE-FTS v2.2 and SMR v2.0 comparisons reported by

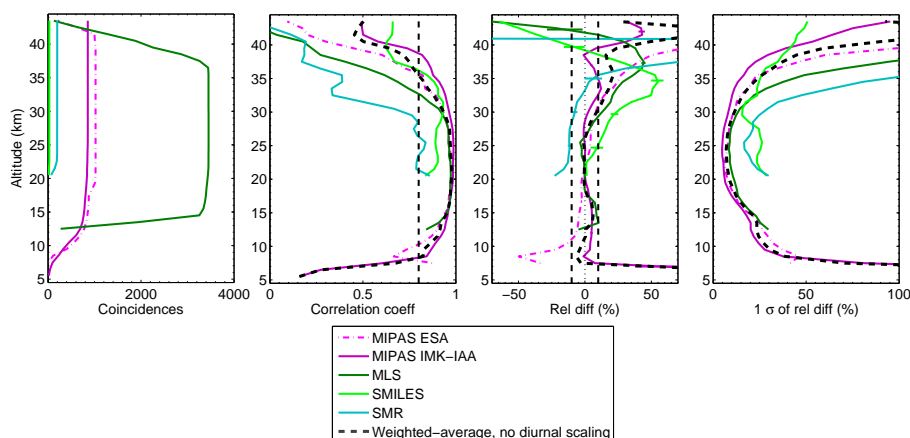


Figure 11. Same as top panel of Fig. 7 (comparisons with no diurnal scaling) except for HNO₃, with coincidence criteria of within 6 h and 100 km. Error bars in the relative difference profiles represent the standard error of the mean (values less than $\sim 1\%$ may not be visible).

Wolff et al. (2008). Mean relative differences between ACE-FTS and SMR are also within $\pm 10\%$ between 30 and 35 km.

There were no major local time biases found in the HNO₃ comparisons. In the altitude range 12–28 km, weighted-average mean relative differences were within $\pm 4\%$ for the local morning comparisons, whereas local evening comparisons yielded weighted-average mean relative differences within $\pm 7\%$ (not shown). Between 16 and 38 km, there was no significant hemispheric bias found in the HNO₃ comparisons, with SMILES data excluded (due to asymmetric hemispheric coverage).

4.2.4 Comparisons of N₂O₅

Before showing the N₂O₅ validation results, it should be noted that a significant difference was found between local morning and local evening MIPAS (both ESA and IMK-IAA) N₂O₅ comparisons with ACE-FTS: the evening comparisons exhibited much worse agreement than the morning comparisons. Figure 12a shows results for comparisons between local evening diurnally scaled ACE-FTS and MIPAS profiles using coincidence criteria of within 3 h and within 100 km. Near 20–25 km, the relative differences are on the order of $\pm 10\%$ with standard deviations of ~ 50 – 80% and correlation coefficients of ~ 0.65 – 0.75 . However, outside of this region, comparison results yield poorer results, with weak correlation, standard deviations greater than 100% , and relative differences beyond $\pm 100\%$. In order to highlight that this poor agreement is not an issue with differences due to diurnal variation, Fig. 12b shows comparisons using non-scaled ACE-FTS profiles and with a much tighter temporal coincidence of within 20 min (and within 200 km). In comparing to both MIPAS data products in this case, there are large systematic differences from ACE-FTS. The MIPAS ESA differences range from approximately -60 to 200% and the MIPAS IMK-IAA differences range from approxi-

mately -130 to 200% . The poor agreement in the evening is mostly due to the low signal-to-noise ratio in the ACE-FTS measurements due to the lower N₂O₅ concentrations at sunset than at sunrise.

Figures 13 and 14 show the results of the morning comparisons. At coincidence criteria of within 3 h and 100 km, with diurnal scaling, ACE-FTS and MIPAS tend to agree best in the altitude range of 22–34 km. In this region, weighted-average correlation coefficients are better than 0.8, weighted-average standard deviations are between 16 and 40% , and weighted-average mean relative differences are typically better than -7% . Above 34 km, ACE-FTS exhibits a positive bias that is within 10% up to 38 km and increases with altitude, up to 33% at 43 km. This positive bias in the upper altitudes is not reduced when tighter temporal coincidence criteria are chosen (down to within 20 min) and exists both with and without diurnal scaling. Also shown in Fig. 14 are the weighted-average comparison results for non-scaled ACE-FTS profiles. It can be seen that using the photochemical box model does improve the comparison results, especially in the 23–38 km region, where it leads to an improvement to the average standard deviations on the order of 5% . Diurnal scaling also reduces the positive bias above 33 km by up to 16% .

Although there was very poor agreement between local evening ACE-FTS and local evening MIPAS N₂O₅ profiles, comparisons between diurnally scaled morning ACE-FTS and evening MIPAS N₂O₅ profiles yield much better agreement. This indicates that the poor agreement seen in the evening data is most likely due to the high level of noise in the evening ACE-FTS N₂O₅ data and is unlikely an issue with the MIPAS data. Figure 15 shows the weighted-average results (scaled) from Fig. 14, along with comparison results between diurnally scaled morning ACE-FTS and evening MIPAS (both ESA and IMK-IAA) profiles using coincidence criteria of within 12 h and within 100 km. Between

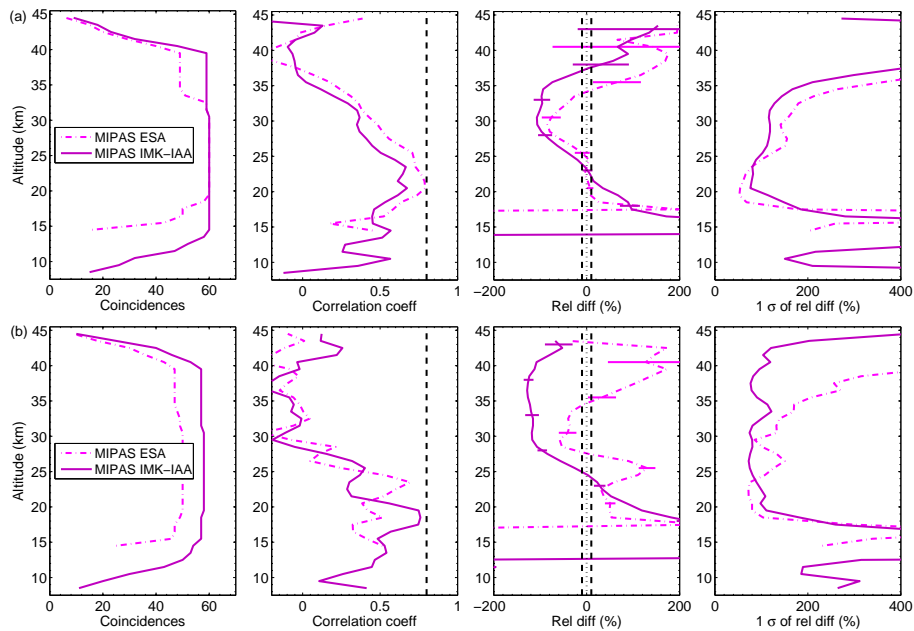


Figure 12. Evening N₂O₅ comparisons between ACE-FTS and MIPAS measurements: **(a)** with diurnal scaling and coincidence criteria of within 3 h and 100 km and **(b)** without diurnal scaling and with coincidence criteria of within 20 min and 200 km. Error bars in the relative difference profiles represent the standard error of the mean.

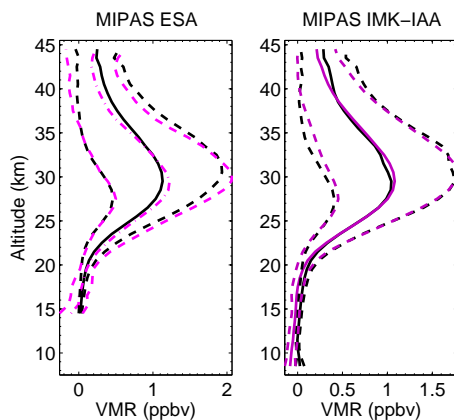


Figure 13. Mean morning N₂O₅ profiles (in ppbv) for coincident diurnally scaled ACE-FTS (black) and MIPAS (coloured) measurements and corresponding measurement standard deviations (dashed lines). Coincidence criteria for all comparisons are within 3 h and 100 km.

22 and 37 km, the morning/evening weighted-average correlation coefficients are greater than 0.8 and the standard deviations are less than 50 %. In this altitude range, the weighted-average relative differences are better than 10 %.

4.2.5 Comparisons of ClONO₂

Figures 16 and 17 show the results of the ACE-FTS and MIPAS, both ESA and IMK-IAA, comparisons. Comparisons

were found to be optimized at coincidence criteria of within 4 h and 100 km. With diurnal scaling, ACE-FTS and MIPAS tend to agree best in the altitude range of 17–34 km. In this region, weighted-average correlation coefficients are better than 0.7, weighted-average standard deviations are between 13 and 32 %, and weighted-average mean relative differences tend to exhibit a negative bias within -1 and -10 %, except at the lower altitudes where the low bias reaches -20 % near 17 km. Also shown in Fig. 17 are the weighted-average comparison results for non-scaled ACE-FTS profiles. It can be seen that using the photochemical box model does improve the comparison results, especially above 26 km, where it leads to an improvement to the average correlation coefficients by up to 0.15 and to the average standard deviations by up to 4 %.

Similar to the case for the MIPAS IMK-IAA NO comparisons in Sect. 4.2.1, separating the coincident MIPAS ClONO₂ data into morning and evening subsets seasonally biases the data. Due to the orbital geometries and the MIPAS retrievals' sensitivity to ClONO₂, there is typically only coincident evening data between February and April in the NH and August and October in the SH (henceforth referred to as “spring” months). In examining the differences between spring morning and evening comparison results, shown in Fig. 18, between 17 and 36 km there are no major differences in the weighted-average relative difference profiles. In the 13–23 km region, where the comparison results are more consistent for the evening results, both the morning and evening results tend to exhibit a -10 % bias. Above 25 km,

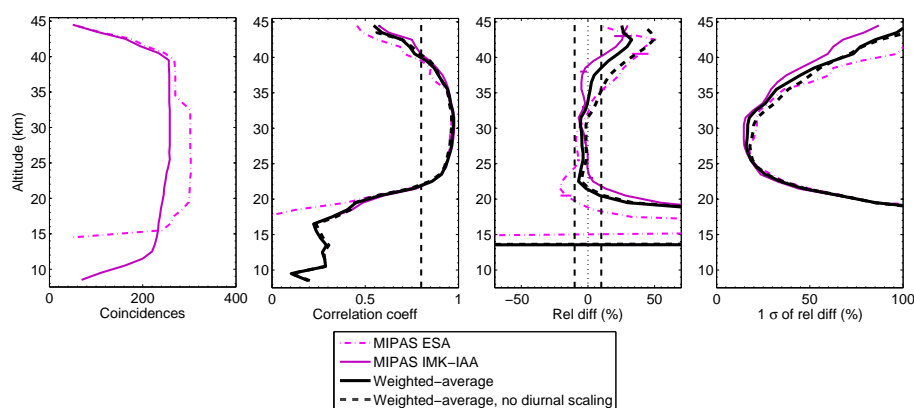


Figure 14. Morning N₂O₅ comparisons between ACE-FTS and MIPAS measurements with coincidence criteria of within 3 h and 100 km. Error bars in the relative difference profiles represent the standard error of the mean.

Table 4. Summary of validated ACE-FTS NO_y systematic differences for two different cases. Case 1: region where the weighted-average correlation coefficient profile is greater than 0.5 and the weighted-average standard deviation of the relative differences profile is less than 100 %. Case 2: region where the weighted-average correlation coefficient profile is greater than 0.8 and the weighted-average standard deviation of the relative differences profile is less than 50 %. Results are for comparisons using all data and the species-dependent optimized coincidence criteria (given in text and Table 3).

Species	$r > 0.5, \sigma < 100\%$		$r > 0.8, \sigma < 50\%$	
	Altitude (km)	Bias (%)	Altitude (km)	Bias (%)
NO HALOE	23–27	–20 to –12	27–53	–15 to 6
	28–48	Better than –10		
	48–57	0 to 17		
NO MIPAS IMK-IAA (Summer only)	21–26	–100 to –10	36–52	–9 to 2
	27–51	Better than –10		
	53–56	–11 to 4		
NO ₂	15–27	25 to 0	17–27	Better than 18
	28–47	Better than –15	28–41	Better than –15
	48–52	Better than 20		
HNO ₃	8–29	Within ±7	9–17	Within ±7
	30–40	7 to 37	18–26	Within ±1
			27–35	1 to 20
N ₂ O ₅ (Morning only)	21–34	Better than –7	22–34	Better than –7
	35–42	0 to 33	35–38	0 to 7
ClONO ₂	14–24	Better than –20	16–24	Better than –20
	21–35	Better than –8	21–33	Better than –8
	36–38	0 to 37		

the comparison results are typically better for the morning results. Between 25 and 33 km the morning relative differences are typically between –5 and 0 % and evening relative differences are typically between –1 and 4 %. Near 36 km, both the morning and evening values are ~ 10 % and then increase with altitude, up to 74 % at 40 km for morning data and 130 % for evening data.

5 Discussion and summary

The ACE-FTS v3.5 NO, NO₂, HNO₃, N₂O₅, and ClONO₂ data sets have been compared to correlative data sets from multiple satellite limb sounders. The comparison used a photochemical box model to scale the ACE-FTS data to the local times of the individual correlative measurements. Results for NO₂, N₂O₅, and ClONO₂ were improved by diurnal scaling. By improved, it is meant that there is both an increase

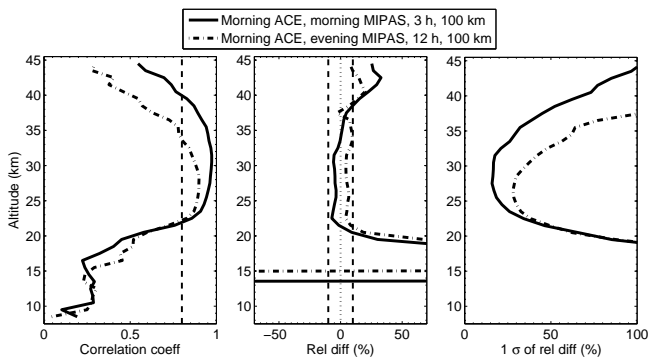


Figure 15. Weighted-average results for morning N_2O_5 comparisons between ACE-FTS and MIPAS (ESA and IMK-IAA) measurements with coincidence criteria of within 3 h and 100 km (solid) and comparisons between diurnally scaled morning ACE-FTS and evening MIPAS measurements with coincidence criteria of within 12 h and 100 km (dot-dash).

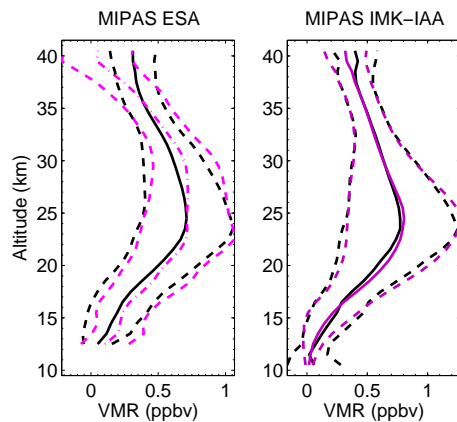


Figure 16. Mean ClONO_2 profiles (in ppbv) for coincident diurnally scaled ACE-FTS (black) and MIPAS (coloured) measurements and corresponding measurement standard deviations (dashed lines). Coincidence criteria for all comparisons are within 4 h and 100 km.

in correlation coefficient values and a decrease in standard deviations of the relative differences.

Table 4 summarizes the average systematic differences between ACE-FTS and the data sets for all other instruments in the regions where the ACE-FTS data have been validated and where there is typically a strong correlation and reasonable standard deviations. The column outlining the systematic differences where average correlation coefficients are better than 0.8 and average standard deviations are typically below 50 % could also be used to determine recommended altitude limits for the different ACE-FTS data sets (with the exception of NO , which was only examined below 60 km, the top altitude of the photochemical model).

In general there is good agreement between ACE-FTS and HALOE NO , but as mentioned above, the diurnal scaling

factors did not help improve the comparison results. Comparisons indicated that ACE-FTS has a negative bias on the order of 0 to -10% in the altitude region of 28–48 km. This is a slight improvement on the ACE-FTS v2.2 NO profiles, which Kerzenmacher et al. (2008) found to have a $\sim 8\%$ bias with HALOE in this region.

ACE-FTS and MIPAS IMK-IAA comparisons suggest that ACE-FTS NO has a negative bias at all altitudes below 60 km, and between 40 and 60 km this bias is approximately -5% . Below 25 km, the bias becomes more negative with decreasing altitude from -15% to beyond -100% , and 10–40 % of this bias is expected to be due to diurnal variations along the ACE-FTS line of sight. Comparisons using only summer data yield similar results. Both summer morning and summer evening comparisons yield negative relative differences at all altitudes, with values more negative than 50 % below ~ 23 km and above ~ 50 km, and within -10 and 0 % in the 32–50 km region.

ACE-FTS v3.5 NO_2 profiles have a clear systematic negative bias with respect to all other instruments at and around the NO_2 peak, ~ 32 km. With diurnal scaling, this negative bias near the peak is $\sim -10\%$ for evening comparisons (which typically yield better results than morning comparisons) and $\sim -12\%$ for morning comparisons. This bias is likely in part due to errors in the characterization of the ACE-FTS instrumental line shape in v3.5 (Boone et al., 2013), but the complete source of this bias is the subject of on-going investigations. Better evening comparison results than morning results are likely attributable to sunrise observations sampling a region of the atmosphere where NO_2 concentrations are not yet in daytime equilibrium. Below 25 km, ACE-FTS tends to exhibit a 5–40 % positive bias with respect to non-solar occultation instruments and HALOE. This bias is expected due to diurnal variation of NO_2 along the ACE-FTS line of sight that is not accounted for in the forward model. No major differences were found between NH comparisons and SH comparisons, but below 25 km the average relative differences were on the order of 8 % in the SH, and on the order of 15 % in the NH. These results are an improvement over the findings of Kerzenmacher et al. (2008), who found that ACE-FTS v2.2 NO_2 had a $\sim 15\%$ low bias near the peak and between 20 and 40 km agreed with correlative data sets to within 40 %.

HNO_3 comparisons near 35 km show that ACE-FTS has a positive bias that on average is $\sim 20\%$. Within the 8–30 km range ACE-FTS and correlative data sets on average are within $\pm 7\%$, and around the HNO_3 peak (~ 20 –26 km) on average ACE-FTS is within $\pm 1\%$ of the other measurements. These results suggest an improvement from ACE-FTS v2.2 comparisons by Wolff et al. (2008), who found that ACE-FTS was typically within $\pm 20\%$ of correlative satellite data sets. No major biases in the HNO_3 comparisons were found due to measurement local time or hemispheric coverage.

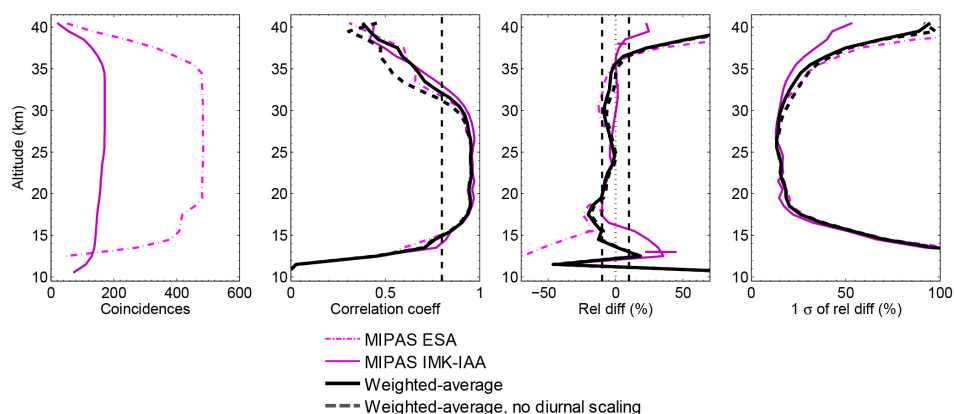


Figure 17. Comparisons between ACE-FTS and MIPAS ClONO₂ measurements with coincidence criteria of within 4 h and 100 km. Error bars in the relative difference profiles represent the standard error of the mean.

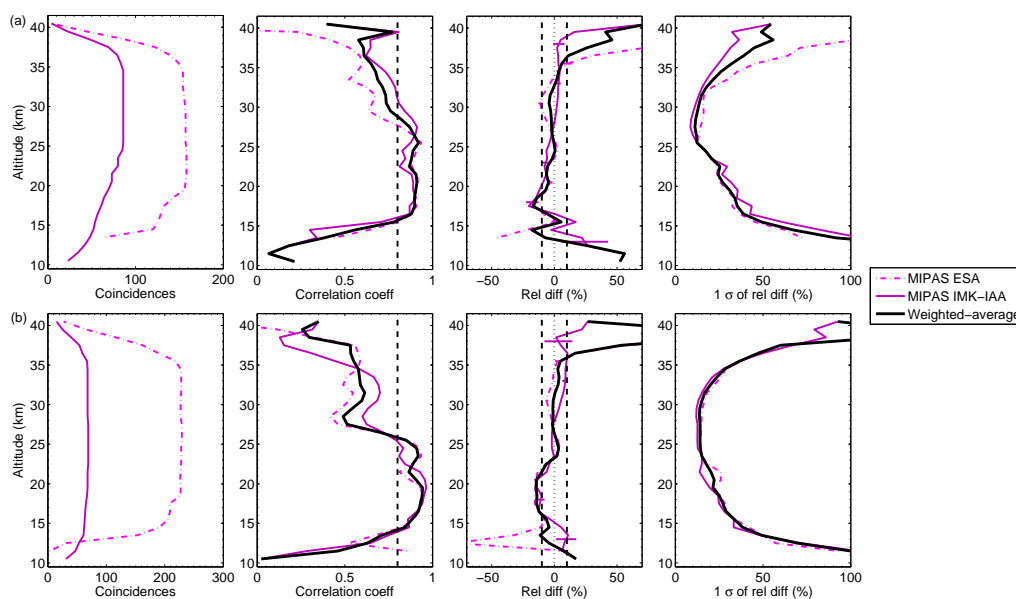


Figure 18. Comparisons of ACE-FTS ClONO₂ profiles with MIPAS data sets (ESA and IMK-IAA) using coincidence criteria of within 4 h and 250 km and only spring months (Feb–Apr in NH, Aug–Oct in SH). (a) Comparisons for local morning and (b) local evening. Error bars in the relative difference profiles represent the standard error of the mean.

Above 35 km, morning ACE-FTS N₂O₅ has a positive bias with respect to MIPAS ESA and IMK-IAA, which reaches 33 % near 42 km. This bias is not an artifact of diurnal mismatch as it still exists when comparing profiles using a temporal coincidence criterion on the order of 20 min (not shown). At these higher altitudes, where the VMR is decreasing with altitude, it is difficult to accurately derive N₂O₅ concentrations given the broad, unstructured N₂O₅ absorption spectrum. Between 22 and 35 km, ACE-FTS tends to exhibit a negative bias, on average better than −7 %.

Evening ACE-FTS N₂O₅ profiles show very poor agreement with evening MIPAS measurements regardless of diurnal scaling, coincidence criteria, and hemisphere. As the coincident ACE-FTS measurements are always evening sun-

set measurements, this is when N₂O₅ is at its least abundant (roughly an order of magnitude less than morning concentrations) and therefore where the ACE-FTS N₂O₅ retrievals suffer from the lowest absorption signals for the molecule. The evening MIPAS retrievals are most likely not equally affected by the low abundance of N₂O₅, as they compare reasonably well with morning ACE-FTS profiles that have been diurnally scaled to match the MIPAS local times. Further investigation into the poorer quality of the ACE-FTS evening N₂O₅ data is needed.

In the 14–35 km region ACE-FTS ClONO₂ exhibits a negative bias with respect to the MIPAS data sets. From 14 to 24 km, the ACE-FTS bias is on average better than −20 %, and in the 21–35 km region better than −8 %. Differences

in morning and evening ACE-FTS–MIPAS comparison results are examined for the spring months. Major differences are only exhibited above 25 km, where the comparison results are typically better for the morning results. In the 25–33 km range, spring morning relative differences on average are -3% and the spring evening relative differences on average are $+2\%$. Below ~ 25 km, these results are slightly worse than those of Wolff et al. (2008), who found that below ~ 25 km ACE-FTS v2.2 ClONO₂ data were typically within 1% of MIPAS IMK-IAA data. Although at higher altitudes, ACE-FTS v2.2 exhibited a positive bias of up to 20% near 33 km, and therefore above the VMR peak v3.5 ClONO₂ has improved.

6 Data availability

The ACE-FTS Level 2 data used in this study can be obtained via the ACE-FTS website (registration required), <http://www.ace.uwaterloo.ca> (ACE-FTS, 2016), or upon request from the corresponding author (kaley.walker@utoronto.ca). The GOMOS data can be obtained via <https://earth.esa.int/web/guest/data-access> (registration required) (ESA, 2016a). The HALOE data can be obtained via <http://haloe.gats-inc.com/download/index.php> (HALOE, 2016). The MIPAS ESA data can be obtained via <https://earth.esa.int/web/guest/data-access> (registration required) (ESA, 2016b). The MIPAS IMK-IAA data can be obtained via <https://www.imk-asf.kit.edu/english/308.php> (registration required) (KIT, 2016). The MLS data are publicly available via <http://disc.sci.gsfc.nasa.gov/Aura/data-holdings/MLS/index.shtml> (registration required) (GES DISC, 2016). The OSIRIS data can be obtained via <http://odin-osiris.usask.ca> (registration required) (University of Saskatchewan, 2016). The POAM III data can be obtained via https://eosweb.larc.nasa.gov/project/poam3/poam3_table (registration required) (NASA, 2016a). The SAGE III data can be obtained via https://eosweb.larc.nasa.gov/project/sage3/sage3_table (registration required) (NASA, 2016b). The SCIAMACHY data can be obtained via <http://www.iup.uni-bremen.de/scia-arc/> (registration required) (IUP, 2016). The SMILES data can be obtained via <https://www.darts.isas.jaxa.jp/iss/smiles/> (DARTS, 2016). The SMR data can be obtained via <http://odin.rss.chalmers.se> (registration required) (Odin/SMR, 2016).

Appendix A

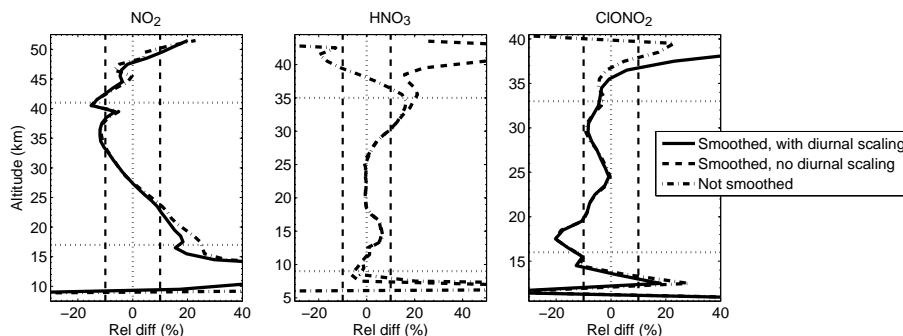


Figure A1. Weighted-average relative difference profiles for vertically smoothed (solid and dashed) and non-smoothed (dot-dashed) NO₂, HNO₃, and ClONO₂ data. The only profiles that do not include diurnal scaling are those for HNO₃. Horizontal dotted lines indicate altitude limits within which the ACE-FTS comparisons yield average correlation coefficients greater than 0.8 and average standard deviations below 50 %.

Figure A1 shows that, away from the upper and lower altitude limits, where retrieval errors are typically largest, the vertical smoothing has little to no effect on the weighted-average relative differences. At altitudes where the weighted-average correlation coefficients are greater than 0.8 and the weighted-average standard deviations are less than 50 % (altitude limits indicated by horizontal dotted lines in Fig. A1), the largest effect on the relative differences is in the NO₂ comparisons near 17 km. In this region the difference between the smoothed and non-smoothed relative difference is less than 9 %, and this difference is mainly due to coarser vertical resolution values for SCIAMACHY, and to a lesser extent MIPAS IMK-IAA, retrievals in this region. Otherwise, within the altitude limits mentioned above, differences between smoothed and non-smoothed relative differences are typically less than 1 %, as the vertical resolutions of most of the retrievals are on the same order.

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References

- ACE-FTS: ACE-FTS Level 2 data, available at: <http://www.ace.uwaterloo.ca>, last access: November 2016.
- Bauer, R., Rozanov, A., McLinden, C. A., Gordley, L. L., Lotz, W., Russell III, J. M., Walker, K. A., Zawodny, J. M., Ladstätter-Weißmayer, A., Bovensmann, H., and Burrows, J. P.: Validation of SCIAMACHY limb NO₂ profiles using solar occultation measurements, *Atmos. Meas. Tech.*, 5, 1059–1084, doi:10.5194/amt-5-1059-2012, 2012.
- Bernath, P. F., McElroy, C. T., Abrams, M. C., Boone, C. D., Butler, M., Camy-Peyret, C., Carleer, M., Clerboux, 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.
- Bogumil, K., Orphal, J., Voigt, S., Bovensmann, H., Fleischmann, O. C., Hartmann, M., Homann, T., Spietz, P., Vogel, A., and Burrows, J. P.: Reference spectra of atmospheric trace gases measured with the SCIAMACHY PFM satellite spectrometer, *Proc. 1st Europ. Sympos. Atmos. Meas. from Space (ESAMS-99)*, 2, 443–447, ESA-ESTEC, Noordwijk, 18–22 January 1999.
- 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., Walker, K. A., and Bernath, P. F.: Version 3 Retrievals for the Atmospheric Chemistry Experiment Fourier Transform Spectrometer (ACE-FTS), *The Atmospheric Chemistry Experiment ACE at 10: A Solar Occultation Anthology*, A. Deepak Publishing, Hampton, Virginia, USA, 103–127, 2013.
- Bracher, A., Sinnhuber, M., Rozanov, A., and Burrows, J. P.: Using a photochemical model for the validation of NO₂ satellite measurements at different solar zenith angles, *Atmos. Chem. Phys.*, 5, 393–408, doi:10.5194/acp-5-393-2005, 2005.
- Brasseur, G. P. and Solomon, S.: *Aeronomy of the middle atmosphere: chemistry and physics of the stratosphere and mesosphere*, Springer, Dordrecht, the Netherlands, 327–346, 2005.
- Brohede, S. M., Haley, C. S., McLinden, C. A., Sioris, C. E., Murtagh, D. P., Petelina, S. V., Llewellyn, E. J., Bazureau, A., Goutail, F., Randall, C. E., Lumpe, J. D., Taha, G., Thomasson, L. W., and Gordley, L. L.: Validation of Odin/OSIRIS stratospheric NO₂ profiles, *J. Geophys. Res.*, 112, D07310, doi:10.1029/2006JD007586, 2007a.
- Brohede, S. M., McLinden, C. A., Berthet, G., Haley, C. S., Murtagh, D., and Sioris, C. E.: A stratospheric NO₂ climatology from Odin/OSIRIS limb-scatter measurements, *Can. J. Phys.*, 85, 1253–1274, doi:10.1139/P07-141, 2007b.
- Bovensmann, H., Burrows, J. P., Buchwitz, M., Frerick, J., Noël, S., Rozanov, V., Chance, K., and Goede, A.: SCIAMACHY: Mission objectives and measurement modes, *J. Atmos. Sci.*, 56, 127–150, doi:10.1175/1520-0469(1999)056<0127:SMOAMM>2.0.CO;2, 1999.
- Burkholder, J. B., Sander, S. P., Abbatt, J. P. D., Barker, J. R., Huie, R. E., Kolb, C. E., Kurylo, M. J., Orkin, V. L., Wilmouth, D. M., and Wine, P. H.: *Chemical kinetics and photochemical data for use in atmospheric studies*, Jet Propulsion Laboratory Publications 15–10, Evaluation Number 18, 2015.
- Burrows, J. P., Hötzle, E., Goede, A., Visser, H., and Fricke, W.: SCIAMACHY – scanning imaging absorption spectrometer for atmospheric chartography, *Acta Astronaut.*, 35, 445–451, doi:10.1016/0094-5765(94)00278-T, 1995.
- Burrows, J. P., Dehn, A., Deters, B., Himmelmann, S., Richter, A., Voigt, S., and Orphal, J.: Atmospheric remote-sensing reference data from GOME: 1. Temperature-dependent absorption cross-sections of NO₂ in the 231–794 nm range, *J. Quant. Spectrosc. Ra.*, 60, 1025–1031, doi:10.1016/S0022-4073(97)00197-0, 1998.
- Burrows, J. P., Weber, M., Buchwitz, M., Rozanov, V., Ladstätter-Weißmayer, A., Richter, A., Debeek, R., Hoogen, R., Bramstedt, K., Eichmann, K.-U., Eisinger, M., and Perner, D.: The Global Ozone Monitoring Experiment (GOME): Mission Concept and First Scientific Results, *J. Atmos. Sci.*, 56, 151–175, doi:10.1175/15200469(1999)056<0151:TGOMEG>2.0.CO;2, 1999.
- Ceccherini, S.: Analytical determination of the regularization parameter in the retrieval of atmospheric vertical profiles, *Optic. Lett.*, 30, 2554–2556, 2005.
- Ceccherini, S., Belotti, C., Carli, B., Raspollini, P., and Ridolfi, M.: Technical Note: Regularization performances with the error consistency method in the case of retrieved atmospheric profiles, *Atmos. Chem. Phys.*, 7, 1435–1440, doi:10.5194/acp-7-1435-2007, 2007.
- Chahine, M. T.: Determination of the temperature profile in an atmosphere from its outgoing radiance, *J. Opt. Soc. Am.*, 58, 1634, doi:10.1364/JOSA.58.001634, 1968.
- DARTS: Data Archives and Transmission System, SMILES data, available at: <https://www.darts.isas.jaxa.jp/iss/smiles/>, last access: November 2016.
- ESA: European Space Agency, GOMOS data, available at: <https://earth.esa.int/web/guest/data-access>, last access: November 2016a.

- ESA: European Space Agency, MIPAS ESA data, available at: <https://earth.esa.int/web/guest/data-access>, last access: November 2016b.
- Fischer, H. and Oelhaf, H.: Remote sensing of vertical profiles of atmospheric trace constituents with MIPAS limb-emission spectrometers, *Appl. Opt.*, 35, 2787–2796, 1996.
- Fischer, H., Birk, M., Blom, C., Carli, B., Carlotti, M., von Clarmann, T., Delbouille, L., Dudhia, A., Ehnhalt, D., Endemann, M., Flaud, J. M., Gessner, R., Kleinert, A., Koopman, R., Langen, J., López-Puertas, M., Mosner, P., Nett, H., Oelhaf, H., Perron, G., Remedios, J., Ridolfi, M., Stiller, G., and Zander, R.: MIPAS: an instrument for atmospheric and climate research, *Atmos. Chem. Phys.*, 8, 2151–2188, doi:10.5194/acp-8-2151-2008, 2008.
- 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., Malat, 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., Wade-falk, N., Zirath, H., Zimmermann, P., and Zimmermann, R.: The Odin satellite. I. Radiometer design and test, *Astron. Astrophys.*, 402, L27–L34, doi:10.1051/0004-6361:20030335, 2003.
- Funke, B., López-Puertas, M., Gil-López, S., von Clarmann, T., Stiller, G. P., Fischer, H., and Kellmann, S.: Downward transport of upper atmospheric NO_x into the polar stratosphere and lower mesosphere during the Antarctic 2003 and Arctic 2002/2003 winters, *J. Geophys. Res.*, 110, D24308, doi:10.1029/2005JD006463, 2005a.
- Funke, B., López-Puertas, M., von Clarmann, T., Stiller, G. P., Fischer, H., Glatthor, N., Grabowski, U., Höpfner, M., Kellmann, S., Kiefer, M., Linden, A., Mengistu Tsidu, G., Milz, M., Steck, T., and Wang, D. Y.: Retrieval of stratospheric NO_x from 5.3 and 6.2 μm nonlocal thermodynamic equilibrium emissions measured by Michelson Interferometer for Passive Atmospheric Sounding (MIPAS) on Envisat, *J. Geophys. Res.*, 110, D09302, doi:10.1029/2004JD005225, 2005b.
- Funke, B., López-Puertas, M., Holt, L., Randall, C. E., Stiller, G. P., and von Clarmann, T.: Hemispheric distributions and interannual variability of NO_y produced by energetic particle precipitation in 2002–2012, *J. Geophys. Res.-Atmos.*, 119, 13565–13582, doi:10.1002/2014JD022423, 2014.
- Fussen, D., Vanhellefont, F., Dodion, J., Bingen, C., Walker, K. A., Boone, C. D., McLeod, S. D., and Bernath, P. F.: Initial intercomparison of ozone and nitrogen dioxide number density profiles retrieved by the ACE-FTS and GOMOS occultation experiments, *Geophys. Res. Lett.*, 32, L16S02, doi:10.1029/2005GL022468, 2005.
- GES DISC: Goddard Earth Sciences Data and Information Services Center, MLS data, available at: <http://disc.sci.gsfc.nasa.gov/Aura/data-holdings/MLS/index.shtml>, last access: November 2016.
- Gordley, L. L., Russell III, J. M., Mickley, L. J., Frederick, J. E., Park, J. H., Stone, K. A., Beaver, G. M., McInerney, J. M., Deaver, L. E., Toon, G. C., Murcray, F. J., Blatherwick, R. D., Gunson, M. R., Abbatt, J. P. D., Mauldin III, R. L., Mount, G. H., Sen, B., and Blavier, J.-F.: Validation of nitric oxide and nitrogen dioxide measurements made by the Halogen Occultation Experiment for UARS platform, *J. Geophys. Res.*, 101, 10241–10266, doi:10.1029/95JD02143, 1996.
- Griffioen, E. and Oikarinen, L.: LIMBTRAN: A pseudo three-dimensional radiative transfer model for the limb-viewing imager OSIRIS on the ODIN satellite, *J. Geophys. Res.*, 105, 29717–29730, doi:10.1029/2000JD900566, 2000.
- Haley, C. S. and Brohede, S.: Status of the Odin/OSIRIS stratospheric O₃ and NO₂ data products, *Can. J. Phys.*, 85, 1177–1194, doi:10.1139/p07-114, 2007.
- HALOE: HALOE data, available at: <http://haloe.gats-inc.com/download/index.php>, last access: November 2016.
- Hanke, M.: A regularizing Levenberg-Marquardt scheme, with applications to inverse groundwater filtration problems, *Inverse Probl.*, 13, 79–95, 1997.
- Höpfner, M., von Clarmann, T., Fischer, H., Funke, B., Glatthor, N., Grabowski, U., Kellmann, S., Kiefer, M., Linden, A., Milz, M., Steck, T., Stiller, G. P., Bernath, P., Blom, C. E., Blumenstock, Th., Boone, C., Chance, K., Coffey, M. T., Friedl-Vallon, F., Griffith, D., Hannigan, J. W., Hase, F., Jones, N., Jucks, K. W., Keim, C., Kleinert, A., Kouker, W., Liu, G. Y., Mahieu, E., Mellqvist, J., Mikuteit, S., Notholt, J., Oelhaf, H., Piesch, C., Reddman, T., Ruhnke, R., Schneider, M., Strandberg, A., Toon, G., Walker, K. A., Warneke, T., Wetzell, G., Wood, S., and Zander, R.: Validation of MIPAS CIONO₂ measurements, *Atmos. Chem. Phys.*, 7, 257–281, doi:10.5194/acp-7-257-2007, 2007.
- IUP: Universität Bremen, SCIAMACHY data, available at: <http://www.iup.uni-bremen.de/scia-arc/>, last access: November 2016.
- Kar, J., McElroy, C. T., Drummond, J. R., Zou, J., Nichitui, F., Walker, K. A., Randall, C. E., Nowlan, C. R., Dufour, D. G., Boone, C. D., Bernath, P. F., Trepte, C. R., and McLinden, C.: Initial comparison of ozone and NO₂ profiles from ACE-MAESTRO with balloon and satellite data, *J. Geophys. Res.*, 112, D16301, doi:10.1029/2006JD008242, 2007.
- Kerzenmacher, T., Wolff, M. A., Strong, K., Dupuy, E., Walker, K. A., Amekudzi, L. K., Batchelor, R. L., Bernath, P. F., Berthet, G., Blumenstock, T., Boone, C. D., Bramstedt, K., Brogniez, C., Brohede, S., Burrows, J. P., Catoire, V., Dodion, J., Drummond, J. R., Dufour, D. G., Funke, B., Fussen, D., Goutail, F., Griffith, D. W. T., Haley, C. S., Hendrick, F., Höpfner, M., Huret, N., Jones, N., Kar, J., Kramer, I., Llewellyn, E. J., López-Puertas, M., Manney, G., McElroy, C. T., McLinden, C. A., Melo, S., Mikuteit, S., Murtagh, D., Nichitui, F., Notholt, J., Nowlan, C., Piccolo, C., Pommereau, J.-P., Randall, C., Raspollini, P., Ridolfi, M., Richter, A., Schneider, M., Schrems, O., Silicani, M., Stiller, G. P., Taylor, J., Tétard, C., Toohey, M., Vanhellefont, F., Warneke, T., Zawodny, J. M., and Zou, J.: Validation of NO₂ and NO from the Atmospheric Chemistry Experiment (ACE), *Atmos. Chem. Phys.*, 8, 5801–5841, doi:10.5194/acp-8-5801-2008, 2008.
- Kikuchi, K., Nishibori, T., Ochiai, S., Ozeki, H., Irimajiri, Y., Kasai, Y., Koike, M., Manabe, T., Mizukoshi, K., Murayama, Y., Nagahama, T., Sano, T., Sato, R., Seta, M., Takahashi, C., Takayanagi, M., Masuko, H., Inatani, J., and Shiotani, M.: Overview and early results of the Superconducting Submillimeter-Wave Limb-Emission Sounder (SMILES), *J. Geophys. Res.*, 115, D23306, doi:10.1029/2010JD014379, 2010.

- KIT: Karlsruhe Institute of Technology, MIPAS IMK-IAA data, available at: <https://www.imk-asf.kit.edu/english/308.php>, last access: November 2016.
- Kyrölä, E., Tamminen, J., Leppelmeiera, G. W., Sofieva, V., Hassinen, S., Bertaux, J. L., Hauchecorne, A., Dalaudier, F., Cotb, C., Korablev, O., Fanton d'Andon, O., Barrot, G., Mangin, A., Théodores, B., Guirlet, M., Etanchaud, F., Snoeij, P., Koopman, R., Saavedra, L., Fraisse, R., Fussen, D., and Vanhellemont, F.: GOMOS on Envisat: an overview, *Adv. Space Res.*, 33, 1020–1028, doi:10.1016/S0273-1177(03)00590-8, 2004.
- Kyrölä, E., Tamminen, J., Sofieva, V., Bertaux, J. L., Hauchecorne, A., Dalaudier, F., Fussen, D., Vanhellemont, F., Fanton d'Andon, O., Barrot, G., Guirlet, M., Mangin, A., Blanot, L., Fehr, T., Saavedra de Miguel, L., and Fraisse, R.: Retrieval of atmospheric parameters from GOMOS data, *Atmos. Chem. Phys.*, 10, 11881–11903, doi:10.5194/acp-10-11881-2010, 2010.
- Livesey, N. J., Van Snyder, W., Read, W. G., and Wagner, P. A.: Retrieval algorithms for the EOS Microwave Limb Sounder (MLS), *IEEE Trans. Geosci. Remote S.*, 44, 1144–1155, doi:10.1109/TGRS.2006.872327, 2006.
- Livesey, N. J., Read, W. G., Froidevaux, L., Lambert, A., Manney, G. L., Pumphrey, H. C., Santee, M. L., Schwartz, M. J., Wang, S., Cofield, R. E., Cuddy, D. T., Fuller, R. A., Jarnot, R. F., Jiang, J. H., Knosp, B. W., Stek, P. C., Wagner, P. A., and Wu, D. L.: EOS MLS Version 3.3/3.4 Level 2 data quality and description document, Tech. Rep., Jet Propulsion Laboratory, Pasadena, CA USA, available at: <http://mls.jpl.nasa.gov> (last access: January 2016), 2013.
- Livesey, N. J., Read, W. G., Wagner, P. A., Froidevaux, L., Lambert, A., Manney, G. L., Millan Valle, L. F., Pumphrey, H. C., Santee, M. L., Schwartz, M. J., Wang, S., Fuller, R. A., Jarnot, R. F., Knosp, B. W., and Martinez, E.: Version 4.2x Level 2 data quality and description document, Rev. B, Jet Propulsion Laboratory, Pasadena, CA USA, available at: <http://mls.jpl.nasa.gov>, (last access: September 2016), 2015.
- Llewellyn, E. J., Lloyd, N. D., Degenstein, D. A., Gattinger, R. L., Petelina, S. V., Bourassa, A. E., Wiens, J. T., Ivanov, E. V., McDade, I. C., Solheim, B. H., McConnell, J. C., Haley, C. S., von Savigny, C., Sioris, C. E., McLinden, C. A., Griffioen, E., Kaminski, J., Evans, W. F. J., Puckrin, E., Strong, K., Wehrle, V., Hum, R. H., Kendall, D. J. W., Matsushita, J., Murtagh, D. P., Brohede, S., Stegman, J., Witt, G., Barnes, G., Payne, W. F., Piché, L., Smith, K., Warshaw, G., Deslauniers, D. L., Marchand, P., Richardson, E. H., King, R. A., Wevers, I., McCreath, W., Kyrölä, E., Oikarinen, L., Leppelmeier, G. W., Auvinen, H., Mégie, G., Hauchecorne, A., Lefèvre, F., de La Noë, J., Ricaud, P., Frisk, U., Sjöberg, F., von Schéele, F., and Nordh, L.: The OSIRIS instrument on the Odin spacecraft, *Can. J. Phys.*, 82, 411–422, doi:10.1139/P04-005, 2004.
- Lucke, R. L., Korwan, D. R., Bevilacqua, R. M., Hornstein, J. S., Shettle, E. P., Chen, D. T., Daehler, M., Lumpe, J. D., Fromm, M. D., Debrestian, D., Neff, B., Squire, M., König-Langlo, G., and Davies, J.: The Polar Ozone and Aerosol Measurement (POAM) III instrument and early validation results, *J. Geophys. Res.*, 104, 18785–18799, doi:10.1029/1999JD900235, 1999.
- Lumpe, J. D., Bevilacqua, R. M., Hoppel, K. W., and Randall, C. E.: POAM III Retrieval Algorithm and Error Analysis, *J. Geophys. Res.*, 107, 4575, doi:10.1029/2002JD002137, 2002.
- McElroy, C. T., Nowlan, C., Drummond, J., Bernath, P., Barton, D., Dufour, D., Midwinter, C., Hall, R., Ogyu, A., Ullberg, A., Wardle, D., Kar, J., Zou, J., Nichitiu, F., Boone, C., Walker, K., and Rowlands, N.: The ACE-MAESTRO instrument on SCISAT: description, performance, and preliminary results, *Appl. Opt.*, 46, 4341–4356, doi:10.1364/AO.46.004341, 2007.
- McLinden, C. A., Olsen, S. C., Hannegan, B., Wild, O., Prather, M. J., and Sundet, J.: Stratospheric ozone in 3-D models: A simple chemistry and the cross-tropopause flux, *J. Geophys. Res.*, 105, 14653–14665, doi:10.1029/2000JD900124, 2002.
- Murtagh, D., Frisk, U., Merino, F., Ridal, M., Jonsson, A., Stegman, J., Witt, G., Eriksson, P., Jiménez, C., Megie, G., de la Noë, J., Ricaud, P., Baron, P., Pardo, J. R., Hauchecorne, A., Llewellyn, E. J., Degenstein, D. A., Gattinger, R. L., Lloyd, N. D., Evans, W. F. J., McDade, I. C., Haley, C. S., Sioris, C., von Savigny, C., Solheim, B. H., McConnell, J. C., Strong, K., Richardson, E. H., Leppelmeier, G. W., Kyrölä, E., Auvinen, H., and Oikarinen, L.: An overview of the Odin atmospheric mission, *Can. J. Phys.*, 80, 309–319, doi:10.1139/P01-157, 2002.
- NASA: POAM III data, available at: https://eosweb.larc.nasa.gov/project/poam3/poam3_table, last access: November 2016.
- NASA: SAGE III data, available at: https://eosweb.larc.nasa.gov/project/sage3/sage3_table, last access: November 2016.
- Odin/SMR: Odin/SMR Project Portal, SMR data, available at: <http://odin.rss.chalmers.se>, last access: November 2016.
- Olsen, S., McLinden, C. A., and Prather, M. J.: Stratospheric N₂O-NO_y system: Testing uncertainties in a three-dimensional framework, *J. Geophys. Res.*, 106, 28771–28784, doi:10.1029/2001JD000559, 2001.
- Pickett, H. M., Poynter, R. L., Cohen, E. A., Delitsky, M. L., Pearson, J. C., and Müller, H. S. P.: Submillimeter, millimeter, and microwave spectral line catalogue, *J. Quant. Spectrosc. Ra.*, 60, 883–890, doi:10.1016/S0022-4073(98)00091-0, 1998.
- Polyakov, A. V., Timofeyev, Y. M., Ionov, D. V., Virolainen, Y. A., Steele, H. M., and Newchurch, M. J.: Retrieval of ozone and nitrogen dioxide concentrations from Stratospheric Aerosol and Gas Experiment III (SAGE III) measurements using a new algorithm, *J. Geophys. Res.*, 110, D06303, doi:10.1029/2004JD005060, 2005.
- Popp, C., Wang, P., Brunner, D., Stammes, P., Zhou, Y., and Grzegorski, M.: MERIS albedo climatology for FRESCO+ O₂ A-band cloud retrieval, *Atmos. Meas. Tech.*, 4, 463–483, doi:10.5194/amt-4-463-2011, 2011.
- Prather, M. J.: Catastrophic loss of stratospheric ozone in dense volcanic clouds, *J. Geophys. Res.*, 97, 10187–10191, doi:10.1029/92JD00845, 1997.
- Randall, C. E., Rusch, W., Bevilacqua, R. M., Hoppel, K. W., and Lumpe, J.: Polar Ozone and Aerosol Measurement (POAM) II stratospheric NO₂, 1993–1996, *J. Geophys. Res.*, 103, 28361–28371, doi:10.1029/98JD02092, 1998.
- Randall, C. E., Lumpe, J. D., Bevilacqua, R. M., Hoppel, K. W., Shettle, E. P., Rusch, D. W., Gordley, L. L., Kreher, K., Pfeilsticker, K., Boesch, H., Toon, G., Goutail, F., and Pommereau, J.-P.: Validation of POAM III NO₂ measurements, *J. Geophys. Res.*, 107, 4432, doi:10.1029/2001JD001520, 2002.
- Randall, C. E., Harvey, V. L., Singleton, C. S., Bailey, S., Bernath, P. F., Codrescu, M., Nakajima, H., and Russell III, J. M.: Energetic particle precipitation effects on the Southern Hemisphere

- stratosphere in 1992–2005, *J. Geophys. Res.*, 112, D08308, doi:10.1029/2006JD007696, 2007.
- Randall, C. E., Harvey, V. L., Siskind, D. E., France, J., Bernath, P. F., Boone, C. D., and Walker, K. A.: NO_x descent in the Arctic middle atmosphere in early 2009, *Geophys. Res. Lett.*, 36, L18811, doi:10.1029/2009GL039706, 2009.
- Raspollini, P., Belotti, C., Burgess, A., Carli, B., Carlotti, M., Ceccherini, S., Dinelli, B. M., Dudhia, A., Flaud, J.-M., Funke, B., Höpfner, M., López-Puertas, M., Payne, V., Piccolo, C., Remedios, J. J., Ridolfi, M., and Spang, R.: MIPAS level 2 operational analysis, *Atmos. Chem. Phys.*, 6, 5605–5630, doi:10.5194/acp-6-5605-2006, 2006.
- Raspollini, P., Carli, B., Carlotti, M., Ceccherini, S., Dehn, A., Dinelli, B. M., Dudhia, A., Flaud, J.-M., López-Puertas, M., Niro, F., Remedios, J. J., Ridolfi, M., Sembhi, H., Sgheri, L., and von Clarmann, T.: Ten years of MIPAS measurements with ESA Level 2 processor V6 – Part 1: Retrieval algorithm and diagnostics of the products, *Atmos. Meas. Tech.*, 6, 2419–2439, doi:10.5194/amt-6-2419-2013, 2013.
- Rodgers, C. D.: *Inverse Methods for Atmospheric Sounding*, World Sci., Hackensack, New Jersey, USA, ISBN-13: 9789810227401, ISBN-10: 981022740X, 2008.
- Rothman, L. S., Jacquemart, D., Barbe, A., Chris Benner, D., Birk, M., Browne, L. R., Carleer, M. R., Chackerian Jr., C., Chance, K., Coudert, L. H., Dana, V., Devi, V. M., Flaud, J.-M., Gamache, R. R., Goldman, A., Hartmann, J.-M., Jucks, K. W., Maki, A. G., Mandin, J.-Y., Massien, S. T., Orphal, J., Perrin, A., Rinsland, C. P., Smith, M. A. H., Tennyson, J., Tolchenov, R. N., Toth, R. A., Vander Auwera, J., Varanasiq, P., and Wagner, G.: The HITRAN 2004 molecular spectroscopic database, *J. Quant. Spectrosc. Ra.*, 96, 139–204, doi:10.1016/j.jqsrt.2004.10.008, 2005.
- Roazanov, A., Bovensmann, H., Bracher, A., Hrechanyy, S., Roazanov, V., Sinnhuber, M., Stroth, F., and Burrows, J. P.: NO₂ and BrO vertical profile retrieval from SCIAMACHY limb measurements: Sensitivity studies, *Adv. Space Res.*, 6, 846–854, doi:10.1016/j.asr.2005.03.013, 2005.
- Russell III, J. M., Gordley, L. L., Park, J. H., Drayson, S. R., Hesketh, W. D., Cicerone, R. J., Tuck, A. F., Frederick, J. E., Harries, J. E., and Crutzen, P. J.: The Halogen Occultation Experiment, *J. Geophys. Res.*, 98, 10777–10797, doi:10.1029/93JD00799, 1993.
- SAGE III ATBD Team: SAGE III Algorithm Theoretical Basis Document (ATBD) for transmission level 1b products version 2.1, Tech. rep., NASA Langley Res. Cent. (LaRC), Hampton, Virginia, 2002a.
- SAGE III ATBD Team: SAGE III Algorithm Theoretical Basis Document (ATBD) Solar and Lunar Algorithm version 2.1, Tech. rep., NASA Langley Res. Cent. (LaRC), Hampton, Virginia, 2002b.
- Sander, S. P., Friedl, R. R., Golden, D. M., Kurylo, M. J., Huie, R. E., Orkin, V. L., Moortgat, G. K., Ravishankara, A. R., Kolb, C. E., Molina, M. J., and Finlayson-Pitts, B. J.: Chemical kinetics and photochemical data for use in atmospheric studies, *Jet Propulsion Laboratory Publications 02-25*, Evaluation Number 14, 2003.
- Santee, M. L., Lambert, A., Read, W. G., Livesey, N. J., Cofield, R. E., Cuddy, D. T., Daffer, W. H., Drouin, B. J., Froidevaux, L., Fuller, R. A., Jarnot, R. F., Knosp, B. W., Manney, G. L., Perun, V. S., Snyder, W. V., Stek, P. C., Thurstans, R. P., Wagner, P. A., Waters, J. W., Muscari, G., de Zafra, R. L., Dibb, J. E., Fahey, D. W., Popp, P. J., Marcy, T. P., Jucks, K. W., Toon, G. C., Stachnik, R. A., Bernath, P. F., Boone, C. D., Walker, K. A., Urban, J., and Murtagh, D.: Validation of the Aura Microwave Limb Sounder HNO₃ measurements, *J. Geophys. Res.*, 112, D24S40, doi:10.1029/2007JD008721, 2007.
- 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.
- Takahashi, C., Ochiai, S., and Suzuki, M.: Operational retrieval algorithms for JEM/SMILES level 2 data processing system, *J. Quant. Spectrosc. Ra.*, 111, 160–173, doi:10.1016/j.jqsrt.2009.06.005, 2010.
- Tikhonov, A. N.: On the solution of incorrectly stated problems and a method of regularization, *Dokl. Acad. Nauk SSSR*, 151, 501, 1963.
- University of Saskatchewan: OSIRIS data, available at: <http://odin-osiris.usask.ca>, last access: November 2016.
- Urban, J., Baron, P., Lautié, N., Dassas, K., Schneider, N., Ricaud, P., and de La Noë, J.: MOLIERE (v5): A versatile forward and inversion model for the millimeter and sub-millimeter wavelength range, *J. Quant. Spectrosc. Ra.*, 83, 529–554, doi:10.1016/S0022-4073(03)00104-3, 2004.
- Urban, J., Lautié, N., Le Flochmoën, E., Jiménez, C., Eriksson, P., de La Noë, J., Dupuy, E., El Amraoui, L., Frisk, U., Jégou, F., Murtagh, D., Olberg, M., Ricaud, P., Camy-Peyret, C., Dufour, G., Payan, S., Huret, N., Pirre, M., Robinson, A. D., Harris, N. R. P., Bremer, H., Kleinböhl, A., Küllmann, K., Künzi, K., Kuttippurath, J., Ejiri, M. K., Nakajima, H., Sasano, Y., Sugita, T., Yokota, T., Piccolo, C., Raspollini, P., and Ridolfi, M.: Odin/SMR limb observations of stratospheric trace gases: Level 2 processing of ClO, N₂O, HNO₃, and O₃, *J. Geophys. Res.*, 110, D14307, doi:10.1029/2004JD005741, 2005.
- Urban, J., Pommier, M., Murtagh, D. P., Santee, M. L., and Orsolini, Y. J.: Nitric acid in the stratosphere based on Odin observations from 2001 to 2009 – Part 1: A global climatology, *Atmos. Chem. Phys.*, 9, 7031–7044, doi:10.5194/acp-9-7031-2009, 2009.
- Verronen, P. T., Ceccherinib, S., Cortesib, U., Kyrölä, E., and Tamminen, J.: Statistical comparison of night-time NO₂ observations in 2003–2006 from GOMOS and MIPAS instruments, *Adv. Space Res.*, 43, 1918–1925, doi:10.1016/j.asr.2009.01.027, 2009.
- von Clarmann, T., Höpfner, M., Kellmann, S., Linden, A., Chauhan, S., Funke, B., Grabowski, U., Glatthor, N., Kiefer, M., Schieferdecker, T., Stiller, G. P., and Versick, S.: Retrieval of temperature, H₂O, O₃, HNO₃, CH₄, N₂O, ClONO₂ and ClO from MIPAS reduced resolution nominal mode limb emission measurements, *Atmos. Meas. Tech.*, 2, 159–175, doi:10.5194/amt-2-159-2009, 2009.
- Wang, D. Y., Höpfner, M., Mengistu Tsidu, G., Stiller, G. P., von Clarmann, T., Fischer, H., Blumenstock, T., Glatthor, N., Grabowski, U., Hase, F., Kellmann, S., Linden, A., Milz, M., Oelhaf, H., Schneider, M., Steck, T., Wetzels, G., López-Puertas, M., Funke, B., Koukouli, M. E., Nakajima, H., Sugita, T., Irie, H., Urban, J., Murtagh, D., Santee, M. L., Toon, G., Gunson, M. R., Irion, F. W., Boone, C. D., Walker, K., and Bernath, P. F.: Validation of nitric acid retrieved by the IMK-IAA processor from MIPAS/ENVISAT measurements, *Atmos. Chem. Phys.*, 7, 721–738, doi:10.5194/acp-7-721-2007, 2007a.

- Wang, D. Y., Höpfner, M., Blom, C. E., Ward, W. E., Fischer, H., Blumenstock, T., Hase, F., Keim, C., Liu, G. Y., Mikuteit, S., Oelhaf, H., Wetzels, G., Cortesi, U., Mencaraglia, F., Bianchini, G., Redaelli, G., Pirre, M., Catoire, V., Huret, N., Vigouroux, C., De Mazière, M., Mahieu, E., Demoulin, P., Wood, S., Smale, D., Jones, N., Nakajima, H., Sugita, T., Urban, J., Murtagh, D., Boone, C. D., Bernath, P. F., Walker, K. A., Kuttippurath, J., Kleinböhl, A., Toon, G., and Piccolo, C.: Validation of MIPAS HNO₃ operational data, *Atmos. Chem. Phys.*, 7, 4905–4934, doi:10.5194/acp-7-4905-2007, 2007b.
- Waters, J. W., Froidevaux, L., Harwood, R. S., Jarnot, R. F., Pickett, H. M., Read, W. G., Siegel, P. H., Cofield, R. E., Filipiak, M. J., Flower, D. A., Holden, J. R., Lau, G. K., Livesey, N. J., Manney, G. L., Pumphrey, H. C., Santee, M. L., Wu, D. L., Cuddy, D. T., Lay, R. R., Loo, M. S., Perun, V. S., Schwartz, M. J., Stek, P. C., Thurstans, R. P., Boyles, M. A., Chandra, K. M., Chavez, M. C., Chen, G. S., Chudasama, B. V., Dodge, R., Fuller, R. A., Girard, M. A., Jiang, J. H., Jiang, Y. B., Knosp, B. W., LaBelle, R. C., Lam, J. C., Lee, K. A., Miller, D., Oswald, J. E., Patel, N. C., Pukala, D. M., Quintero, O., Scaff, D. M., Van Snyder, W., Tope, M. C., Wagner, P. A., and Walch, M. J.: The Earth observing system microwave limb sounder (EOS MLS) on the Aura Satellite, *IEEE Trans. Geosci. Remote S.*, 44, 1075–1092, doi:10.1109/TGRS.2006.873771, 2006.
- Wetzels, G., Bracher, A., Funke, B., Goutail, F., Hendrick, F., Lambert, J.-C., Mikuteit, S., Piccolo, C., Pirre, M., Bazureau, A., Belotti, C., Blumenstock, T., De Mazière, M., Fischer, H., Huret, N., Ionov, D., López-Puertas, M., Maucher, G., Oelhaf, H., Pommereau, J.-P., Ruhnke, R., Sinnhuber, M., Stiller, G., Van Roozendaal, M., and Zhang, G.: Validation of MIPAS-ENVISAT NO₂ operational data, *Atmos. Chem. Phys.*, 7, 3261–3284, doi:10.5194/acp-7-3261-2007, 2007.
- Wetzels, G., Oelhaf, H., Friedl-Vallon, F., Kleinert, A., Maucher, G., Nordmeyer, H., and Orphal, J.: Long-term intercomparison of MIPAS additional species ClONO₂, N₂O₅, CFC-11, and CFC-12 with MIPAS-B measurements, *Ann. Geophys.-Italy*, 56, Fast Track-1, doi:10.4401/ag-6329, 2013.
- Wolff, M. A., Kerzenmacher, T., Strong, K., Walker, K. A., Toohey, M., Dupuy, E., Bernath, P. F., Boone, C. D., Brohede, S., Catoire, V., von Clarmann, T., Coffey, M., Daffer, W. H., De Mazière, M., Duchatelet, P., Glatthor, N., Griffith, D. W. T., Hannigan, J., Hase, F., Höpfner, M., Huret, N., Jones, N., Jucks, K., Kagawa, A., Kasai, Y., Kramer, I., Küllmann, H., Kuttippurath, J., Mahieu, E., Manney, G., McElroy, C. T., McLinden, C., Mébarki, Y., Mikuteit, S., Murtagh, D., Piccolo, C., Raspollini, P., Ridolfi, M., Ruhnke, R., Santee, M., Senten, C., Smale, D., Tétard, C., Urban, J., and Wood, S.: Validation of HNO₃, ClONO₂, and N₂O₅ from the Atmospheric Chemistry Experiment Fourier Transform Spectrometer (ACE-FTS), *Atmos. Chem. Phys.*, 8, 3529–3562, doi:10.5194/acp-8-3529-2008, 2008.