

Validation of NO₂ and HNO₃ measurements from the Improved Limb Atmospheric Spectrometer (ILAS) with the version 5.20 retrieval algorithm

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[1] The Improved Limb Atmospheric Spectrometer (ILAS) on board the Advanced Earth Observing Satellite (ADEOS) measured nitrogen dioxide (NO₂) and nitric acid (HNO₃) profiles from November 1996 to June 1997 at high latitudes in both hemispheres. The ILAS NO₂ profiles (version 5.20) are compared with those obtained by balloon-borne and satellite measurements to validate ILAS NO₂ data. Comparisons with balloon-borne measurements indicate that ILAS NO₂ at 25–30 km has a positive bias of 0.3–0.4 ppbv (6–11%). The random difference in NO₂ at 25–30 km is 0.2–0.3 ppbv (3–9%). The random error in the ILAS NO₂ measurements is larger than 100% below 20 km and above 45 km, where the NO₂ mixing ratios were less than 1.0 ppbv. It is possible that ILAS NO₂ values were lowered by optically thick aerosols with aerosol extinction coefficients at 780 nm of greater than 0.001 km⁻¹. The lack of diurnal correction along the line of sight contributes to the positive bias in the ILAS NO₂ values below 25 km. Agreement of the ILAS NO₂ values with those by the Polar Ozone and Aerosol Measurement (POAM) II instrument is within 10–30% at 25–35 km. The agreement with the Halogen Occultation Experiment (HALOE) is as good as ±10% at 25–40 km. ILAS HNO₃ (version 5.20) agrees with balloon-borne HNO₃ to within 0.1 ppbv (0–1%), and the random difference is within 10% at 25–30 km. *INDEX TERMS*: 0340 Atmospheric Composition and Structure: Middle atmosphere—composition and chemistry; 0341 Atmospheric Composition and Structure: Middle atmosphere—constituent transport and chemistry (3334); 0394 Atmospheric Composition and Structure: Instruments and techniques

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1. Introduction

[2] It is well known that NO_x (= NO + NO₂) destroys stratospheric ozone via the following catalytic cycle [*Crutzen*, 1970]:



The NO_x catalytic cycle is the dominant mechanism of ozone loss in the middle stratosphere [*Jucks et al.*, 1996; *Osterman et al.*, 1997] and also in the lower stratosphere during polar summer [*Fahey et al.*, 2000]. In addition, NO₂ regulates the stratospheric ozone budget via reactions with radicals (ClO, BrO, OH, and NO₃) to form reservoir species (ClONO₂, BrONO₂, HNO₃, and N₂O₅). Therefore, the

Table 1. List of Balloon Experiments

Experiment	Date	Location at 20 km	SZA	Distance, ^a km	Time Difference, ^a hours
LPMA	Feb. 26	66.9°N, 20.5°E	~90°	570	0.6
SAOZ	Feb. 24	67.4°N, 19.2°E	~90°	1010	1.8
SAOZ	March 20	66.9°N, 17.6°E	~90°	620	0.9
MIPAS-B2	March 24	69.6°N, 30.1°E	~105°	200	3.6
MkIV	May 8	68.6°N, 146.3°W	~90° ^b	760	6.4

^a Values between the ILAS and balloon-borne measurement locations.

^b MkIV measurement was made in the morning, whereas the ILAS measurement was made in the evening.

observation of NO₂ in the stratosphere is important in studies regarding ozone depletion.

[3] The Improved Limb Atmospheric Spectrometer (ILAS) on board the Advanced Earth Observing Satellite (ADEOS) is a solar occultation sensor. ILAS measured concentrations of atmospheric constituents including NO₂ in the stratosphere at high latitudes in the Northern Hemisphere (NH) and Southern Hemisphere (SH) from November 1996 through June 1997. A balloon-borne measurement campaign for the validation of ILAS data was conducted at Kiruna, Sweden (68°N, 21°E) in February and March 1997, and Fairbanks, Alaska (65°N, 148°W) in May 1997 [Kanzawa *et al.*, 1997]. Vertical profiles of NO₂ were obtained during this campaign.

[4] In the present paper, NO₂ profiles obtained by ILAS are compared with those obtained by several balloon-borne and satellite measurements, and the quality of ILAS NO₂ retrieved using the version 5.20 algorithm is described. Because of diurnal variations in NO₂ concentration, the comparisons are made under similar instantaneous sunlight conditions. To minimize the effect of the differences in the measurement locations, the comparisons of NO₂ for winter and early spring are made only when the ILAS and correlative measurements were made at similar potential vorticity (PV) values. To make consistent evaluations of ILAS reactive nitrogen measurements, similarly to NO₂, the quality of version 5.20 ILAS HNO₃ measurements is briefly reported in the appendix, while Koike *et al.* [2000] have already reported the quality of version 3.10 ILAS HNO₃ data.

2. ILAS

[5] The ADEOS satellite carrying ILAS was launched into a 98.6°-inclination Sun-synchronous polar orbit, resulting in 14 ILAS measurements in each hemisphere per day. Routine ILAS measurements at sunset (SS) were performed from November 1996 to June 1997 in NH, and from November 1996 to March 1997 in SH, and at sunrise (SR) from April to June 1997 in SH [Sasano *et al.*, 1999a], with some measurements in September and October 1996. The latitudes of 57°–72°N and 64°–89°S were covered during these periods. The solar zenith angle (SZA) of the measurements was 90.2 ± 0.2° at 20–50 km. The horizontal width of the ILAS sampling volume is calculated to be 13 km from its field of view. Air density-weighted absorption of solar radiation occurred over an effective path length of about 220 km. Vertical resolutions estimated from the instantaneous field of view are 1.9, 2.5, 3.0, and 3.5 km at altitudes of 15, 25, 35, and 55 km, respectively [Yokota *et al.*, 2002]. The NO₂ mixing ratio was retrieved up to 50 km every 1 km using an onion-

peeling method [Yokota *et al.*, 2002]. NO₂ absorption at wavelengths between 6.2 and 6.3 μm was measured by ILAS.

[6] The sum of the systematic and random errors in retrieving NO₂ mixing ratios with the version 5.20 algorithm was estimated primarily from the errors in the spectral fitting and atmospheric temperature. The percentage value is about 20% at 30 km and becomes progressively larger at higher and lower altitudes [Yokota *et al.*, 2002]. Multiplying by the average of all NO₂ profiles measured by ILAS, the absolute values of the errors at 15–30 km and 30–50 km are calculated to be 0.5–1.5 and 1.5–2.5 ppbv, respectively. The systematic error of ILAS NO₂ data will be estimated from the comparisons with balloon-borne measurements in the present study.

3. Comparisons With Balloon-Borne Measurements

3.1. Approach

[7] The balloon-borne measurements used in this study are summarized in Table 1. The ILAS NO₂ measurements are compared with 4 balloon-borne measurements in 1997 (5 NO₂ profiles): (1) the Limb Profile Monitor of the Atmosphere (LPMA) measurement performed on 26 February, (2) the balloon-borne Système d'Analyse d'Observations Zénithales (SAOZ) measurements performed on 24 February and 20 March, (3) the Michelson Interferometer for Passive Atmospheric Sounding-Balloon-borne version 2 (MIPAS-B2) measurement performed on 24 March, and (4) the MkIV measurement performed on 8 May. These measurements in the validation campaign acquired NO₂ profiles on the same UT days as the ILAS measurements and at locations within ~1000 km distance of ILAS measurement points. A coincidence criterion in PV was taken as within ±10% of the PV at the ILAS measurements. Applying this PV criterion to the measurement points for each altitude, the altitude levels at which the difference in PV for the balloon and ILAS measurements

Table 2. Performance of the Balloon-Borne Measurements at 25 km

Experiment	Vertical Resolution, ^a km	Precision, ^b %	Accuracy, ^b %
LPMA	1.5	10	15
SAOZ	1.2	8	13
MIPAS-B2	1.2	11	15
MkIV	1.5	4	8

^a Estimated from the instantaneous field of view.

^b One-σ values of the volume mixing ratios.

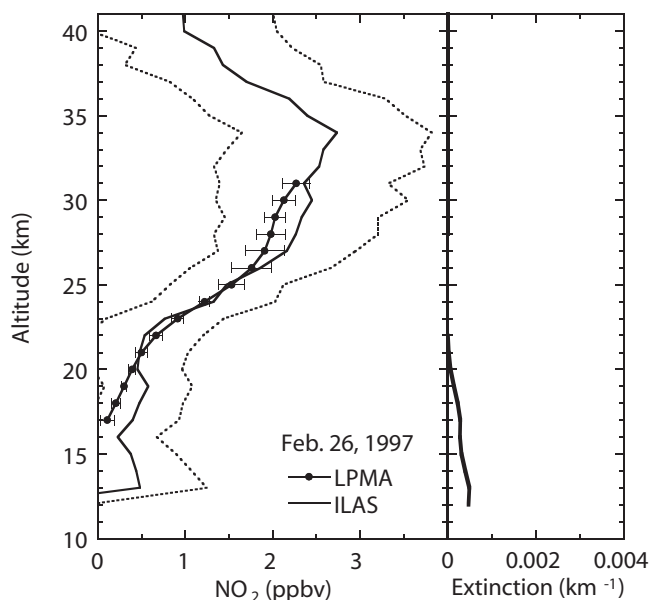


Figure 1. Comparison of ILAS with LPMA NO₂ measurements on 26 February 1997. Left panel: Thick line indicates the ILAS NO₂ profile. Dashed lines indicate the error of the ILAS NO₂ measurements. The balloon-borne NO₂ profile at SS is shown by the line with solid symbols. Associated bars indicate the error of the balloon-borne measurements. Right panel: Profile of the ILAS aerosol extinction coefficient at 780 nm.

was larger than 10% were rejected. Since the PV differences between equatorward and poleward edges of the Arctic vortex boundary region were usually about 20–30% in NH during the winter considered here [e.g., Koike *et al.*, 2000], this criterion is narrow enough to exclude the comparison of air masses having different characteristics, such as between air masses inside and outside the vortex. To make comparisons under the same SZA conditions with distinction of SS and SR as the ILAS measurements, MIPAS-B2 and MkIV profiles were diurnally corrected by a model calculation as described in sections 3.2.3 and 3.2.4, respectively.

[8] Heterogeneous reactions on the surface of polar stratospheric cloud (PSC) particles convert NO₂ to HNO₃, lowering the NO₂ concentration. Air masses sampled by balloon-borne instruments and ILAS were sometimes colder than the nitric acid trihydrate condensation temperature (T_{NAT}). The effect of heterogeneous reactions has been assessed by the period when these air masses were cooled to below T_{NAT} . Using the European Centre for Medium-Range Weather Forecasts (ECMWF) wind and temperature fields, 10-day isentropic back trajectories of air masses at ILAS and balloon-borne measurement points were calculated. T_{NAT} was calculated following the method described by Hanson and Mauersberger [1988].

[9] For all comparisons except for SAOZ on 24 February, the balloon-borne and ILAS air masses experienced temperatures below T_{NAT} for periods as short as 0–1 day. Air masses sampled by ILAS at 16–24 km and SAOZ at 15–23 km on 24 February experienced temperatures below T_{NAT} for 43 hours over 10 days, suggesting similar reduc-

tions in NO₂ concentrations by heterogeneous reactions on PSCs for ILAS and SAOZ measurements. Therefore, the present study does not make any corrections for heterogeneous reactions on PSCs.

[10] The ILAS NO₂ mixing ratios used for the comparison were linearly interpolated to the potential temperature levels corresponding to altitudes at which balloon-borne measurements were made. The United Kingdom Meteorological Office (UKMO) pressure and temperature were used for this calculation.

3.2. NO₂ Profiling by Balloon-Borne Measurements

3.2.1. LPMA

[11] The LPMA instrument developed at Laboratoire de Physique Moléculaire et Applications is a Fourier transform infrared (FTIR) spectrometer that measures the vertical profiles of various species from solar occultation spectra [Camy-Peyret, 1995]. The LPMA NO₂ vertical profile measurement was performed on 26 February 1997 near Kiruna, Sweden, during SS (SZA $\sim 90^\circ$). The 170 spectra recorded during the flight showed sufficient absorption by NO₂ molecules for the precise retrieval in the appropriate microwindow around 3.4 μm (2915 cm^{-1}). A global fit algorithm associated with an efficient minimization algorithm of the Levenberg-Marquardt type allows the retrieval of vertical profiles of NO₂ from the occultation spectra [Payan *et al.*, 1998].

3.2.2. SAOZ

[12] The SAOZ sonde is a UV-visible diode array spectrometer derived from the ground-based SAOZ designed for the monitoring of ozone and NO₂ total columns [Pommereau and Goutail, 1988]. SAOZ NO₂ vertical profile measurements were performed on 24 February and 20 March 1997 from Kiruna, Sweden, during SS (SZA $\sim 90^\circ$). NO₂ was measured by differential spectroscopy over a large spectral range of 410–530 nm using the absorption cross sections at 220 K of Vandaele *et al.*

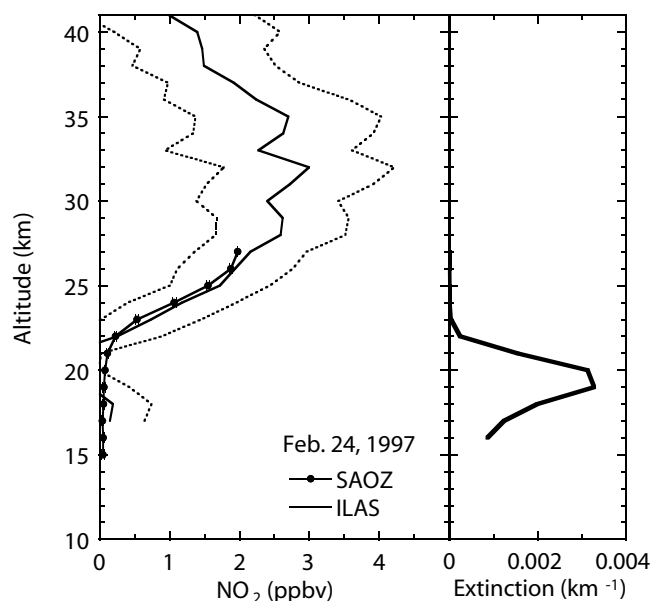


Figure 2. Same as Figure 1, but for the comparison with the SAOZ NO₂ measurements on 24 February 1997.

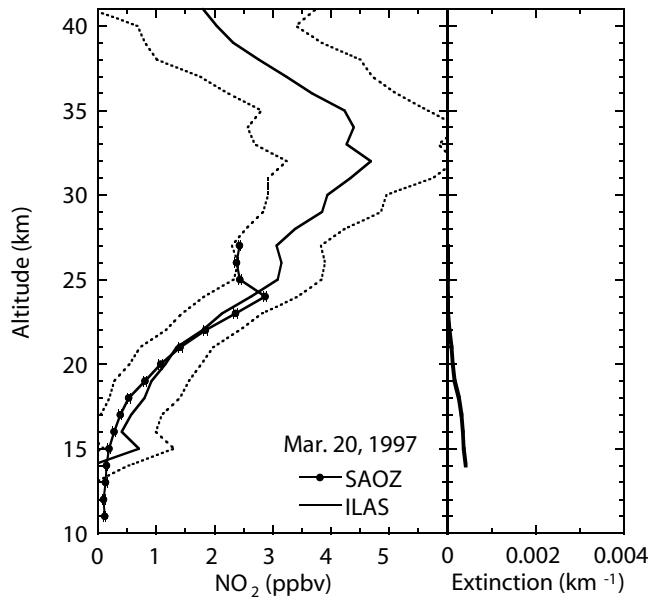


Figure 3. Same as Figure 1, but for the comparison with the SAOZ NO₂ measurements on 20 March 1997.

[1998]. Concentration profiles were retrieved by a linear tangent ray inversion technique (onion peeling).

3.2.3. MIPAS-B2

[13] The MIPAS-B2 instrument is a cryogenic FTIR spectrometer that measures atmospheric thermal emissions from the limb [Oelhaf et al., 1996; Friedl-Vallon et al., 1999]. The MIPAS-B2 NO₂ vertical profile measurement was performed on 24 March 1997 near Kiruna, Sweden, during the nighttime (SZA ~105° at 20 km). NO₂ was measured using the ν₃ band between 6.2 and 6.3 μm (1585

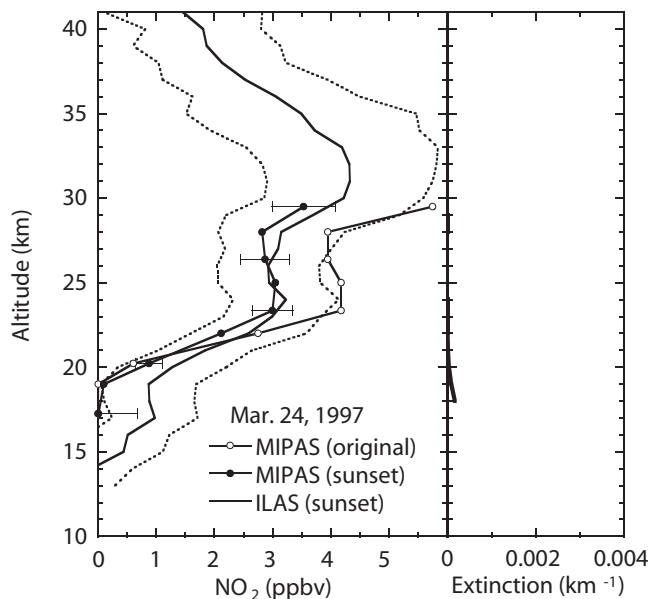


Figure 4. Same as Figure 1, but for the comparison with the MIPAS-B2 NO₂ measurements on 24 March 1997. The original balloon-borne NO₂ profile (SZA ~105°) is shown by the line with open symbols for reference.

and 1615 cm⁻¹). The vertical profile of NO₂ mixing ratios was retrieved by an onion-peeling technique and a multiparameter nonlinear least squares fitting procedure [Wetzel et al., 2002].

[14] The 3-dimensional Karlsruhe Simulation Model of the Middle Atmosphere (KASIMA) [Ruhnke et al., 1999] was used to convert the nighttime NO₂ values to those at SS at the ILAS measurement point, by multiplying the MIPAS NO₂ values with the ratio of the model NO₂ values for the time and location of ILAS and MIPAS-B2 measurements. The model uses the 6-hourly analyses of the ECMWF wind and temperature data that are dynamically interpolated in space and time to the model environment up to a pressure level of 10 hPa [Reddmann et al., 2001]. The model was initialized on 10 December 1996 using data from a 2-dimensional model (for details, see Wetzel et al. [2002]). The chemistry module consists of 58 chemical species and families, which are involved in 101 gas-phase reactions, 39 photodissociation reactions, and 10 heterogeneous reactions taking place on the surface of polar stratospheric clouds and liquid sulfuric acid aerosols. The rate constants of the gas phase reactions and the surface reaction probabilities are taken from the compilation of DeMore et al. [1997] with the update of Sander et al. [2000].

3.2.4. MkIV

[15] The MkIV instrument, which is a solar occultation FTIR spectrometer, measures the concentrations of various chemical species simultaneously [Toon, 1991; Sen et al., 1998]. The MkIV NO₂ vertical profile measurement was performed on 8 May 1997 from Fairbanks, Alaska, during SR (SZA ~90°), where the ILAS measurement in NH was made during SS. NO₂ was measured using the wavelengths of 6.1–6.3 and 3.4–3.5 μm. From a series of spectra measured at different tangent altitudes, the vertical profiles of chemical species were retrieved.

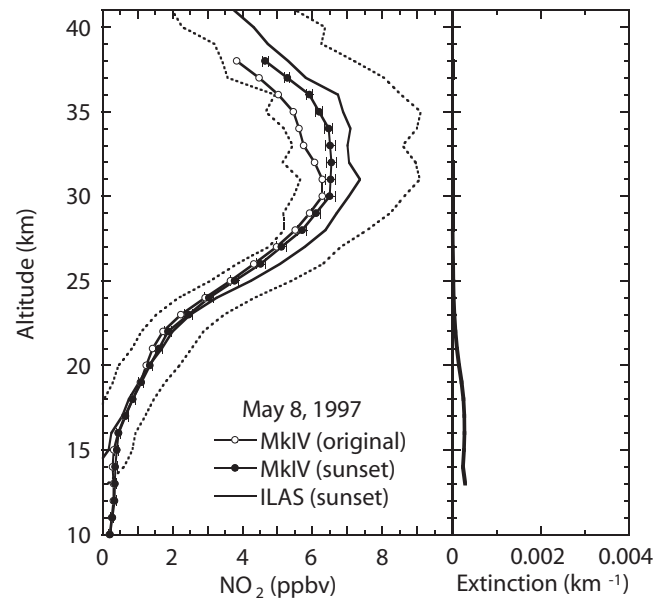


Figure 5. Same as Figure 1, but for the comparison with the MkIV NO₂ measurements on 8 May 1997. The original balloon-borne NO₂ profile (SZA ~90° in the morning) is shown by the line with open symbols for reference.

Table 3. Altitudes Used for the Comparison and Altitudes Where the $|\Delta\text{NO}_2|$ Value Was Greater Than 0.5 ppbv

Experiment	Date	Altitudes for Comparison	$ \Delta\text{NO}_2 > 0.5$ ppbv
LPMA	Feb. 26	17–19 km (boundary) 24–31 km (inside)	–
SAOZ	Feb. 24	16–27 km (inside)	19–21 km (ILAS low)
SAOZ	March 20	14–26 km (boundary)	25–26 km (ILAS high)
MIPAS-B2	March 24	18–29 km (inside)	18–20 km (ILAS high)
MkIV	May 8	15–38 km (outside)	25–37 km (ILAS high)

Altitudes used for the comparison have been chosen with PV differences less than 10%. Measurements in the vortex boundary and inside and outside the vortex are denoted by “boundary,” “inside,” and “outside,” respectively.

[16] A photochemical model [Osterman *et al.*, 1999, and references therein] was used to convert NO₂ values at SR to those at SS. The model is constrained by temperature, pressure, O₃, H₂O, CH₄, CO, NO_y, and Cl_y, inferred from MkIV measurements. The input O₃ and temperature profiles are taken from the measurements, including MkIV [Sen *et al.*, 1998]. The abundance of radical (e.g., NO, NO₂) and reservoir (e.g., HNO₃, N₂O₅) gases were calculated allowing for diurnal variation and assuming a balance between production and loss rates of each species integrated over a 24-hour period, for the latitude and temperature of the MkIV observation. Reaction rates and absorption cross sections were adopted from the JPL 97-4 compendium [DeMore *et al.*, 1997], except for the reaction HNO₃ + OH. The rate of HNO₃ + OH was adopted from Brown *et al.* [1999].

3.3. Performance of the Measurements

[17] Table 2 shows the vertical resolutions of the balloon-borne measurements used in the present study. These were estimated from the instantaneous field of view of the instrument to be 1.2–1.5 km at 25 km. Since the vertical resolution of the ILAS instrument (~ 2.5 km at 25 km) is of

the same order as the balloon-borne instruments, the influence of the difference in vertical resolutions on this comparison study is negligible.

[18] Table 2 also shows the precision and absolute accuracy of the balloon-borne measurements. The 1- σ precision was estimated primarily from the errors in the spectral fitting and the atmospheric temperature. The 1- σ absolute accuracy of the retrieval profiles was estimated primarily from the errors in the line parameters or the cross section of NO₂. It is noted that the error estimates for each instrument were determined independently by each group. They were not performed in a consistent manner and therefore do not represent a consensus or a basis for comparing instrument performance. This is because the method of estimating the error depends on the nature of the measurements (i.e., thermal emission, solar transmission, etc.) as well as on the retrieval technique employed. Consequently, the predominant error sources of error were sometimes different for the different measurements.

3.4. Difference in NO₂

[19] The ILAS NO₂ profiles and coincident balloon-borne NO₂ profiles are shown in Figures 1–5. In each figure, the right panel shows the profile of ILAS aerosol extinction coefficients at a wavelength of 780 nm measured simultaneously with NO₂. The left panel is for the comparison of NO₂ profiles. For comparisons with MIPAS-B2 and MkIV, the profiles with and without the correction for SZA

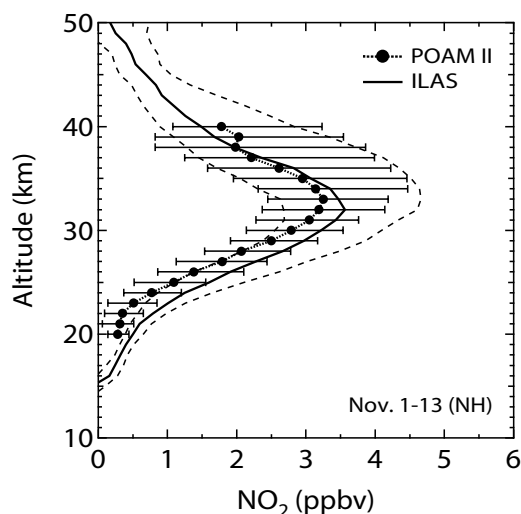


Figure 6. Comparison with the POAM II NO₂ measurements during 1–13 November in NH. The ILAS median NO₂ profile is shown with the solid line. Dashed lines indicate the 67% central values of ILAS NO₂. The POAM II median NO₂ profile is shown by the dotted line with symbols. Associated bars indicate the 67% central values of POAM II NO₂.

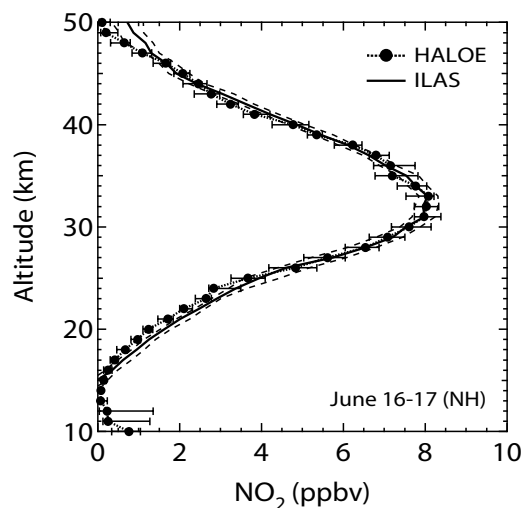


Figure 7. Same as Figure 6, but for the comparison with the HALOE NO₂ measurements during 16–17 June in NH.

Table 4. Time Period, Latitude, and Number of Profiles (*N*) Used for the Comparison of ILAS with POAM II

Time Period	Latitude		<i>N</i>	
	ILAS	POAM II	ILAS	POAM II
Nov. 1–13	70.2°–67.9°N	69.2°–67.3°N	154	122
Nov. 1–13 (outside)	74.8°–70.7°S	74.1°–70.0°S	16–87 ^a	11–60 ^a
Nov. 1–13 (boundary)	74.8°–70.7°S	74.1°–70.0°S	11–37 ^a	8–27 ^a
Nov. 1–13 (inside)	74.8°–70.7°S	74.1°–70.0°S	34–111 ^a	15–60 ^a

Measurements in the vortex boundary and inside and outside the vortex are denoted by “boundary,” “inside,” and “outside,” respectively.

^aThe number of profiles depends on the altitude, due to the vortex condition at each altitude.

by the model are denoted as “sunset” and “original,” respectively.

[20] The comparisons are made only for altitudes chosen with the PV criteria as discussed above. The altitudes used for comparisons are listed in Table 3. Table 3 also shows the altitudes where the $|\Delta\text{NO}_2|$ values are greater than 0.5 ppbv (~absolute error in retrieving the ILAS NO₂ during the winter), where ΔNO_2 is defined as NO₂ (ILAS) – NO₂ (correlative measurement).

[21] Some NO₂ mixing ratios measured by ILAS at 18–37 km were larger than the balloon measurements by 0.5–0.8 ppbv (Table 3). However, the ILAS NO₂ values at 19–21 km on 24 February were negative (Figure 2) and much smaller than those by SAOZ (Table 3).

[22] ILAS aerosol extinction coefficients indicate that optically thick aerosols (most likely PSC particles) were present at 16–25 km and centered at 19–20 km (Figure 2). Since aerosols have a continuous absorption that overlaps the infrared wavelength region for retrieving ILAS NO₂ mixing ratios [Yokota *et al.*, 2002], the retrieval algorithm must separate the contribution of aerosols from the total transmittance. The simulations with climatological NO₂ profiles and the theoretical extinction spectra, however, show the optically thick aerosols cause the apparent artificial reduction in ILAS NO₂ values because of the inadequate correction of aerosol effects employed in the operational data processing [Yokota *et al.*, 2002]. Consequently, the reduction in NO₂ at 19–21 km (Table 3) is attributed to interference from the optically thick aerosols, with extinction greater than 0.001 km⁻¹ (Figure 2).

4. Comparisons With Satellite Measurements

4.1. Approach

[23] The ILAS NO₂ measurements are compared with 2 satellite solar occultation measurements: (1) the Polar

Ozone and Aerosol Measurement (POAM) II in NH and SH during 1–13 November 1996 and (2) the Halogen Occultation Experiment (HALOE) in SH on 24 November, 15–16 December 1996 and 18–19 February 1997, and in NH on 25–31 March and 16–17 June 1997. For each period, these measurements were performed at locations within 1000 km in distance and 1° in latitude of the ILAS measurement points.

[24] For each altitude during winter and early spring, the daily PV values delineating the equatorward and poleward edges of the vortex boundary region are calculated from the maximum convex and concave curvature in the PV values plotted against equivalent latitude, following the method described by Nash *et al.* [1996]. On the basis of this calculation, NO₂ data at each altitude are classified as inside the vortex, vortex boundary region, and outside the vortex. This classification is made for data obtained at altitudes below 40 km, where the UKMO data are available.

[25] Median profiles for the NO₂ mixing ratios measured by ILAS, POAM II, and HALOE in each period and vortex condition are compared. For each comparison, the 67% central values of the median NO₂ mixing ratios measured by ILAS agree with those by POAM II and HALOE measurements to within 10% at 20–40 and 15–45 km, respectively (e.g., Figures 6 and 7), indicating the comparisons of median profiles are valid.

4.2. NO₂ Profiling by Satellite Measurements

4.2.1. POAM II

[26] The POAM II instrument was launched into a 98.7°-inclination orbit on board the French SPOT-3 satellite on 26 September 1993. POAM II made measurements using solar occultation with nine channels between 353 and 1060 nm until the middle of November 1996. The NO₂ value is derived from the differential signal

Table 5. Time Period, Latitude, and Number of Profiles (*N*) Used for the Comparison of ILAS with HALOE

Time Period	Latitude		<i>N</i>	
	ILAS	HALOE	ILAS	HALOE
Nov. 24	67.9°–68.1°S	68.4–68.9°S	7	11
Dec. 15–16	64.4°–64.6°S	63.9°–65.5°S	16	18
Feb. 18–19	75.0°–75.8°S	74.6°–76.1°S	24	18
March 25–31 (40–50 km)	68.1°–68.8°N	67.2°–69.1°N	92	90
March 25–31 (outside)	68.1°–68.8°N	67.2°–69.1°N	14–27 ^a	14–29 ^a
March 25–31 (boundary)	68.1°–68.8°N	67.2°–69.1°N	7–32 ^a	6–33 ^a
March 25–31 (inside)	68.1°–68.8°N	67.2°–69.1°N	19–73 ^a	28–89 ^a
June 16–17	56.9°–57.0°N	56.1°–57.7°N	23	11

Measurements in the vortex boundary and inside and outside the vortex are denoted by “boundary,” “inside,” and “outside,” respectively.

^aThe number of profiles depends on the altitude, due to the vortex condition at each altitude.

Table 6. Altitudes with $|\Delta\text{NO}_2| > 0.5$ ppbv for POAM II Measurements

Period	ILAS > POAM II	ILAS < POAM II
Nov. 1–13 (NH)	23, 25–26, and 28 km	–
Nov. 1–13 (SH, outside)	20–29 and 31 km	–
Nov. 1–13 (SH, boundary)	20, 23–26, 28–29, and 31–32 km	38 km
Nov. 1–13 (SH, inside)	21–22 km	36–37 km

Measurements in the vortex boundary and inside and outside the vortex are denoted by “boundary,” “inside,” and “outside,” respectively.

between two narrow-band channels at 442 and 448 nm. The retrieval altitude ranged between ~ 20 and 40 km. The ILAS NO₂ profiles are compared with those by POAM II (version 6) for 1–13 November 1996 in NH and SH as listed in Table 4.

4.2.2. HALOE

[27] The HALOE instrument was launched on board the Upper Atmosphere Research Satellite (UARS) on 12 September 1991. Routine HALOE observations started on 11 October 1991. Vertical profiles of NO₂ and several other trace species are derived using the solar occultation technique. Fifteen vertical profiles in each hemisphere are obtained each day. The ILAS NO₂ profiles are compared with those by HALOE (version 19) for five time periods: (1) 24 November, (2) 15–16 December 1996, (3) 18–19 February 1997 in SH, (4) 25–31 March, and (5) 16–17 June 1997 in NH (Table 5).

4.2.3. Difference in NO₂

[28] Figures 6 and 7 show representative comparisons of the ILAS NO₂ median profiles with those by POAM II (1–13 November in NH) and HALOE (16–17 June in NH), respectively. Table 6 shows the altitudes where the $|\Delta\text{NO}_2|$ values for the POAM II are greater than 0.5 ppbv for each comparison. Some ILAS NO₂ mixing ratios below 32 km are greater than the POAM II NO₂ by >0.5 ppbv, whereas some ILAS NO₂ mixing ratios above 36 km are smaller than the POAM II NO₂ by >0.5 ppbv.

[29] Table 7 shows the altitudes where the $|\Delta\text{NO}_2|$ values for HALOE are greater than 0.5 ppbv for each comparison. Some ILAS NO₂ mixing ratios above 37 km are greater than HALOE NO₂ by >0.5 ppbv, whereas some ILAS NO₂ mixing ratios below 13 km and 29–31 km are smaller than HALOE NO₂ by >0.5 ppbv. At 14–26 km,

the ILAS NO₂ mixing ratios are sometimes larger or smaller than HALOE NO₂ values by 0.5 ppbv.

[30] The ΔNO_2 values outside the vortex are compared with those inside the vortex for the POAM II and HALOE data. In Table 8, for each correlative measurement, the ΔNO_2 values averaged in every 5-km layer are shown for inside and outside the vortex. Also shown are the δNO_2 values defined as the ΔNO_2 divided by the NO₂ mixing ratios measured by a correlative measurement. For POAM II data, the ΔNO_2 values outside the Antarctic vortex are 0.9 ppbv (23%) and 0.5 ppbv (7%) at 25 and 30 km, respectively. The ΔNO_2 values outside the vortex are 0.5 ppbv (6%) larger than those inside the vortex at 25 and 30 km. For HALOE data, similarly, the ΔNO_2 values outside the Arctic vortex are <0.3 ppbv (3%) larger than those inside the vortex at 20 and 25 km, and smaller by <0.3 ppbv (7%) at 30 and 35 km. Considering that the estimated error of the ILAS NO₂ measurements is 0.5–1.5 ppbv (20%) at 20–30 km, as discussed in section 2, however, the differences in the ΔNO_2 and δNO_2 values between inside and outside the vortex are within the uncertainty of the ILAS measurements. Table 8 also shows the ΔNO_2 and δNO_2 values of HALOE data in NH and SH for the same season. The differences in the ΔNO_2 and δNO_2 values between NH and SH are <0.4 ppbv (10%) at 20–45 km, and are within the uncertainty of the measurements.

5. Discussion

[31] The ΔNO_2 and δNO_2 profiles for the comparisons with each balloon-borne and satellite measurement are shown in Figures 8 and 9 respectively. Table 9 shows the average δNO_2 values in each 5-km layer. For SAOZ data, the ΔNO_2 and δNO_2 values at each altitude were averaged from comparisons on 24 February and 20 March. In this calculation, the comparison at 16–25 km on 24 February was excluded because of the presence of optically thick aerosols, as discussed above. For the comparisons with profiles observed by MIPAS-B2 and MkIV, the profiles corrected by a model calculation were used. For the HALOE and POAM II measurements, profiles classified as outside the vortex, vortex boundary region, and inside the vortex were used below 40 km. Since the differences in the ΔNO_2 and δNO_2 values between inside and outside the vortex and between NH

Table 7. Altitudes with $|\Delta\text{NO}_2| > 0.5$ ppbv for HALOE Measurements

Period	ILAS > HALOE	ILAS < HALOE
Nov. 24 (SH)	39–41 km	10–14, 19, 24–25, and 29–30 km
Dec. 15–16 (SH)	16–17, 37–38, 41–43, and 48–50 km	10–12 and 30–31 km
Feb. 18–19 (SH)	14–19, 21, 23–26, 40–43, and 47–50 km	10–11 km
March 25–31 (NH, 40–50 km)	–	–
March 25–31 (NH, outside)	20 and 24 km	31 km
March 25–31 (NH, boundary)	–	–
March 25–31 (NH, inside)	–	12 km
16–17 June (NH)	24 and 48–50 km	11–14 km

Measurements in the vortex boundary and inside and outside the vortex are denoted by “boundary,” “inside,” and “outside,” respectively.

Table 8. The Comparison of the ΔNO_2 (ppbv) and δNO_2 (%) Values Between Outside and Inside the Vortex and Between NH and SH

Experiment	20 km	25 km	30 km	35 km	40 km	45 km
POAM II ^a (outside)	...	0.91 23%	0.52 7%	-0.20 -3%
POAM II ^a (inside)	...	0.43 17%	0.03 1%	-0.20 -3%
HALOE ^b (outside)	0.45 (69%) ^c	0.29 10%	-0.36 -6%	-0.04 -1%
HALOE ^b (inside)	0.19 (21%) ^c	0.18 7%	-0.09 -2%	0.18 6%
HALOE ^d (NH)	0.32 (27%) ^c	0.19 7%	0 0%	0 0%	0.17 5%	0.06 (4%) ^c
HALOE ^e (SH)	0.24 (13%) ^c	0.07 3%	-0.43 -5%	0.32 5%	0.53 15%	0.33 (20%) ^c

Measurements in the vortex boundary and inside and outside the vortex are denoted by “boundary,” “inside,” and “outside,” respectively.

^aOn 1–13 November in SH.

^bOn 25–31 March in NH.

^cThese values are very sensitive to the HALOE NO₂ values due to low NO₂ mixing ratios.

^dOn 16–17 June in NH.

^eOn 15–16 December in SH.

and SH were within the uncertainty of the ILAS measurements, as discussed above, the ΔNO_2 and δNO_2 values at each altitude were averaged for POAM II and HALOE data.

[32] Table 10 shows the systematic and random differences in NO₂ between the ILAS and balloon-borne measurements. These systematic and random differences are the average and standard deviation of the ΔNO_2 (δNO_2) values for all comparisons with the balloon-borne NO₂ data, except for the comparison with SAOZ at 16–25 km on 24 February.

[33] Random differences in NO₂ at 25 and 30 km were estimated to be 0.32 ppbv (9%) and 0.22 ppbv (3%), respectively (Table 10 and Figure 8). Systematic differences in NO₂ at 25 and 30 km were estimated to be 0.25 ppbv (6%) and 0.44 ppbv (11%), respectively, within the uncertainty of the balloon measurements (Table 2).

[34] NO₂ concentrations vary along the line of sight for occultation measurements because of the changing SZA. Sen *et al.* [1998] have compared NO₂ profiles with and without its diurnal correction using MkIV NO₂ measurements performed during SS at midlatitudes. The uncorrected NO₂ mixing ratio at 20 km was \sim 10% higher than the corrected mixing ratio, with progressively smaller differences at higher altitudes. No diurnal correction along the line of sight has been made in ILAS NO₂, whereas the correction has been made in HALOE NO₂. For 20–25 km, ILAS NO₂ is greater than HALOE by 6–27%, with progressively smaller differences at higher altitudes (Table 9 and Figure 9b). This result indicates that the lack of diurnal correction of NO₂ accounts for part of the positive bias in ILAS NO₂ below 25 km.

[35] Random differences between the ILAS and balloon measurements at 25 and 30 km are as small as 3–9% (Table 10). Furthermore, the δNO_2 values for HALOE data are within \pm 10% at 25–40 km (Table 9), suggesting that the lack of diurnal correction is not a significant source of uncertainty above 25 km.

[36] Although the absolute value of random differences between ILAS and balloon-borne measurements at 20 km is as small as 0.3 ppbv (Table 10 and Figure 8a), the percentage value at 20 km is as large as 400% (Table 10

and Figure 8b). The δNO_2 values for SAOZ and MkIV data at 20 km are smaller than those for LPMA and MIPAS-B2 (Table 9). NO₂ values at 20 km measured by SAOZ and MkIV were greater than 1.0 ppbv (Figures 3 and 5), whereas those measured by LPMA and MIPAS-B2 were smaller than 1.0 ppbv (Figures 1 and 4). On the other hand, for HALOE data, the δNO_2 value at 45 km, where the NO₂ concentration is often lower than 1–2 ppbv, is larger than at 25–40 km (Table 9). At 45 km, the standard deviation of the δNO_2 values is as large as 100% (Table 9). Therefore, the ILAS NO₂ measurements below 20 km and above 45 km were most likely degraded by low (<1.0 ppbv) NO₂ concentrations.

[37] The δNO_2 values for the POAM II data at 30–35 km are within \pm 10%, with progressively larger δNO_2 at lower altitudes (Table 9 and Figure 9b). This result is very similar to that obtained by Danilin *et al.* [2002], who have compared the ILAS NO₂ data with those measured by POAM II in SH in November 1996 using the trajectory hunting technique with large statistics. The δNO_2 value for POAM II data is 28% at 25 km, where the δNO_2 is 6% for HALOE. Using the ILAS NO₂ data as a reference, the difference of POAM II data from HALOE at 25 km was calculated to be -17%. On the other hand, Randall *et al.* [1998] have made the direct comparisons of NO₂ profiles measured by POAM II and HALOE. These comparisons indicate that POAM II data are 5–15% smaller than HALOE at 25 km, accounting for the difference in δNO_2 values between POAM II and HALOE at 25 km.

6. Summary

[38] The results of the comparisons of ILAS NO₂ data with those by balloon-borne (LPMA, SAOZ, MIPAS-B2, and MkIV) and satellite (POAM II and HALOE) measurements are summarized in Tables 9 and 10. The best agreement of the ILAS NO₂ with the balloon-borne measurements, within \sim 10%, was obtained at 25–30 km in NH in the winter and spring.

[39] For the comparisons with POAM II measurements in NH and SH in November and HALOE in NH and SH in spring and summer, the δNO_2 values, defined as (ILAS –

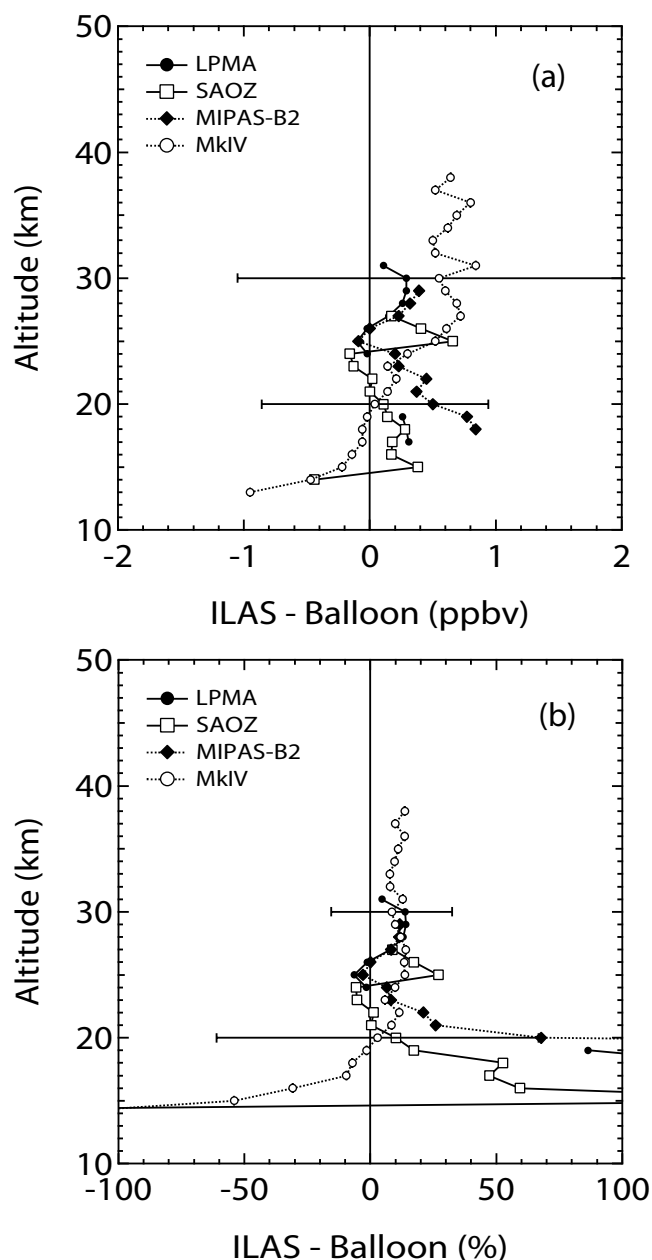


Figure 8. (a) The ΔNO_2 and (b) δNO_2 values for the balloon-borne NO_2 mixing ratios as a function of altitude. The values for LPMA, SAOZ, MIPAS-B2, and MkIV are shown by lines with solid circles, squares, diamonds, and open circles, respectively. Error bars indicate the combined uncertainties of the ILAS and balloon measurements, and are shown only for the comparisons with the MkIV profile at 20 and 30 km.

correlative measurement)/(correlative measurement), were calculated for each altitude. Since the differences in the δNO_2 values between inside and outside the vortex and between NH and SH were within the uncertainty of the ILAS measurements, the averages of all the δNO_2 values for each altitude were used for the comparisons with POAM II and HALOE data. At 25–40 km, the average δNO_2 values for HALOE data were within $\sim\pm 10\%$. For POAM II data,

the δNO_2 values were within $\sim\pm 10\%$ at 30–35 km, while the δNO_2 value at 25 km was 30%. The difference in the δNO_2 values for HALOE and POAM II data at 25 km is accounted for by the result of the NO_2 difference between HALOE and POAM II as described by *Randall et al.* [1998].

[40] The systematic difference between the ILAS and balloon NO_2 data was as large as 170% at 20 km, where the NO_2 concentrations were lower than 1.0 ppbv. The δNO_2 values for HALOE data increased to 27% and 40% at 20 and 45 km, respectively, where the NO_2 mixing ratios were significantly lower than those at 25–40 km.

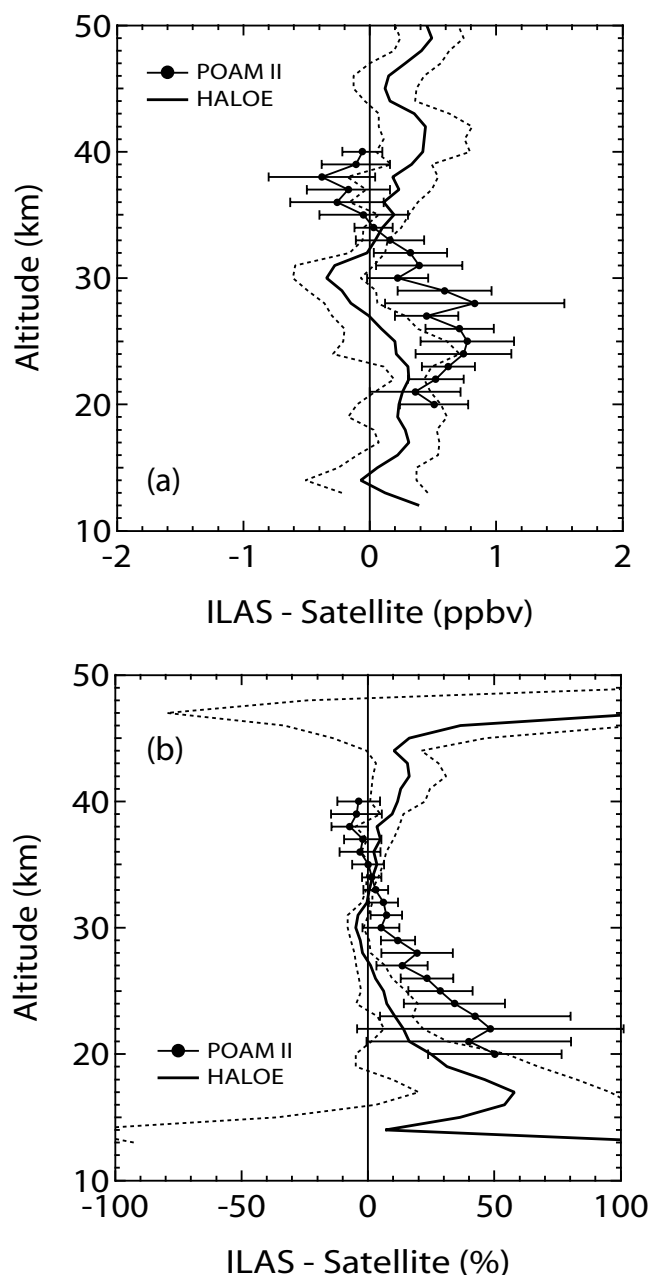


Figure 9. (a) ΔNO_2 and (b) δNO_2 values for POAM II and HALOE NO_2 mixing ratios as a function of altitude. Bars and dashed lines indicate 1- σ standard deviations of the values for POAM II and HALOE, respectively.

Table 9. The Average δNO_2 Values in Each 5-km Layer^a

Experiment	15 km	20 km	25 km	30 km	35 km	40 km	45 km
LPMA	...	155 ± 40	0 ± 6	11 ± 5
SAOZ	1 ± 214	16 ± 21	8 ± 14
MIPAS-B2	...	520 ± 742	4 ± 5	12 ± 0
MkIV	-99 ± 107	3 ± 7	11 ± 4	10 ± 2	10 ± 2
POAM II	28 ± 21	10 ± 9	0 ± 6
HALOE	45 ± 290	27 ± 28	6 ± 9	-3 ± 3	3 ± 4	10 ± 10	39 ± 94

^a Average difference ± standard deviation.

[41] Analyses of stratospheric chemistry using ILAS NO₂ require the careful consideration of the following effects: (1) negative biases in ILAS NO₂ values resulting from interference that was not completely eliminated even with the correction for optically thick PSCs, where aerosol extinction coefficients at 780 nm were greater than 0.001 km⁻¹, and (2) the lack of diurnal correction contributing to the positive bias in ILAS NO₂ below 25 km.

[42] ILAS version 5.20 HNO₃ data achieved excellent agreement (systematic differences <±0.1 ppbv) with balloon-borne HNO₃ at 20–35 km, as described in the appendix. This indicates small errors in the height registration at these altitudes in the version 5.20 products.

Appendix A. Validation of ILAS Version 5.20 HNO₃ Data

[43] The quality of version 5.20 ILAS HNO₃ data is evaluated here in the same way as has been done for the version 3.10 ILAS data by *Koike et al.* [2000]. The HNO₃ data used for the comparison with ILAS were obtained from balloon-borne measurements with a chemiluminescence detector, a cold atmospheric emission spectral radiometer, LPMA, MIPAS-B2, a far-infrared spectrometer, and MkIV. At 12 km, the NO_y measurement from the Deutsches Zentrum für Luft- und Raumfahrt Falcon research aircraft is also used. The ILAS HNO₃ data used were obtained nearest to the location where the balloon-borne measurements were made. The systematic differences in HNO₃ (ILAS minus balloon) were ±15%, -10%, and -19% at 15–25, 30, and 35 km, respectively, for the version 3.10 ILAS data. Random differences were 35%, 10%, and 14–34% at 15, 20–25, and 30–35 km, respectively. It has been suggested that the altitude change in the systematic difference of HNO₃ (Figure 10) was caused by an error in the height registration in the version 3.10 ILAS retrieval algorithm [*Koike et al.*, 2000], consistent with the analysis of the version 3.10 ILAS O₃ data [*Sasano et al.*, 1999b].

Table 10. Systematic and Random NO₂ Differences Between the ILAS and Balloon Measurements

	20 km	25 km	30 km
Systematic difference			
Percent	172	6	11
ppbv	0.26	0.25	0.44
Random difference			
Percent	439	9	3
ppbv	0.26	0.32	0.22

[44] Figure 10 shows the results of comparisons of the version 5.20 HNO₃ data with balloon-borne HNO₃ measurements. The ILAS version 5.20 HNO₃ data achieved excellent agreement (systematic differences <±0.1 ppbv) with the balloon-borne HNO₃ data in the 20–35 km range (Figure 10). The ILAS HNO₃ value at 15 km has a positive bias of ~0.5 ppbv. Systematic differences at 15, 20–25, 30, and 35 km were estimated to be 13%, 0–1%, 4%, and 25%, respectively. The random difference remained unchanged from the algorithm versions 3.10 through 5.20. Most of the altitude change in the systematic difference disappeared (Figure 10), indicating that the error in the height registration has become smaller, especially at 20–25 km.

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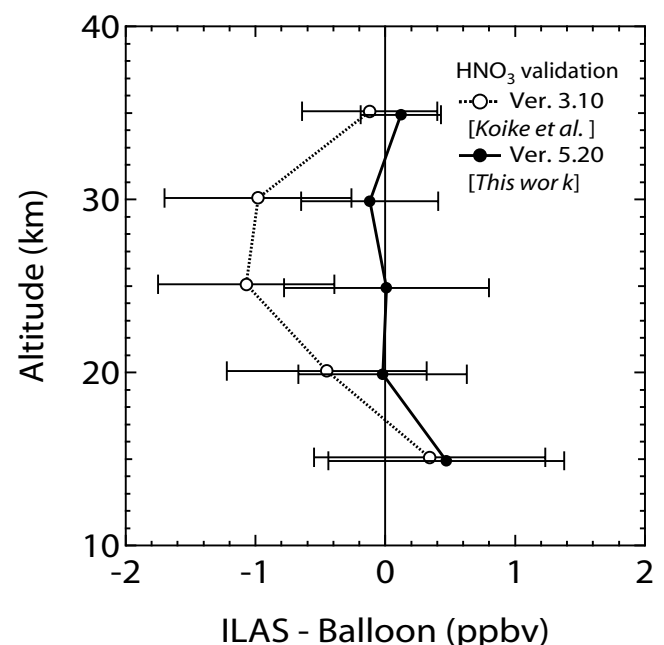


Figure 10. Systematic differences between the ILAS and balloon-borne HNO₃ mixing ratios for the version 3.10 and 5.20 ILAS data. Bars indicate random differences between ILAS and balloon-borne HNO₃ mixing ratios.

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