

Global HCFC-22 measurements with MIPAS: retrieval, validation, global distribution and its evolution over 2005–2012

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Abstract. We report on HCFC-22 data acquired by the Michelson Interferometer for Passive Atmospheric Sounding (MIPAS) in the reduced spectral resolution nominal observation mode. The data cover the period from January 2005 to April 2012 and the altitude range from the upper troposphere (above cloud top altitude) to about 50 km. The profile retrieval was performed by constrained nonlinear least squares fitting of modelled spectra to the measured limb spectral radiances. The spectral ν_4 -band at $816.5 \pm 13 \text{ cm}^{-1}$ was used for the retrieval. A Tikhonov-type smoothing constraint was applied to stabilise the retrieval. In the lower stratosphere, we find a global volume mixing ratio of HCFC-22 of about 185 pptv in January 2005. The rate of linear growth in the lower latitudes lower stratosphere was about 6 to 7 pptv/yr in the period 2005 – 2012. The profiles obtained were compared with ACE-FTS satellite data v3.5, as well as with MkIV balloon profiles and cryosampler balloon measurements. Between 13 km and 22 km, average agreement within -3 to +5 pptv (MIPAS - ACE) with ACE-FTS v3.5 profiles is demonstrated. Agreement with MkIV solar occultation balloon-borne measurements is within 10 – 20 pptv below 30 km and worse above, while in situ cryosampler balloon measurements are systematically lower over their full altitude range by 15–50 pptv below 24 km and less than 10 pptv above 28 km. MIPAS HCFC-22 time series below 10 km altitude are shown to agree mostly well to corresponding time series of near-surface abundances from NOAA/ESRL and AGAGE networks, although a more pronounced seasonal cycle is obvious in the satellite data. This is attributed to tropopause altitude fluctuations and subsidence of polar winter stratospheric air into the troposphere. A parametric model consisting of constant, linear, quasi-biennial oscillation (QBO) and several sine and cosine terms with different periods has been fitted to the temporal variation of stratospheric HCFC-22 for all 10° -latitude/1-to-2-km-altitude bins. The relative linear variation was always positive, with relative increases of 40 – 70 %/decade in the tropics and global lower stratosphere, and up to 120 %/decade in the upper stratosphere of the northern

polar region and the southern extratropical hemisphere. In the middle stratosphere, between 20 and 30 km, the observed trend is inconsistent with the trend at the surface (corrected for the age of stratospheric air). There exists a stronger positive trend in HCFC-22 in the southern hemisphere and a more muted positive trend in the northern hemisphere, implying a potential change in the stratospheric circulation over the observation period.

1 Introduction

HCFC-22 (CHClF_2) is a chlorine source gas and a greenhouse gas (IPCC, 2014). The sources of HCFC-22 are anthropogenic emissions due to its use as a propellant and refrigerant. The gas is removed from the atmosphere by photolysis and by reactions with $\text{O}(^1\text{D})$, Cl^- , and the OH radical. The chemical lifetime of HCFC-22 in the stratosphere is 165 years (12 years for its global total atmospheric lifetime), according to SPARC(2013), but has been estimated to a significantly longer span of 260 ± 25 years by Moore and Remedios (2008). The radiative forcing potential of HCFC-22 is $0.208 \text{ Wm}^{-2} \text{ ppbv}^{-1}$, and its ozone depletion potential is about 20 times lower than that of CFC-12 (0.04) (World Meteorological Organization (WMO), 2014). Production and import of HCFC-22 is limited by the Montreal Protocol on Substances that Deplete the Ozone Layer and will be banned by 2030 for dispersive uses in developed countries (United Nations Environment Programme, 2009). The 2007 Adjustment to the Protocol asks for a 100% reduction by 2030 for all countries, including developing nations – albeit with a 2.5% allowance for servicing of refrigeration and air conditioning equipment existing on 1 January 2030 for the period 2030–2040 and subject to review in 2015. Therefore the overall reduction shall already be 97.5 - 100% by 2030.

Ambient HCFC-22 was first measured by Rasmussen et al. (1980) by air sampling techniques. Atmospheric HCFC-22 abundances are typically measured on site by gas-chromatographic techniques or by collecting samples in flasks or by balloon-borne air-sampling measurements (Engel et al., 1997) followed by subsequent gas-chromatographic analysis in a central laboratory (Montzka et al., 2009; O'Doherty et al., 2004; Yokouchi et al., 2006). Further, there exist remote measurements by infrared spectroscopy from ground-based (Rinsland et al., 2005b; Zander et al., 2005; Gardiner et al., 2008), balloon-borne (Murray et al., 1975; Williams et al., 1976; Goldman et al., 1981), or space-borne (Zander et al., 1987; Rinsland et al., 2005a; Moore and Remedios, 2008) platforms in solar absorption geometry. Among recently flying space-borne instruments, only the Atmospheric Chemistry Experiment – Fourier Transform Spectrometer (ACE-FTS; solar occultation) and the Michelson Interferometer for Passive Atmospheric Sounding (MIPAS; limb emission) have been pro-

viding measurements of HCFC-22 (Kolonjari et al., 2012; Park et al., 2014; Moore and Remedios, 2008). The history of measurements is summarized in von Clarmann (2013).

In this paper we present and discuss HCFC-22 distributions and time series as retrieved with the MIPAS data processor developed and operated by the Institute for Meteorology and Climate Research at the Karlsruhe Institute of Technology (KIT-IMK) in Germany in cooperation with the Instituto de Astrofísica de Andalucía (IAA, CSIC) in Granada, Spain. In the next section we provide a description of the MIPAS instrument and measurements. The retrieval strategy and error estimation are summarized in Section 3. Section 4 reports on the validation of this data set, and Section 5 presents global distributions of HCFC-22 and an assessment of temporal variations including a linear trend. The derived estimates of trends will be compared to those from long-term surface data records. Section 6 contains the discussion of the results and the summary.

2 MIPAS data

MIPAS measured the thermal emission of the atmosphere, and thus provided data during day and night. It was a cryogenic limb emission Fourier transform spectrometer (FTS) designed for measurement of trace species from space (European Space Agency, 2000; Endemann and Fischer, 1993; Endemann et al., 1996; Fischer and Oelhaf, 1996). MIPAS was one of 10 instruments aboard the Environmental Satellite (Envisat). Envisat was launched into a sun-synchronous polar orbit on 1 March 2002 at approximately 800 km, with an orbital period of about 101 min (Fischer et al., 2008), resulting in more than 14 orbits per day. The end of the Envisat mission was declared on 9 May 2012 after loss of communication with the satellite on 8 April 2012.

MIPAS sounded the atmosphere tangentially to the Earth in the infrared spectral range (4.15–14.6 μm) covering tangent altitudes from about 7 to 72 km in its nominal observation mode. The instrument's field of view was approximately 3 km (vertically) \times 30 km (horizontally). MIPAS operated from July 2002 to March 2004 with full spectral resolution as specified: 0.05 cm^{-1} in terms of full width at half maximum, after apodisation with the 'strong' function suggested by Norton and Beer (1976). The full resolution mode was stopped in March 2004. Starting from January 2005 and up to the end of the mission, the spectral resolution of MIPAS was degraded from 0.05 cm^{-1} to 0.12 cm^{-1} (apodised). These later measurements are referred to as 'reduced resolution mode'.

The data analysis reported in this paper relies on the ESA-provided so-called level-1b data product which includes calibrated phase-corrected and geolocated radiance spectra (Nett et al., 1999). The versions of ESA level-1b data used are IPF 5.02–5.06. All spectra under consideration here were recorded according to the nominal reduced resolution mea-

surement mode, including per limb sequence 27 tangent altitudes between about 7 and 72 km, with the tangent altitude adjustment following roughly the tropopause altitude over latitudes. The vertical distance between adjacent tangent heights varies between 1.5 km in the upper troposphere/lower stratosphere up to 4.5 km in the mesosphere. One limb scan is recorded per each 410 km increment along the polar sun-synchronous orbit, leading to a dense horizontal sampling which is independent of the season or latitude band. As a result, MIPAS covers all latitudes during one orbit.

3 Retrieval

The retrieval of HCFC-22 profiles presented here was performed with a MIPAS data processor dedicated for research applications, which has been developed at the Institut für Meteorologie und Klimaforschung (IMK) and complemented by components relevant to treatment of non-local thermodynamic equilibrium at the Instituto de Astrofísica de Andalucía (IAA). The non-LTE components are not used in this study because they are not relevant to the retrieval of HCFC-22.

The IMK retrieval processor consists of the radiative transfer algorithm KOPRA (Stiller, 2000) and the retrieval algorithm RCP (Retrieval Control Program). Local spherical homogeneity of the atmosphere is assumed here, i.e., atmospheric state parameters related to one limb sounding sequence are assumed not to vary with latitude or longitude but only with altitude. An exception is temperature for which horizontal gradients are considered in the retrieval (Kiefer et al., 2010). The general strategy of the IMK/IAA data processing has been documented in von Clarmann et al. (2003).

3.1 Retrieval of HCFC-22

HCFC-22 is retrieved by a constrained multi-parameter non-linear least-squares fitting of modelled to measured spectra. Spectral data from all tangent altitudes are analysed within one inversion process, as suggested by Carlotti (1988). Volume mixing ratio (vmr) vertical profiles are retrieved on a fixed, i.e. tangent height independent altitude grid which is finer than the tangent height spacing (1-km steps from 4 to 35 km; then 5-km grid width from 35 to 50 km; 10 km grid width from 50 to 100 km; 120 km). In order to obtain stable profiles, the profiles have been constrained such that the first order finite difference quotient $\Delta \text{vmr} / \Delta \text{altitude}$ at adjacent altitude grid points was minimised, similar as proposed by Tikhonov (1963). An altitude-constant profile of zero mixing ratio throughout was taken as a priori, while the initial guess profile was from the MIPAS climatology (Kiefer et al., 2002). For the retrieval of HCFC-22 we have used 4 microwindows of the MIPAS spectrum (803.500 cm^{-1} to 804.750 cm^{-1} , 808.250 cm^{-1} to 809.750 cm^{-1} , 820.500 cm^{-1} to 821.125 cm^{-1} and 828.750 cm^{-1}

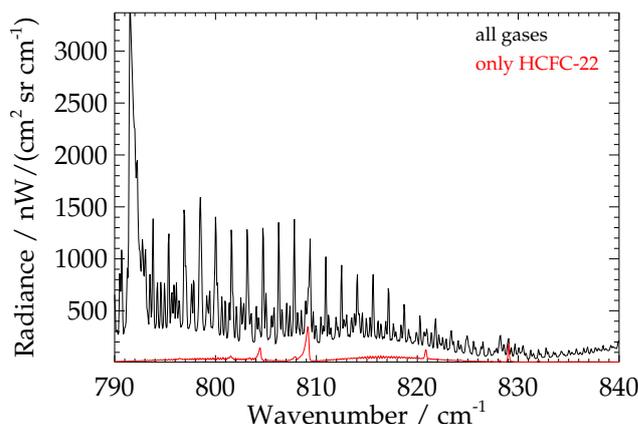


Figure 1. Typical spectrum of HCFC-22 (red curve) and combined spectrum of 24 gases (HCFC-22 included) (black curve) at 16 km tangent altitude.

to 829.500 cm^{-1}). For HCFC-22, the HITRAN2K spectroscopic database (Rothman et al., 2003) was used, with some updates for interfering species.

Fig. 1 shows the atmospheric limb emission radiance spectrum at 16 km tangent altitude including contributions from H_2O , CO_2 , O_3 , NO_2 , NH_3 , HNO_3 , ClO , OCS , HCN , CH_3Cl , C_2H_2 , C_2H_6 , COF_2 , C_2H_4 , HNO_4 , CFC-11 , CCl_4 , CFC-113 , ClONO_2 , CH_3CCl_3 , CH_3OH , $\text{C}_2\text{H}_3\text{NO}_5$ (peroxyacetyl nitrate, PAN), $\text{C}_3\text{H}_6\text{O}$ and HCFC-22 (black curve). In addition, the sole contribution of HCFC-22 is shown in red.

For some quantities, information from preceding retrieval steps was used for the retrieval of HCFC-22: a correction of the spectral shift caused by a less-than-perfect frequency calibration; the tangent altitudes of the limb measurements; temperature; and finally the mixing ratios of the species O_3 , H_2O , HNO_3 , ClO , ClONO_2 , CFC-11 , HNO_4 as well as C_2H_6 . The vmr of CO_2 and all other remaining gases were taken from a climatological database, and the temperature was retrieved from CO_2 lines (von Clarmann et al., 2003, 2009b).

Simultaneously with HCFC-22 and with adequate regularisation, we jointly fitted the following quantities: the mixing ratio of PAN; a wavenumber-independent continuum absorption coefficient per microwindow and altitude accounting for aerosol emission; and an additive radiative offset for each microwindow. The data versions are V5r_F22_220 and V5r_F22_221. The only difference between these versions is the source of temperature analysis data used as a priori for the preceding temperature retrieval. Results are equivalent and different version numbers are used only to guarantee full traceability of the retrievals.

3.2 Diagnostics

Diagnostic quantities to characterise the HCFC-22 measurements include estimates of measurement noise, of retrieval errors caused by uncertainties in ancillary parameters used in the radiative transfer modelling, and the averaging kernel (AK) matrix (Rodgers, 2000).

Table 1 shows the estimated total retrieval error of a HCFC-22 profile measured at 18.6°S and 111.6°W on 9 January 2009. This error estimate is considered to be roughly representative for the entire data set but is rather conservative, due to the typically low lower stratospheric temperatures at tropical latitudes, which are associated with a lower signal. The total error is the square root of the quadratic sum of noise error and parameter errors. Its most important components are the uncertainty of the elevation pointing of the line of sight (LOS), the uncertainty of pre-retrieved O₃ mixing ratios, the gain calibration uncertainty, the residual spectral shift uncertainty, and the instrument line shape (ILS) uncertainty. The spectroscopic error (not reported in the table) is about 5%. In the given altitude range, errors resulting from uncertainties of interfering species contribute to the total error by less than 1%.

The percentage of non-converged profiles is about 0.01 to 0.02%. The strength of the regularisation, i.e. the weight of the constraint, has been chosen to be altitude-dependent with a scheme proposed by Steck (2002) such that the retrieved profile represents approximately 5 degrees of freedom, corresponding to a typical altitude resolution of 3 km at 10 km height and 7 km at 30 km height, and further increasing with height (Fig. 2). Here the vertical resolution is provided in terms of full width at half maximum of a row of the averaging kernel matrix. The rows of the AK-matrix show how much information from other atmospheric altitudes contribute to the vmr on the given retrieval altitude. An example of the rows of the HCFC-22 averaging kernel matrix is shown in Fig. 3. The largest peaks of the averaging kernels are generally found in the upper troposphere. This is because the retrieval is more strongly regularised at higher altitudes. In general, the retrieval is well-behaved in a sense that the averaging kernels peak at their nominal altitudes (marked by diamonds in Fig. 3) between 12 and 50 km. The integral over the averaging kernels in this altitude range are, at good accuracy, unity, thus our choice of the constraint does not impose any bias.

The information evaluated by a limb retrieval is not located in one single point but spread horizontally along the line of sight direction. The horizontal information smearing of the HCFC-22 measurement is estimated by using the method of von Clarmann et al. (2009a). In terms of full width at half maximum of the rows of the horizontal averaging kernel matrix, the horizontal information smearing of a MIPAS retrieval typically varies between about 210 and 680 km for most species, altitudes and atmospheric conditions. For HCFC-22 the horizontal information smearing, calculated as

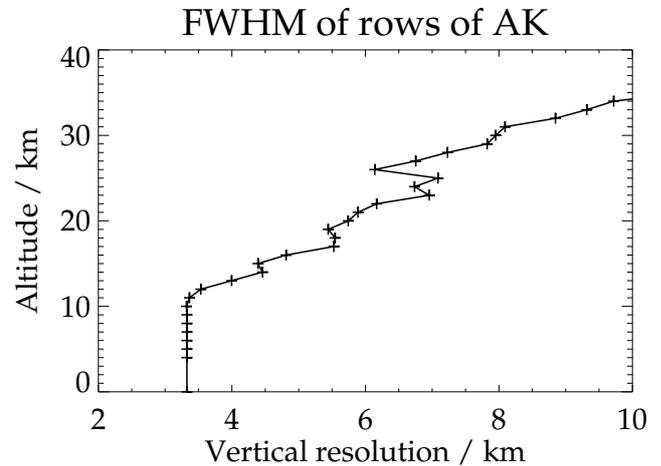


Figure 2. Altitude resolution in terms of full width at half maximum (FWHM) of a row of the averaging kernel matrix.

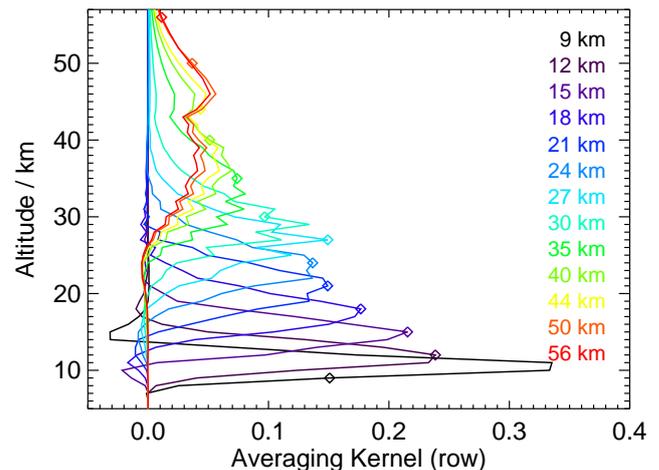


Figure 3. Rows of averaging kernel of HCFC-22 measurements for reduced spectral resolution nominal mode. The diamonds represent the nominal altitudes (e.g. the diagonal value of the averaging kernel matrix). For clarity, only every third kernel is shown.

Table 1. Error budget of a V5r HCFC-22 (nominal mode, reduced resolution) retrieval on 9 January 2009, 18.6°S latitude and 111.6°W longitude degree, orbit 35874, for selected altitudes. The errors are given in units of mixing ratios (pptv), and additionally, in parentheses, in percentage units (%).

V5r HCFC-22 Height	Total Error	Noise	Parameter Error	O ₃	LOS	Shift	Gain	ILS
40 km	14(17.3)	13(16.1)	5.7(7.0)	0.3(0.3)	1.7(2.1)	1.9(2.3)	0.3(0.3)	2.3(2.8)
35 km	12(11.2)	11(10.2)	6.1(5.7)	0.2(0.1)	1.4(1.3)	2.3(2.1)	0.5(0.5)	1.5(1.4)
30 km	12(9.0)	9.4(7.1)	7.3(5.5)	<0.1(<0.1)	1.1(0.8)	2.3(1.7)	1.0(0.7)	0.4(0.3)
25 km	13(8.0)	9.0(5.6)	9.1(5.6)	0.2(0.1)	3.2(2.0)	1.2(0.7)	0.6(0.3)	0.8(0.5)
20 km	14(7.0)	8.2(4.1)	12(6.0)	0.3(0.2)	5.2(2.6)	0.5(0.2)	0.5(0.2)	2.4(1.2)
15 km	13(5.6)	7.1(3.1)	11(4.7)	0.3(0.1)	1.2(0.5)	0.8(0.3)	1.6(0.7)	2.8(1.2)
10 km	13(6.8)	8.6(4.5)	9.6(5.0)	<0.1(<0.1)	1.8(0.9)	0.1(<0.1)	2.1(1.1)	2.0(1.0)

Table 2. Horizontal averaging kernels (full width at half maximum) calculated according to von Clarman et al. (2009a) for retrieval altitudes between 40 km and 10 km for HCFC-22. Positive sign means displacement towards the satellite.

Height	FWHM	Displacement
40 km	554 km	-121 km
35 km	549 km	5 km
30 km	528 km	62 km
25 km	521 km	55 km
20 km	608 km	90 km
15 km	300 km	133 km
10 km	327 km	126 km

the half width of the horizontal component of the 2D averaging kernel, is approximately 300 km at altitudes below 15 km, 608 km at altitude 20 km and approximately 550 km above (see Tab. 2).

The information displacement is defined as the horizontal distance between the point where the most information comes from and the nominal geolocation of the limb scan, which is defined as the geolocation of the tangent point of the middle line of sight in a MIPAS limb scan. The information displacement in case of HCFC-22 varies between -121 km at 40 km altitude (the negative sign refers to displacement beyond the tangent point with respect to the satellite) and 133 km at 15 km altitude, and is lowest and positive (i.e. displacement towards the satellite) in the middle stratosphere (see Tab. 2).

4 Validation

Validation of the HCFC-22 MIPAS IMK profiles is performed by comparison to coincident independent measurements. The availability of reference measurements for the validation of MIPAS HCFC-22 in the stratosphere is quite limited: the only space-borne instrument measuring the ver-

tical profiles of HCFC-22 at the same time as MIPAS is ACE-FTS. We also perform the comparison with MkIV balloon profiles and with measurements at different heights performed with the balloon-borne cryosampler flown by the University of Frankfurt. Although there exist aircraft measurements of HCFC-22 (e.g. Xiang et al., 2014), these have not been used, because of unsolved problems caused by the different altitude resolutions of MIPAS and the air sampling measurements. The application of averaging kernels to homogenise profiles of different altitude resolution as suggested by Connor et al. (1994) is only possible if the better resolving instrument provides vertical profiles but not if values at a single altitude level are available only.

4.1 Comparison with ACE-FTS

The Atmospheric Chemistry Experiment Fourier Transform Spectrometer (ACE-FTS) is a solar occultation instrument flying on the SCISAT satellite platform since August 2003 (Bernath et al., 2005). It takes measurements from the upper troposphere to about 150 km altitude. Temperature, pressure, atmospheric aerosol extinction, and the concentrations of a large number of atmospheric species are retrieved from these measurements with a vertical resolution in the order of 4 km. The SCISAT flies on a highly inclined circular orbit (650 km altitude), which implies that more than half of the ACE-FTS measurements occur at high (over 60°N and S) latitudes. It takes approximately three months for all latitudes (~82°N–82°S) to be sampled by ACE-FTS.

HCFC-22 is retrieved by the algorithm described in Boone et al. (2005) and Boone et al. (2013) from a single window of width 25 cm⁻¹ centered at 817.5 cm⁻¹. The ACE-FTS retrieved HCFC-22 profiles extend from the upper troposphere to about 30 km height, do not include any formal a priori information, and the reported errors are in the order of 3 to 5%, going up to 10% at the lowest and 8% at the highest altitude limits of the retrieval. Spectroscopic data from HITRAN 2004 (Rothman et al., 2005) were used for HCFC-22 but for

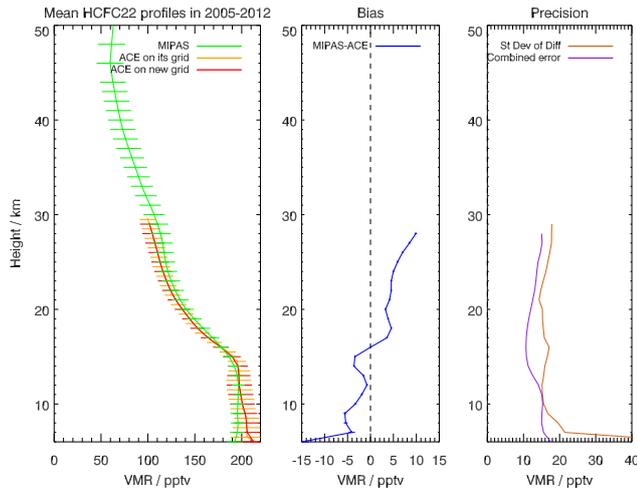


Figure 4. Mean profiles, bias, and standard deviation of the differences versus estimated combined retrieval error for ACE-FTS and MIPAS retrievals of HCFC-22. The original altitude grid on which the ACE-FTS data were provided uses the same grid spacing as MIPAS of 1-km, but is shifted by 0.5 km. Thus, the data were re-sampled on the MIPAS grid. The error bars in the left panel are the typical errors of a single profile, estimated as the mean error over the sample. The tiny error bars in the middle panel (hardly discernible) are the standard errors of the mean differences.

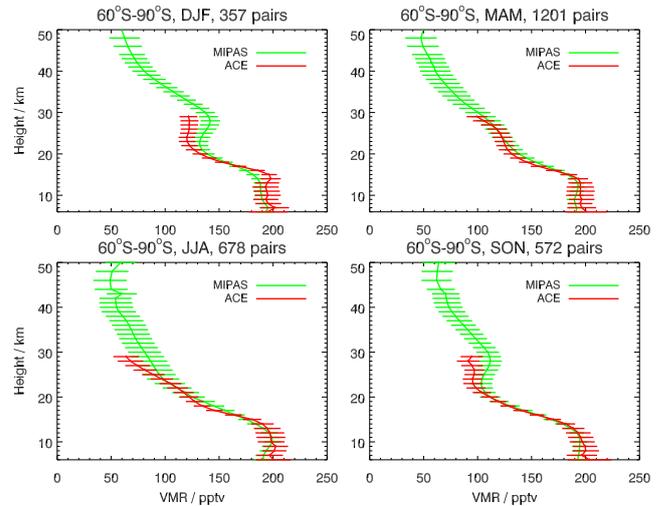


Figure 5. Seasonal mean vmr values (in pptv) of ACE-FTS (red) and MIPAS (green) in 2005-2012 at southern polar latitudes. Error bars represent the typical precision of a single profile, estimated as the mean error over the sample.

this species no updates have been made with respect to the
 HITRAN2K version used for MIPAS.

The analysis was performed on January 2005 – April 2012
 data of ACE-FTS version 3.5, with collocation criteria of 500
 km and 5 hours; this leads to a comparison subset of 8393 co-
 located measurements. In the case of multiple matches, only
 the closest MIPAS profile was used.

The comparison of global mean MIPAS and ACE-FTS
 HCFC-22 profiles (left panel of Fig. 4) reveals a high MIPAS
 bias of 5 to 10 pptv at 17–29 km altitudes and a low MIPAS
 bias (between 0 and -3 pptv) at 10–13 km. Between 13 and
 22 km altitude, the mean difference is -3 to +5 pptv. The bias
 is significant at all altitude levels. Analysis of the scatter of
 the differences versus the estimated combined precision of
 the instruments (Fig. 4, right panel) indicates an underesti-
 mation of the combined uncertainty, i.e. one or both instru-
 ments underestimate the random component of their errors
 (c.f. von Clarmann (2006)). It should, however, be kept in
 mind that the reported fitting error estimates of ACE-FTS in-
 clude only measurement noise and do not include randomly
 varying parameter errors, which implies that perfect coinci-
 dence of the two curves on the right panel of Fig. 4 cannot be
 expected. Further, atmospheric variability within the radius
 defined by the coincidence criteria can contribute to these dif-
 ferences. This is particularly true because many of the ACE-
 FTS measurements occur at higher northern latitudes where
 atmospheric variability is quite pronounced.

The seasonality of the differences between ACE-FTS and
 MIPAS for southern polar latitudes is analysed in Fig. 5.
 Most pronounced differences occur at the top end of the
 ACE-FTS profiles, with ACE-FTS always lower than MI-
 PAS, while the seasonality in the differences comes mainly
 from a very steep vertical gradient in the ACE-FTS profiles
 for polar summer between 16 and 20 km altitude (top left
 panel) which is not in the same way reproduced by MIPAS
 profiles. The very steep vertical gradient in this particular
 case leads to the highest bias of ACE-FTS vs. MIPAS of
 about 15 pptv around 16 km and a low bias around 20 km
 and above.

The correlation plots of ACE-FTS versus MIPAS (Fig. 6)
 corroborate the findings so far: the ACE-FTS and MIPAS
 data points fall into two clusters that match around 16 km
 altitude. Below this altitude (red and yellow colours) the re-
 gression line is slightly steeper than unity indicating a bias
 proportional to the absolute values, while above 16 km ACE-
 FTS has a small and almost constant low bias versus MIPAS.

The histogram plots (Fig. 7) represent the distribution of
 measured volume mixing ratios over latitude and time for a
 fixed altitude level. The histograms for ACE-FTS and MI-
 PAS at 23 km are very similar in shape and position of the
 peak value, while at 16 km the ACE-FTS distribution of
 measured vmrs is somewhat wider. The latter is attributed
 to the fact that the ACE-FTS retrieval does not use any re-
 gularisation, which leads to greater scatter. Nevertheless, both
 peak and extreme values match quite well and the histograms
 suggest a robust agreement between the two instruments, al-
 beit with an altitude-dependent systematic offset as discussed
 above. The bimodal distribution at 23 km altitude is caused

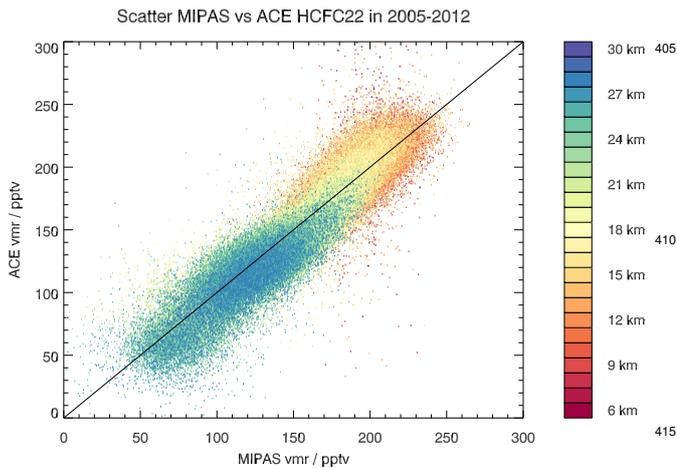


Figure 6. Scatter plot of ACE-FTS versus MIPAS HCFC-22 data points.

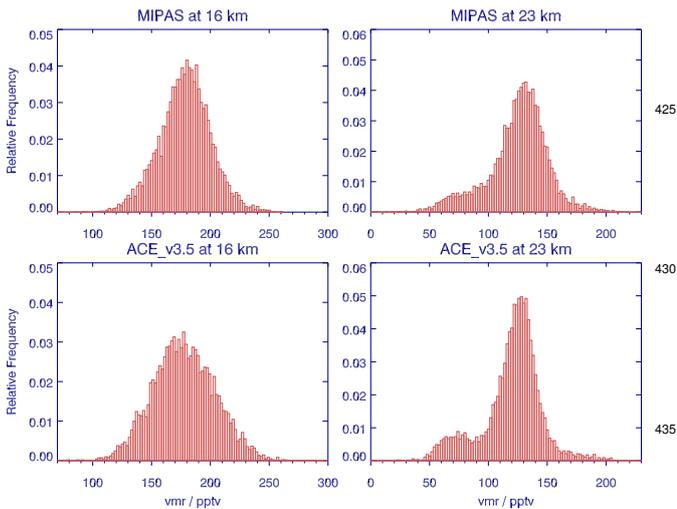


Figure 7. Histograms of MIPAS (upper panels) versus ACE-FTS (lower panels) HCFC-22 mixing ratios at 16 km (left panels) and 23 km (right panels) altitude.

by differences in the air masses sounded by each instrument, where the polar air masses correspond to the lower mode and the mid-latitude air masses to the higher mode. The number of tropical collocations is small and has thus minor impact on the histogram. The ACE-FTS modes are more clearly separated than the MIPAS modes because of the differences in the North-South component of the lines of sight of each measurement technique. Since the mixing ratio gradients in North-South direction are typically larger than those in East-West direction, the modes of MIPAS that observes roughly in the orbit plane are more smeared than those of ACE-FTS whose line of sight is directed towards the sun.

In summary, although minor differences between MIPAS and ACE-FTS HCFC-22 measurements have been detected, the comparison justifies confidence in the data sets. Besides identified small biases in the order of <10 pptv the two data sets compare very well in absolute values, latitude distributions, and seasonalities.

4.2 Comparison with cryosampler profiles

The cryogenic whole air sampler, deployed on stratospheric balloons, is operated by the University of Frankfurt. The instrument collects high volume whole air samples which are frozen out by means of liquid neon. After the flight, the air is left to evaporate which provides high pressure whole air samples from different altitudes (Engel et al., 1997). A wide range of halocarbons are then analysed in these samples using a gas chromatograph coupled to a mass-spectrometer (Laube et al., 2008). The precision of the individual data points of the cryosampler measurements is typically in the order of 0.5%. The measurements were referenced to a standard provided by the National Oceanic and Atmospheric Administration (NOAA) (e.g. Montzka et al., 2003) and the data were reported on the NOAA-2006 calibration scale, which shows excellent agreement (within 1%) to most other calibration scales (Hall et al., 2014; World Meteorological Organization (WMO), 2014).

Cryosampler measurements do not represent contiguous vertical profiles but rather individual independent point measurements. Hence, no regridding has been applied in this case: the cryosampler measurements were just reported as they were on the height where they had been taken.

Coincidences between MIPAS and cryosampler measurements were considered within a spatial distance of 1000 km and a time window around the cryosampler measurement time of ± 24 hours. These coincidence criteria may seem large. However, in their work on the validation of CFC-11/CFC-12 and CH_4 measurements from MIPAS, where data from the same MIPAS geolocations and balloon flights were used, Eckert et al. (2015) and Laeng et al. (2015) found remarkably good agreement between MIPAS and the cryosampler data, except for biases confirmed by other validation instruments. Thus, it is rather unlikely that any differences found in the HCFC-22 comparison can be attributed to the relatively large spatial or temporal distance.

For the first two flights (first two panels of Fig. 8) that took place in June 2005 in the tropics the agreement between MIPAS profiles and cryosampler measurements above 30 km altitude is better than 10 pptv. Nevertheless, the MIPAS profile with the closest coincidence differs significantly from the other coincident profiles in both cases, hinting towards inhomogeneous situations in the atmosphere. Below 30 km, for the first flight neither the closest coincident profile, nor the mean of coincidences agrees with the cryosampler measurements. MIPAS data are higher than the cryosampler measurements between 15 pptv (mean coincident profile) and 35 pptv

(closest profile). For the second flight, the cryosampler data points fall mostly within the error bars of the closest MIPAS profile.

460 The third flight (third panel of Fig. 8) of the cryosampler instrument provided only four measurements, none of which being situated between 18 and 32 km. For all four data points, MIPAS is higher by 15–30 pptv. The cryosampler data of the fourth flight (fourth panel) show a narrow layer with very low HCFC-22 abundances around 23 km; within this layer the abundances are mostly far lower than the closest MIPAS profile. Pronounced oscillations appear also in some but not all individual MIPAS profiles, but have, where present, a significant high bias (20 to 50 pptv) compared to the cryosampler data. Other individual MIPAS profiles, in particular the profile of the closest coincidence, are rather smooth. The oscillating cryosampler data are attributed to an unusual atmospheric situation on this particular flight with a narrow lamina of HCFC-22 poor polar vortex air, and this situation led to strong small-scale variability. This hypothesis is corroborated by the large spread of the co-located MIPAS profiles. The lamina itself is either not encountered by the MIPAS measurements, or it is not resolved; MIPAS measurements represent for each profile point an air parcel of about 400 km in length times 30 km in width times 4 km in height. Below 18 km, the MIPAS profiles have a high bias of 15 to 20 pptv.

470 The last flight (bottom panel of Fig. 8) stands out by a pronounced HCFC-22 minimum (also present in the profiles of other tracers measured during this flight) in MIPAS data at approximately 28 km. Below this altitude, the cryosampler measurements coincide nicely with the MIPAS data but the altitude coverage of the cryosampler data set on this day does not allow to confirm the positive mixing ratio gradient above.

480 In summary, comparison to cryosampler data from five different profiles from tropical and northern polar winter atmosphere reveal a high bias of MIPAS HCFC-22 data of 15 to 50 pptv below 24 km; while above 28 km, the high bias is reduced to less than 10 pptv. The analysis of the comparisons is partly complicated by very inhomogeneous atmospheric situations and the enormous difference between sampling volumes of cryosamplers versus satellite remote sensing instruments.

4.3 Comparison with MkIV balloon interferometer profiles

500 The MkIV interferometer from Jet Propulsion Laboratory (JPL) is a high-resolution solar absorption spectrometer for measurement of over 30 atmospheric constituents which is deployed on stratospheric balloon platforms with a typical float altitude of 37 km (Toon, 1991). MkIV measured HCFC-22 using two spectral windows, centered at the ν_4 Q-branch at 809.19 cm^{-1} and the $2\nu_6$ Q-branch at 829.14 cm^{-1} . The widths of the microwindows were 1.28 cm^{-1} and 0.72 cm^{-1} , respectively. Pseudolines derived from spectroscopic measurements (McDaniel et al., 1991; Varanasi,

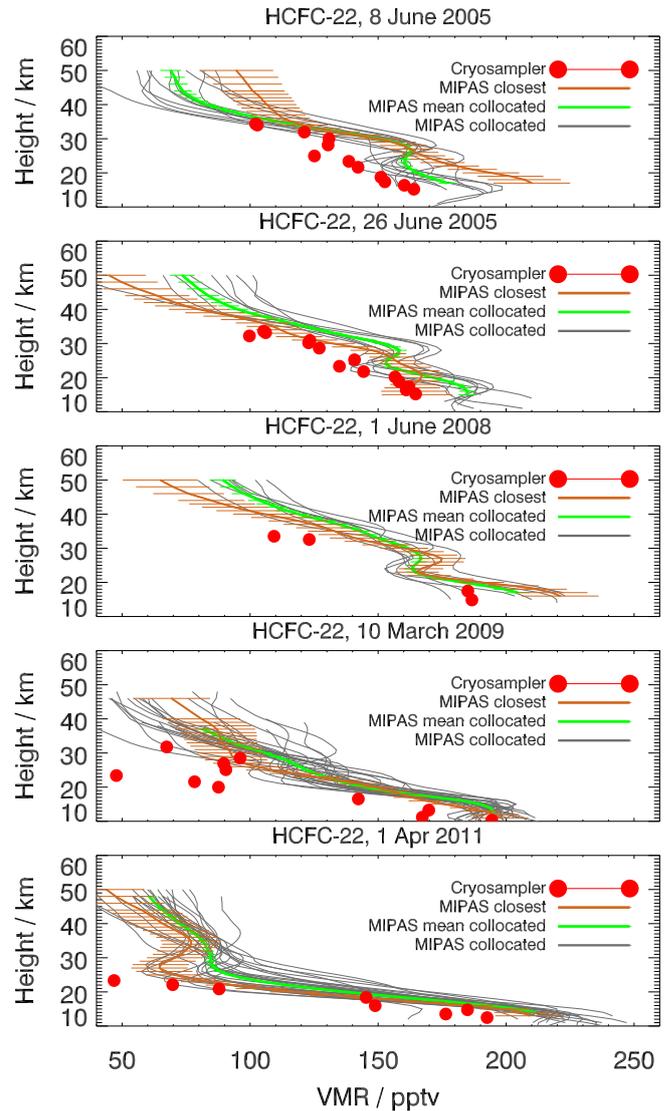


Figure 8. Five cryosampler profiles (red filled circles) and MIPAS HCFC-22 vmr profiles - all co-located (grey), closest collocations (orange), and the mean profiles from all collocations (green).

1992; Varanasi et al., 1994) have been used as spectroscopic data (see <http://mark4sun.jpl.nasa.gov/pseudo.html>). The instrument obtains vmr vertical profiles of HCFC-22 between cloud top and balloon altitude. MkIV measured two HCFC-22 vmr vertical profiles co-located by MIPAS reduced resolution measurements. The dataset is provided on a 1-km altitude grid between 10 and 40 km. The vertical resolution of the MkIV balloon profiles varies between 2–4 km. Data used here were measured during balloon flights from Fort Sumner, NM, with tangent altitude geolocations from 34.0°N–35.7°N and 108.8°W–114.1°W.

Figure 9 presents the two MkIV balloon profiles within the MIPAS reduced resolution period. The first MkIV profile, from September 22, 2007, was measured when MIPAS was temporarily inactive and no matches were found within 24 hours and 1000 km. The MkIV profile from 22 September 2007 was compared to the MIPAS 30°N–40°N monthly mean of September 2007. These observations were made at the end of a summer of easterly stratospheric winds. Since the flow is zonal during the summer and wave activity is negligible, little zonal variation in composition is to be expected, which justifies comparison with a zonal monthly mean of the same year. For the profile from the sunrise of September 23, 2007, three co-located MIPAS profiles were found (grey lines).

Below 25 km the MIPAS profiles agree with the MkIV profiles within the error bars, with MIPAS HCFC-22 being in tendency lower for the co-located profiles, (right panel) but in good agreement for the September 2007 monthly mean. Above 25 km the vertical gradients of mixing ratios from MkIV and MIPAS diverge, with MkIV profiles decreasing stronger with altitude than those of MIPAS. Up to 32 km, the agreement is still well within the error bars. For the 22 September 2007 comparison the profiles agree even up to the highest MkIV altitude levels.

4.4 Summary of the intercomparisons

The comparisons to the three available reference data sets, namely ACE-FTS, balloon-borne cryosampler data, and MkIV balloon measurements, do not provide a unique picture on MIPAS biases. Below about 20 km, MIPAS has either a low (ACE-FTS), a high (cryosampler) or no (MkIV) bias. Above 25 km the MIPAS bias is either clearly positive (ACE-FTS and MkIV) or small, i.e. less than +10 pptv (cryosampler). Between 20 and 25 km the bias can range from -30 pptv (MkIV) to +50 pptv (cryosampler). In summary we state that there is no clear indication of a bias, and MIPAS does not stand out as particularly high or low. MIPAS HCFC-22 data are found to be within ± 15 pptv (1σ) of reference data sets between 10 and 35 km altitude. The different size of the reference data sets needs to be taken into account. The large number of co-locations with ACE-FTS leads to a high statistical significance of deviations with respect to noise. Errors of systematic nature, however, do not cancel out by averaging,

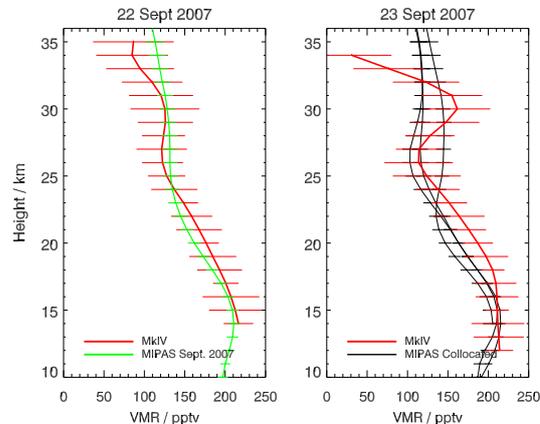


Figure 9. MkIV and MIPAS co-located and monthly mean vertical HCFC-22 profiles along with error bars. The mean profile is taken in the 30°N–40°N latitude band where the two balloon flights took place.

ing, and with respect to these, the balloon measurements are considered equally useful despite the small number of co-locations. In addition, cryosampler data do not rely on spectroscopic measurements as the other three data sets and, thus, are the most independent reference data within the intercomparisons. Disagreement with cryosampler data hints towards a potential bias of all spectral measurements due to incorrect spectroscopic information.

5 Global distributions and temporal evolution

5.1 Zonal means

Figures 10 and 11 show monthly zonal means of HCFC-22 vmr for December 2005 and 2010, and July 2006 and 2010, respectively, for all latitudes and altitudes up to 50 km. The months have been selected to be approximately in the same phase of the quasi-biennial oscillation (QBO). The typical distribution of a tropospheric source gas with photolytic sinks in the stratosphere is observed, with higher values in the tropics and lower altitudes, and lower values in the upper stratosphere and higher latitudes. Within the troposphere, MIPAS sees larger HCFC-22 abundances in the northern hemisphere (NH) than in the southern hemisphere (SH), owing to the global distribution of emissions, which is in agreement with previous results from Xiang et al. (2014), while in the stratosphere this kind of asymmetry is not observed. Furthermore, a substantial increase of HCFC-22 from the year 2005/2006 to the year 2010 can be derived from the MIPAS results.

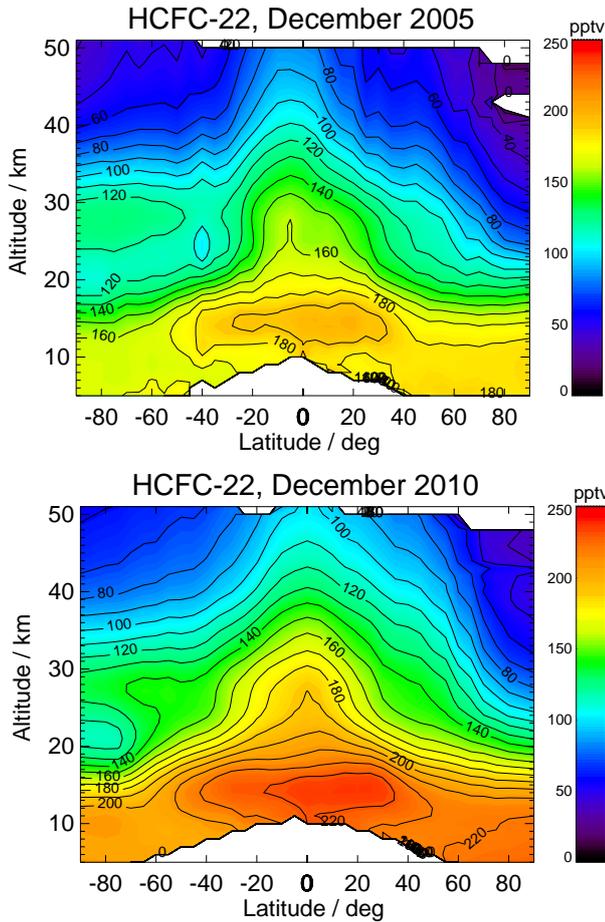


Figure 10. Monthly zonal means of HCFC-22 volume mixing ratios from 5 to 50 km for December 2005 (top) and December 2010 (bottom). The months have been selected to be approximately in the same QBO phase.

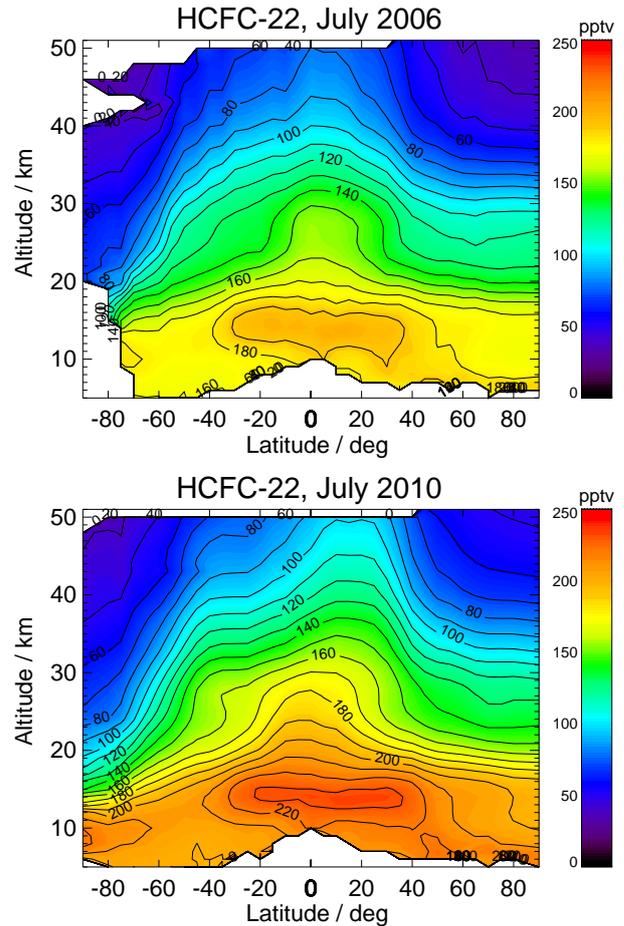


Figure 11. Same as Fig. 10, but for July 2006 (top) and 2010 (bottom)

The HCFC-22 distributions show a maximum in the tropical upper troposphere which is, at first glance, not expected for a source gas emitted at the ground. Closer inspection, however, reveals that in 2D distributions the situation is different, if we assume that there are extratropical localized sources and a localized uplift region and, after a certain altitude has been reached, injection into the zonal transport direction of the tropics. A source gas maximum as observed by MIPAS then can be explained by the following mechanism: Due to the local uplift of polluted air from localized sources the enhancement of the pollutant averages out to a certain degree when the zonal mean values for the lower altitudes are calculated. At higher altitudes the transport direction turns to zonal and averaging happens along the transport direction, involving summation over a series of enhanced values, and the reduction of the zonal mean by averaging over polluted and clean airmasses no longer takes place.

In the case of MIPAS HCFC-22, highest vmrs of up to 240 pptv are observed at the end of the observation period, and occur at altitudes between 10 and 15 km at 30 - 50°N during boreal summer and coincide with the position of the Asian monsoon anticyclone (Fig. 12, upper panel). After the break-down of the Asian monsoon anticyclone, during boreal fall, these HCFC-22 enhancements are spread over all longitudes and transported into the TTL, where they are subsequently distributed over the tropics, providing the isolated maximum layer of up to 225 pptv at low latitudes (Fig. 12, middle panel). This behaviour is nicely consistent with the general explanation scheme outlined above: HCFC-22 production was not restricted for developing countries during the relevant time, and some of these countries, e.g. China, have been large HCFC producers for years, while HCFC-22 production is controlled by the Montreal Protocol for industrialised countries. Indeed, Saikawa et al. (2012) found a surge in HCFC-22 emissions between 2005 and 2009 from developing countries in Asia with the largest emitting region including China and India. Also Montzka et al. (2009) sug-

gested a shift from high to low latitude emissions during this period, consistent with these assertions. High HCFC-22 abundances from these industrial regions of Asian developing countries are transported upwards into the Asian monsoon anticyclone during summer. Transport calculations based on meteorological analyses confirm that emissions from these regions indeed feed the Asian monsoon system (Vogel et al., 2015). After having been lifted into close-to-tropopause levels inside the Asian monsoon anticyclone, the high HCFC-22 abundances are transported into the tropical tropopause region (Fig. 12, lower panel). Transport from the Asian monsoon anticyclone into the tropical tropopause layer as a dominant source of tropical seasonality was suggested by Ploeger et al. (2012) based on model experiments, and by Randel and Jensen (2013). Once intruded into the TTL, the enhanced HCFC-22 abundances are distributed over all the tropical longitudes and generate an isolated maximum layer between appr. 10 km and the tropopause. This process also may explain why the isolated HCFC-22 vmr maximum observed by MIPAS above about 10 km exceeds abundances measured at the surface at remote sites in mid-latitudes by both the Global Monitoring Division of NOAA's Earth System Research Laboratory (NOAA/GMD) and the Advanced Global Atmospheric Gases Experiment (AGAGE) (c.f. Section 5.4).

5.2 Time series analysis of HCFC-22 for various altitudes and latitudes

The MIPAS HCFC-22 data presented here cover the so-called reduced resolution phase from January 2005 to April 2012. The instrument did not measure continuously in the nominal mode. This leads to frequent data gaps, but they are relatively short on average (1 to 2 days, occasionally longer). Some days were filtered out as well, since they contain too few measurements, and for certain phases there are data gaps in certain latitudes due to calibration measurements always performed at the same latitudes.

The averaged global time series of HCFC-22 volume mixing ratio from January 2005 to April 2012, including all latitudes from the South to the North Pole is shown in Figure 13. The global mean HCFC-22 volume mixing ratio increases at all altitudes with time. The global mean HCFC-22 volume mixing ratio at, for example, 16 km altitude was about 161 pptv in January 2005, and it increased up to about 210 pptv by April 2012. This provides us with a rough estimate of the increase of HCFC-22 content: it had increased by 49 pptv in 7 years, which is roughly 7 pptv per year.

For a more detailed analysis, we consider the mean mixing ratios in 20° latitude bands (Fig. 14 and 15). In these figures, it is again visible that the zonal mean abundance of HCFC-22 over the equator is much higher than over the poles at similar altitudes. In addition to the latitude dependence of the absolute volume mixing ratios, the oscillations due to the seasonal cycle are more pronounced at higher latitudes for

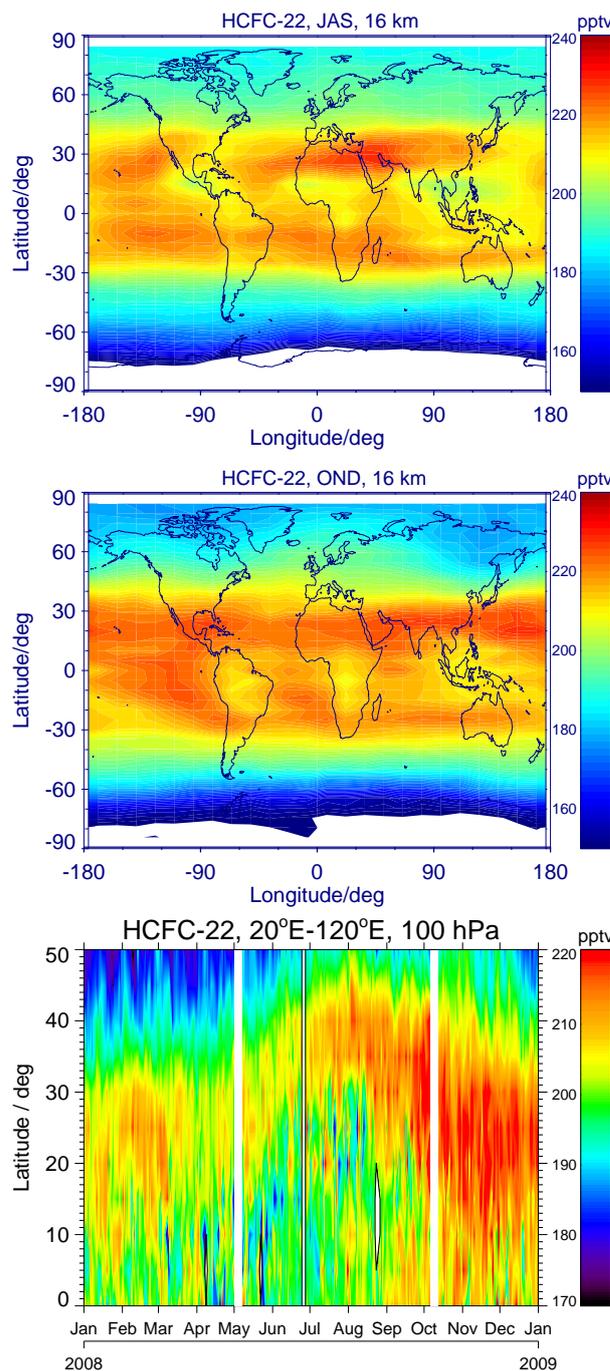


Figure 12. Top panel: Global HCFC-22 distribution in July/August/September for the years 2005 to 2011 at 16 km altitude. An enhancement over the Asian monsoon region is clearly visible. Middle panel: Global HCFC-22 distribution in October/November/December for the years 2005 to 2011 at 16 km altitude. The maximum previously located over the monsoon region has spread over all longitudes now. Bottom panel: The temporal development of HCFC-22 in the Asian monsoon region (i.e. daily averaged over 20°E to 120°E and shown for 0° to 50°N in 5° bins) at 100 hPa for the year 2008. Enhanced values are seen to propagate towards the tropics from August on.

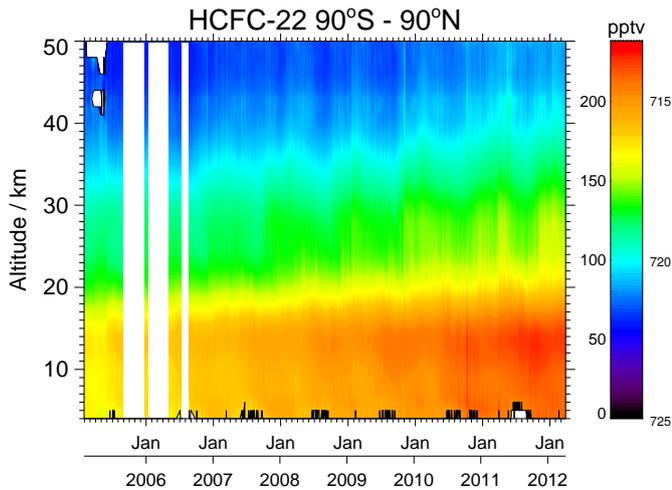


Figure 13. The globally averaged time series of HCFC-22 volume mixing ratio from January 2005 to April 2012.

680 the stratosphere, while a pronounced seasonal cycle in the 735
 upper troposphere can also be found at 30°N–50°N. In the
 stratosphere at the poles, any displacement of the polar vor-
 tex from the poles or its deformation will, in a 2D repre-
 sentation, cause numerical mixing, and by zonal averaging
 685 the sharp vortex boundaries are smeared out. In contrast, a 740
 breakup of the polar vortex would finally lead to physical
 mixing. In a 2D representation, physical and numerical mix-
 ing cannot be distinguished in any obvious way, but both pro-
 cesses would explain a rapid increase of stratospheric zonal
 690 mean polar HCFC-22 mixing ratios. Indeed, the onset of po- 745
 lar stratospheric wave activity has been verified to coincide
 in time with the observed increase of polar stratospheric
 HCFC-22 abundances. Interestingly, the sudden increase of polar
 stratospheric HCFC-22 seems to take place at all altitudes
 695 at almost the same time for the northern polar region, while 750
 for the southern polar region the increase of HCFC-22 is ob-
 served to start at around 30 km and to slowly move down.
 Thus low HCFC-22 abundances are observed to last longer at
 lower altitudes. This is explained by the fact that the increase
 700 at the northern latitudes is caused by wave activity along with
 numerical mixing as discussed above, while in the southern
 hemisphere it is caused by the breakdown of the polar vortex.
 The first kind of processes happens almost at the same time
 705 at higher altitudes and to proceed downwards. 755

In this context it is also interesting to see that a local max-
 imum of HCFC-22 appears just after austral vortex break-up
 around 30 km altitude (most pronounced in the years 2009
 to 2011), which indicates that young HCFC-22-rich air from
 710 low latitudes is rapidly transported into the polar region at 760
 these altitudes. A similar observation has been made within

the analysis of global distributions of mean age of strato-
 spheric air (AoA) (Stiller et al., 2012).

In mid-latitudes in the middle and upper stratosphere,
 there is no clear seasonal cycle visible, while at lower al-
 titudes, in the lower stratosphere (below 20 km), the max-
 ima are shifted towards hemispheric late summer/fall. This
 is consistent with the current picture of the phases of the
 Brewer-Dobson circulation, where the extratropical lower
 stratosphere is thought to be flooded by young tropical air
 during summer when the subtropical jet forms a weak mix-
 ing barrier only (Bönisch et al., 2011; Birner and Bönisch,
 2011; Stiller et al., 2012).

Figure 16 provides the time series of HCFC-22 as latitude-
 time cross sections for certain altitude levels. At 10 and 14
 km, high seasonality in the northern subtropical latitudes
 with maxima during summer and caused by the Asian mon-
 soon uplift is obvious. Around 20 km a clear seasonal cycle
 at high latitudes can be seen. Around 30 km, a QBO varia-
 tion in the tropics modulates the seasonal variation, while
 at 44 km the pattern is dominated by the semi-annual oscil-
 lation.

Revisiting Figure 13, it is obvious that the globally aver-
 aged distribution of HCFC-22 vmr in the stratosphere re-
 veals also seasonal oscillations. While one might expect that
 NH and SH seasonal cycles average perfectly out on a global
 scale, this seems not to be the case. This is due to the fact
 that the SH seasonal cycle is not quite in the opposite phase
 to the cycle in the NH (see Fig. 16). In addition, the HCFC-
 22 seasonal cycle in the SH is more pronounced than in the
 NH, which is attributed to the more stable polar vortex which
 dominates the SH seasonal cycle. Hence, the seasonal cycle
 in the SH has a higher amplitude and thus causes a resid-
 ual seasonal cycle on a global scale. The hemispheric tropo-
 spheric means of HCFC-22 vmr (Figure 16, top panel) for
 the NH are generally higher than for the SH. This is due to
 higher industrial production of HCFC-22 in the NH.

5.3 Trend of HCFC-22

We have analysed time series of monthly means at specific
 altitudes for 10°-wide latitude bands by fitting the following
 regression function to the data:

$$vmr(t) = a + bt + c_1 qbo_1(t) + d_1 qbo_2(t) + \sum_{n=2}^9 (c_n \sin \frac{2\pi t}{l_n} + d_n \cos \frac{2\pi t}{l_n}), \quad (1)$$

where t is time, a is the axis intercept, and b represents
 the linear component of the temporal variation, which for
 reasons of simplicity we call “trend”, without claiming
 that it has any climatological meaning beyond the time
 window under investigation. qbo_1 and qbo_2 are QBO indices,
 and the terms under the sum are 8 sinusoidal functions
 of the period length l_n . The terms qbo_1 and qbo_2 are the
 normalized Singapore winds at 30 and 50 hPa as provided
 by the Free University of Berlin via <http://www.geo.fu->

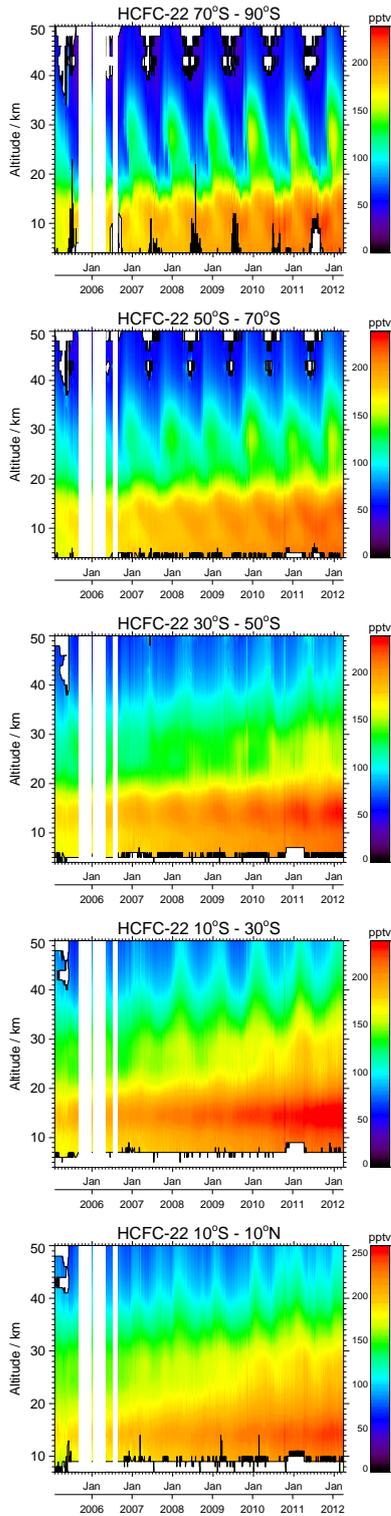


Figure 14. Time series of HCFC-22 vmrs from January 2005 to April 2012 for 70°S to 90°S, 50°S to 70°S, 30°S to 50°S, 10°S to 30°S and 10°S to 10°N (from top to bottom), generated from daily zonal means. White areas in the plots represent MIPAS data gaps. At low altitudes these are caused by clouds, while at high altitudes they are caused by a too low sensitivity of MIPAS. Data gaps covering all altitudes are associated with times when MIPAS did not measure.

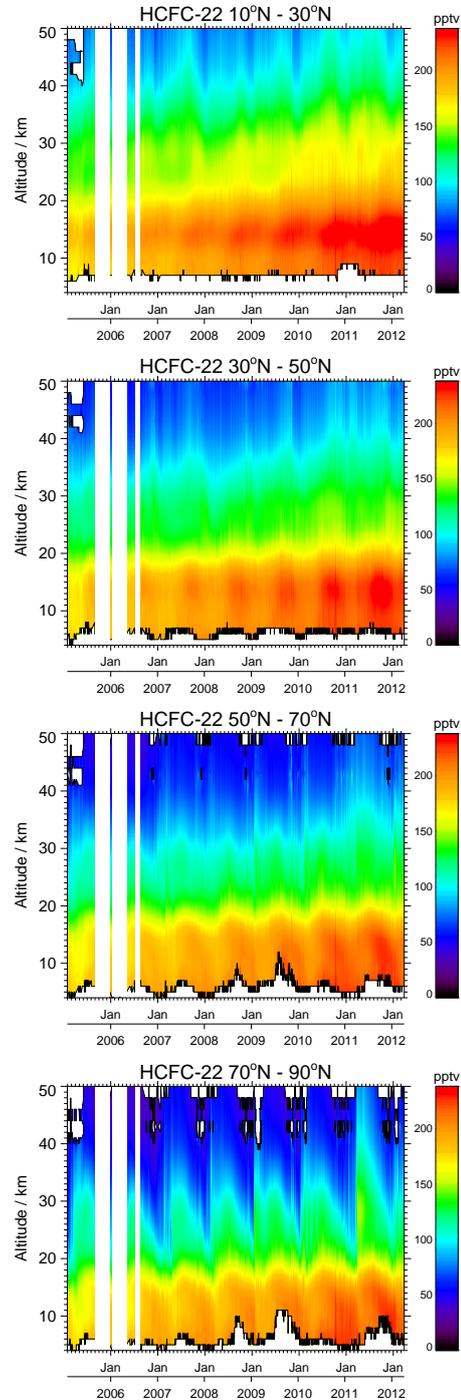


Figure 15. As Fig. 14, but for 10°N to 30°N, 30°N to 50°N, 50°N to 70°N and 70°N to 90°N (from top to bottom).

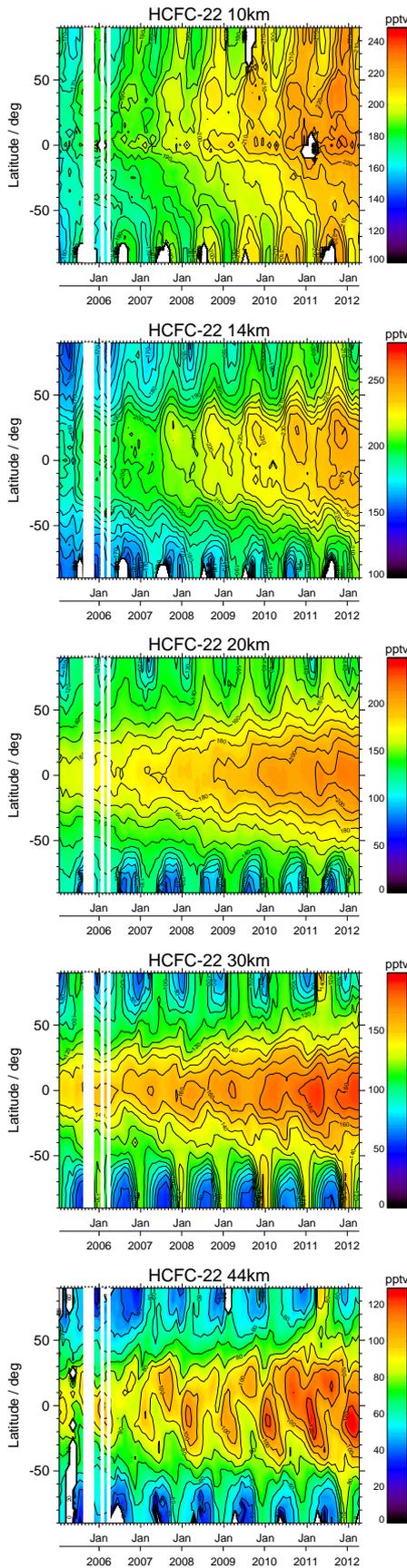


Figure 16. Latitude - time cross sections of HCFC-22 vmrs from January 2005 to April 2012 for altitudes of (from top to bottom) 10 km, 14 km, 20 km, 30 km, and 44 km, generated from monthly zonal mean data. Note different colour bars for different panels.

berlin.de/met/ag/strat/produkte/qbo/index.html. qbo_1 and qbo_2 are approximately orthogonal such that their combination can emulate any QBO phase shift (Kyrölä et al., 2010). Coefficients a , b , c_1 , ..., c_9 , d_1 , ..., d_9 are fitted to the data using the method by von Clarmann et al. (2010), where the full error covariance matrix of the HCFC-22 data is considered, with the squared standard errors of the mean (SEM) of the monthly zonal means as the diagonal terms. Further, a constant model uncertainty error term has been added to the data error covariance matrix, which represents the deficiencies of the regression model with respect to the true atmospheric variation and was, within an iterative procedure, scaled such that the resulting $\chi^2_{reduced}$ of the trend fit was close to unity, corresponding to combined data and model uncertainties consistent to the fit residuals. Since we cannot exclude that these perturbations to be accounted for by this additional error term have a typical duration of more than one month, covariance terms between adjacent data points were also considered in order to account for the resulting autocorrelation. Phase shifts of the variations are represented by common use of sine and cosine functions of the same period length. The first and the second sinusoidal functions represent the seasonal and the semi-annual cycles, and have the periods of 12 and 6 months, respectively. To model the deviations of the temporal variation from pure sine or cosine shapes, i.e. to allow for irregular shapes like sawtooth shapes etc., the period lengths of the remaining 6 terms under the sum are chosen to be equal to 3, 4, 8, 9, 18 and 24 months. The general strategy of this particular fitting has been described in Stiller et al. (2012).

Figure 17 provides some examples for time series and their fits for 3 different latitude bins and two altitudes. (40°S to 50°S , 0° to 10°S , and 50°N to 60°N , at 20 and 30 km, respectively). The lower panel of each figure provides the residuals between measured and fitted time series. The simple model is able to represent the observations very well in most cases. Besides the linear increase, for all altitude/latitude bins, a more or less pronounced seasonal cycle is the dominant feature of the time series. In the southern mid-latitudes, a clear QBO signal is also present. For the northern mid-latitudes at 30 km, we see also the impact of the semi-annual variation. The highest amplitudes in the temporal variation are reached in the 60° regions; in the inner tropics, the amplitudes are lower, particularly in the lower stratosphere. A strong linear increase is present in all altitude/latitude bins and varies considerably.

Fig. 18 summarizes the derived decadal trends for all latitude/altitude bins. The trend is positive for all latitude and altitude bins and highly significant (significance of 5σ or more). In the lower stratosphere below 20 km the trend is between 40 and 50 % per decade, while it varies between 30% per decade in the northern middle and high latitudes between 20 and 30 km, 50 to 60% per decade in the southern mid-latitudes from 20 to 50 km, and 70 to more than 100% per decade in the northern upper stratosphere at middle and high

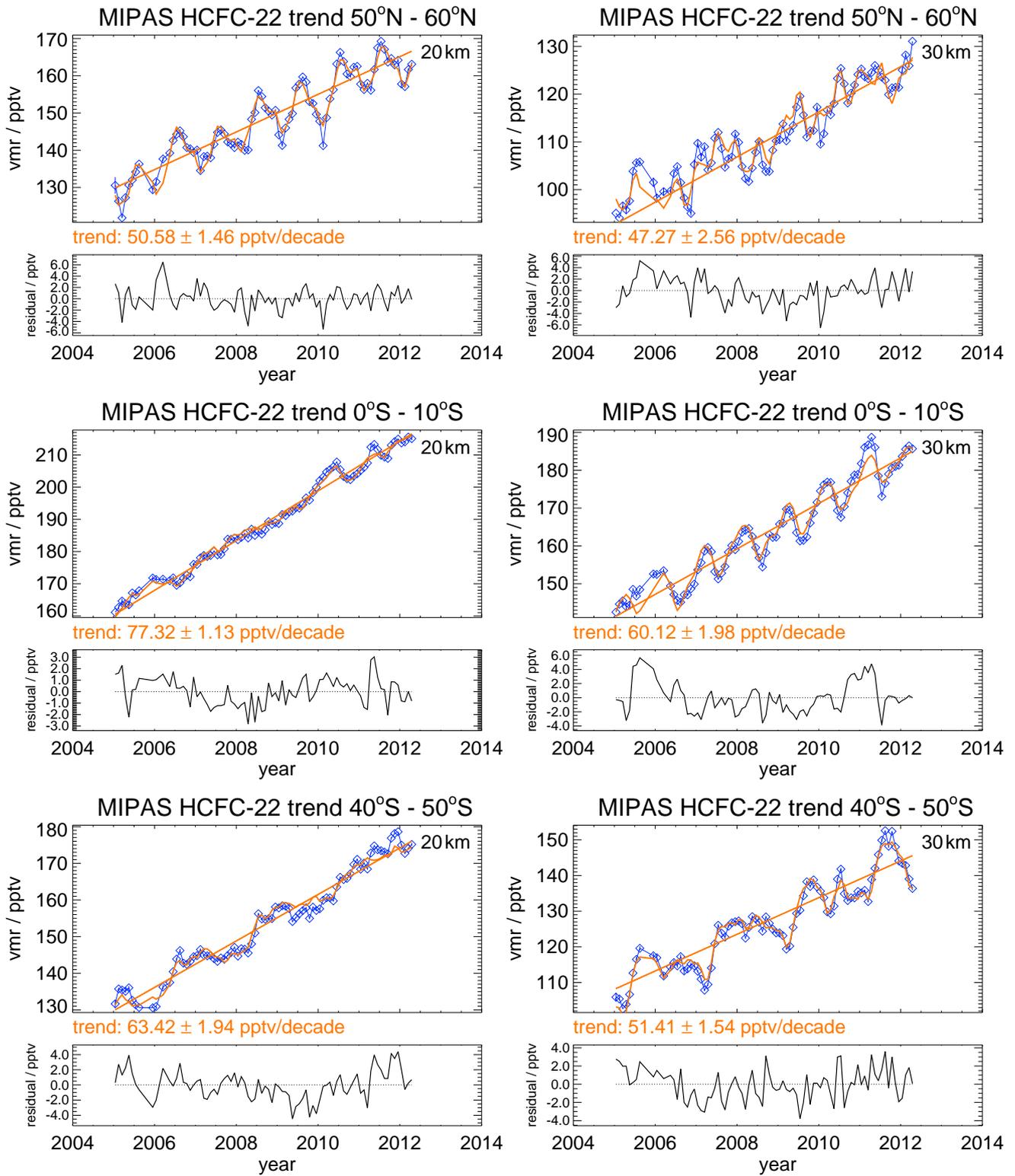


Figure 17. Time series of HCFC-22 volume mixing ratios (blue symbols) and their fits with the regression function described in the text (orange curves) at 20 km (left) and 30 km (right) altitude in three latitude bins. The linear trend component of the multi-parameter fit (straight orange line) is also shown.

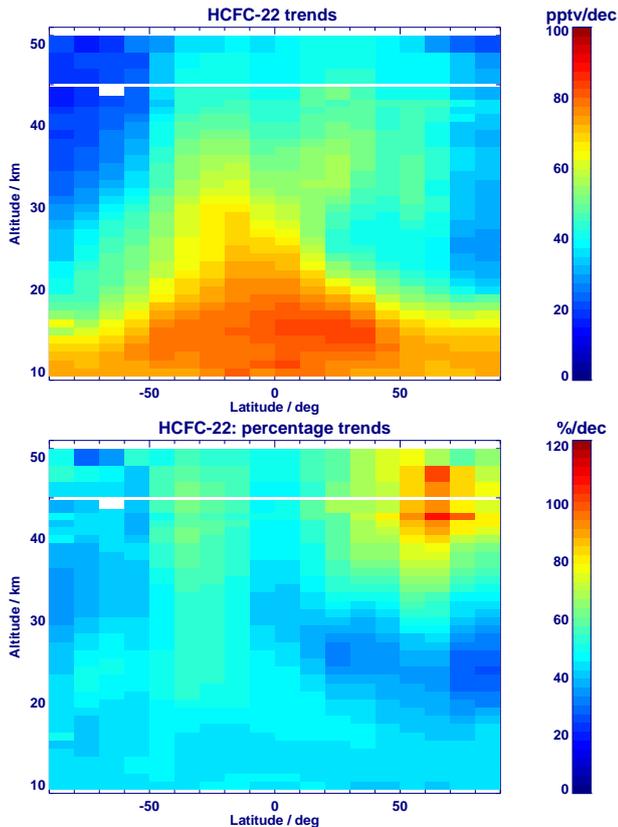


Figure 18. Trend of HCFC-22 volume mixing ratio for all latitude/altitude bins, in absolute units (pptv/decade) (top) and as percentage per decade relative to the background distribution in 2005 (i.e. the constant term of the regression analysis, see Eq. 1) (bottom). For the percentage trends, autocorrelations have been considered in the fit.

latitudes. The maximum of absolute trends is in the northern subtropics around 15 km, supporting our hypothesis that this region is fed by the strongly increasing East Asian emissions. In contrast, the relative trends are not highest in this region, due to the strongly enhanced volume mixing ratios measured there.

Recent HCFC-22 trends derived by earlier works for various time periods, altitudes and latitudes are summarized in Table 3 and compared to MIPAS-derived trends. It has to be noted that the relative trends are based on varying reference vmrs, most of which are not even reported. For this reason we discuss here the comparison of absolute trends only. In general, the trends derived from MIPAS observations are higher than the trends already published. The only exception are the trends at high southern latitudes by Moore and Remedios (2008) that were derived from a combination of ATMOS and MIPAS observations, and that are somewhat higher than our MIPAS-derived trends. One reason for the higher trends found by MIPAS is the inclusion of the Asian monsoon re-

gion in the zonally averaged trend data that are compared to the trends derived by other works, while those trends usually refer to single locations at remote sites.

5.4 Comparisons between upper tropospheric and surface growth rates

Two networks perform regular, long-term and highly precise near-surface measurements of various tracers, among them HCFC-22: these are the NOAA Earth System Research Laboratory (ESRL) Global Monitoring Division (GMD) and the Advanced Global Atmospheric Gases Experiment (AGAGE). Although not directly comparable, because MIPAS observations do not reach the ground, we compare here mean tropospheric values (below 10 km altitude) retrieved from MIPAS data with the surface observations from NOAA/GMD and AGAGE, making use of the fact that within each hemisphere the free troposphere can be considered well-mixed in the altitude domain. This assumption is confirmed by aircraft measurements (Xiang et al., 2014). In this context it should be noted that surface measurements are reported as dry air mole fraction, while MIPAS measurements are reported in mixing ratios where the air with its actual water vapour content is the reference. Since the stratosphere is very dry where MIPAS measures, this makes no discernable difference. Near surface, however, this difference has to be taken into account but since air dries during uplift, the surface dry air mole fraction is exactly the quantity which is comparable to the MIPAS results.

5.4.1 The Surface Networks

NOAA/GMD: NOAA/GMD runs flask measurements at remote site since 1992 (Montzka et al., 2009, 2015). These data are reported on the NOAA–2006 scale. These data are available via <ftp://ftp.cmdl.noaa.gov/hats/hcfc22/flasks/>. Global HCFC-22 data from NOAA/GMD in units of mole fraction in dry air at ground level are shown in Figure 19 for various measurement sites from the South Pole to the high Arctic. The data are not filtered for any pollution events, resulting in some enhanced values for the stations Trinidad Head and Mace Head that are occasionally influenced by nearby emissions. The MIPAS HCFC-22 monthly mean values for latitude bands selected to match the latitudes of the stations, averaged over all altitudes below 10 km, are shown for comparison with the same colour code as the respective station data.

AGAGE: AGAGE provides in situ measurements of a wide range of ozone depleting compounds and greenhouse gases, including HCFC-22, from several ground stations (Prinn et al., 2000, 2013; O’Doherty et al., 2004). AGAGE measurements used here are obtained using in situ gas chromatography with mass spectrometry (GC-MS) detection technique and are reported on the SIO-2005 calibration scale. NOAA flask results and AGAGE in situ data are compared

Table 3. Recent trends of HCFC-22 depending on time period, altitude and latitude from earlier publications, compared to MIPAS-derived trends at the respective altitude/latitude.

Source	Time interval	Latitude/Altitude	Absolute trend / pptvyr ⁻¹	Relative trend / %yr ^{-1a}	MIPAS-derived trend / pptvyr ⁻¹	MIPAS-derived trend / %yr ⁻¹
Moore and Remedios (2008)	1994 – 2004	20°N–50°N / 20 km	5.4 ± 0.7	3.5 ± 0.4 ^b	6.15 ± 0.24	4.24 ± 0.16
Rinsland et al. (2005a)	1994 – 2003	60°S–80°S / 20 km	6.0 ± 0.7	4.3 ± 0.5 ^b	4.88 ± 0.10	4.55 ± 0.09
	1985	30°N / lower strat.		14.57 ± 4.1 ^c		
	1994	30°N / lower strat.		6.35 ± 2.24 ^c		
	2004	30°N / lower strat.		3.92 ± 2.08 ^c	7.60 ± 0.21	4.56 ± 0.12
Rinsland et al. (2005b)	1987–2002	30°N / 2–10 km	5.66 ± 0.15	6.47 ± 0.17 ^b	7.54 ± 1.53	4.32 ± 0.87
Brown et al. (2011)	2004–2010	30°S–30°N / 8–17 km	6.56 ± 0.20	3.7 ± 0.1 ^b	8.02 ± 0.03	4.48 ± 0.02

^a The reference vmrs on which the relative trends are based vary from data set to data set.

^b Linear fit.

^c Derivatives at the respective years from a parabolic fit.

every six months at common sites. Comparison of HCFC-22 from NOAA flasks, using the NOAA–2006 calibration scale to AGAGE in situ measurements based on the SIO-2005 calibration scale at Cape Grim, Samoa, Trinidad Head and Mace Head reveal the following differences: The average difference (NOAA minus AGAGE) across the 4 sites is -0.7 ± 0.5 ppt or $0.35 \pm 0.25\%$. The differences are also relatively constant with time. These results are also very consistent to those found in the International Halocarbons in Air Comparison Experiment (Hall et al., 2014).

5.4.2 Comparison with MIPAS

Although NOAA/GMD and AGAGE provide independent data sets, no appreciable differences were found in any respect discussed here relative to MIPAS. Despite quite different approaches to making the measurements (in situ high frequency at relatively few sites compared to low frequency flask measurements at more sites), the surface data show the same broad features, distributions, seasonality, and trends. Thus, both data sets are discussed relative to MIPAS results together.

Overall, surface data from both networks and MIPAS upper tropospheric mixing ratios of HCFC-22 show a good agreement and similar trends (Figs. 19). The interhemispheric differences of the surface data are clearly visible, with mole fractions in the northern hemisphere being larger by about 10% than SH values, due to the main sources of HCFC-22 being located in the NH. MIPAS upper tropospheric mixing ratios agree best with tropical surface measurements, reflecting the fact that uplift of surface air is dominated by tropical processes.

In the SH, tropospheric MIPAS values are mostly significantly higher than the surface values at the same latitudes and reach these, if any, only during their seasonal minima (Fig. 19, top and middle panel). The reason is roughly this. MIPAS, whose measurements refer to the upper troposphere and above, sees more advected air from the tropical outflow,

and the signal is modulated by a pronounced seasonal cycle, while the related ground-based measurements are clean-air measurements. This hypothesis is in tendency confirmed by SH aircraft measurements (e.g. Xiang et al., 2014), where an indication of higher mixing ratios at higher altitudes is found, which is attributed to transport of NH air into the SH at higher altitudes in lower latitudes. It is interesting to see that among the SH time series, the seasonal cycle in MIPAS data is strongest for the southern polar latitude band (see Fig. 19, top panel), while the minima reached within this time series at the end of Antarctic summer are the lowest among all latitude bands. This strong seasonal cycle is also visible in Fig. 16 with maxima in polar southern spring and indicates flooding of the southern polar UTLS region with low-latitude air around the time of the polar vortex breakdown.

The MIPAS measurements show smaller differences between the hemispheres than the surface measurements. This is explained by the fact that MIPAS observes air at altitudes where the outflow of the intertropical convergence contributes to the composition of air. Assuming that air uplifted within the tropical pipe is mixed between both hemispheres offers an explanation for the reduced hemispheric contrast in the MIPAS data. Also these findings are in agreement with those of (Xiang et al., 2014).

Finally, the most obvious difference between surface time series and zonally averaged MIPAS upper troposphere time series is the pronounced seasonal cycle in the latter, with minimum values during NH spring and SH summer (see Fig. 19, top and middle panel). It is far more pronounced in the NH mid-latitudes than in the SH mid-latitudes and its amplitude increases towards high latitudes.

Xiang et al. (2014) observed a seasonality in the surface measurements of HCFC-22 with minima in northern summer and attributed this to increased scavenging through the OH radical reaction and seasonality in the transport. The best explanation of the observed seasonality, however, relies on an additional seasonality of the emissions of refrigerants with maxima in summer. This seasonality of emissions was de-

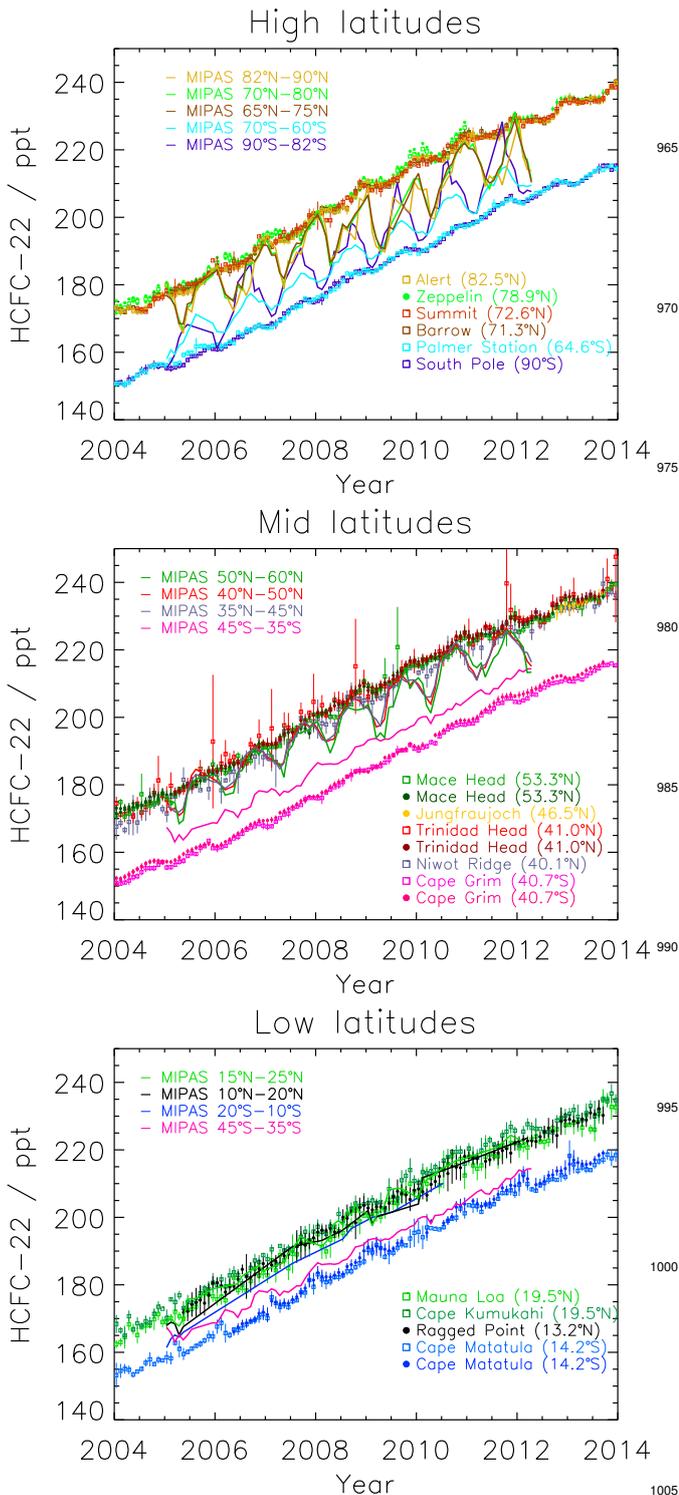


Figure 19. Comparison of HCFC-22 time series from several stations of the NOAA/GMD program and the AGAGE program with corresponding MIPAS values. Open squares: NOAA/GMD monthly mean data from several stations (see legend for colour code); filled circles: AGAGE monthly mean data from several stations (see legend for colour code); solid lines without symbols: MIPAS monthly mean data for 10° latitude bands around the respective station latitude; the colour coding matches that of the stations. The tick marks at the horizontal axis mark the beginning of the years. Top panel: high latitudes (between 60° and 90° N/S); middle panel: mid-latitudes (between 30° and 60° N/S); bottom panel: low latitudes (between 30° S and 30° N.)

rived by inverse modelling of (Xiang et al., 2014) aircraft measurements.

MIPAS maximum values fit perfectly to those of surface stations in the NH. While OH scavenging and seasonal variations in transport could possibly explain the summer minimum observed by MIPAS in the SH, the springtime minimum in the NH is more probably related to intrusion of HCFC-22-poor stratospheric air at the end of the polar winter and during polar vortex breakdown. Similar springtime minima were also observed for other tropospheric source gases and have been attributed to stratospheric air intrusions (Nevison et al., 2004, 2011). The fact that the minima in the UTLS in the MIPAS time series at higher altitudes precede those at lower altitudes (c.f. Fig 15 lowermost panel and Fig. 16, top two panels) supports this. The amplitude of the seasonal cycle seems to be even more enhanced due to transport of high HCFC-22 abundances uplifted within the Asian monsoon anticyclone to higher latitudes during fall (compare Fig. 16, top panel).

The SH upper tropospheric annual cycle is not exactly in the opposite phase of its northern counterpart. The latitudes of the SH stations shown in this comparison are, however, generally lower than the latitudes of the NH stations. At these low latitudes the upper tropospheric air is not modulated that much by polar processes but by the change of the position of the intertropical convergence zone and the Hadley cell. In local summer these latitudes are affected by the outflow of the tropical pipe, providing young HCFC-22-rich air. This hypothesis is corroborated by the fact that the MIPAS time series at 45° S– 35° S (pink solid line in Fig. 19) follows nicely the Cape Matatula (Samoa) (14.2° S, blue open squares and filled circles) time series, while it appears to be uncorrelated with the Cape Grim (Tasmania) (40.7° S) data (pink open squares and filled circles).

While according to MIPAS the HCFC-22 global mean below 10 km altitude (not shown) increased from 161 pptv to 211 pptv between January 2005 and April 2012, the NOAA ground-based global mean for these months were 164 pptv and 216 pptv, providing an increase of 50 (MIPAS) and 52 (NOAA) ppt, respectively. The growth rate derived from NOAA/GMD data for the NH is only slightly higher than that inferred from MIPAS measurements below 10 km (52 vs. 49 ppt over 7 years).

5.5 Unexplained stratospheric trends

Differences between the MIPAS stratospheric percentage trends (relative to the HCFC-22 vmr in 2005) and the tropospheric trends corrected for AoA from NOAA/GMD are shown in Fig. 20. We used the NOAA/GMD global trends, extended linearly with a slope of 57 ppt/decade before 1992, and AoA from Stiller et al. (2012), according to the method described in Kellmann et al. (2012). We would like to stress that we do not necessarily expect the stratospheric trends to reproduce the tropospheric ones with a time lag depending

on the AoA, because stratospheric circulation could change, affecting the time lag and possibly the level of HCFC-22 change in the air parcel observed. Furthermore the AoA spectrum could, in the case of a non-linearly increasing species, render the mean AoA non-representative. However, for an atmosphere without changes in stratospheric dynamics and δ -shaped age spectra, the differences shown in Fig. 20 are expected to be zero throughout. In contrast, we find positive and negative trend differences of up to 40 percent. The HCFC-22 trend in the northern mid-latitude lower stratosphere (15–30 km) is slightly smaller (up to -10%) than the trend at the ground at the time the air started its travel into the stratosphere, while in the southern mid-latitudes, the HCFC-22 trend is larger than the trend observed at surface level over all the stratosphere. In the polar regions, the trend is up to 40% lower at the South pole, and up to 40% higher at the North pole and the northern mid-latitudes. This pattern of trend differences is in agreement with the patterns found for CFC-11 and CFC-12 (Kellmann et al., 2012), and is in accordance with the AoA trends found by Stiller et al. (2012). Positive AoA trends in the northern mid-latitudes below 30 km and the southern polar region lead to increasingly longer exposure of HCFC-22 to loss processes in the stratosphere, and thus to a reduction of the positive HCFC-22 trend, while the general positive HCFC-22 trend is further increased in regions where the AoA becomes younger, leaving less time for stratospheric HCFC-22 loss processes. It should be noted that any incorrect assumption on AoA as such, used within the correction of surface trends, cannot explain both the positive and negative signs in the trend differences. The surface trend was slightly increasing all over the observation period covered by NOAA/GMD measurements, thus the correction by the AoA should always lead to smaller trends. Any incorrect assumption on the AoA would lead to incorrect differences, which, however, had always the same positive or negative sign. Thus, the occurrence of positive and negative trend differences is an unambiguous sign that the circulation must have changed over the MIPAS observation period. Our results further show that these circulation changes were different in the northern and southern mid-latitudes, as was concluded from the AoA data inferred from the MIPAS SF₆ data record (Stiller et al., 2012). These results confirm the hypothesis of Stiller et al. (2012) of interhemispheric asymmetries in circulation changes, for which further evidence has been found by Eckert et al. (2014) (interhemispheric asymmetries in decadal MIPAS ozone trends), Nedoluha et al. (2015) (interhemispheric asymmetries in decadal trends of N₂O from the Microwave Limb Sounder (MLS)), and Mahieu et al. (2014) (HCl trends and modelled age-of-air trends).

6 Summary and conclusions

HCFC-22 data from MIPAS for the period 2005 to 2012 were produced and analysed. Version 5 level-1b reduced res-

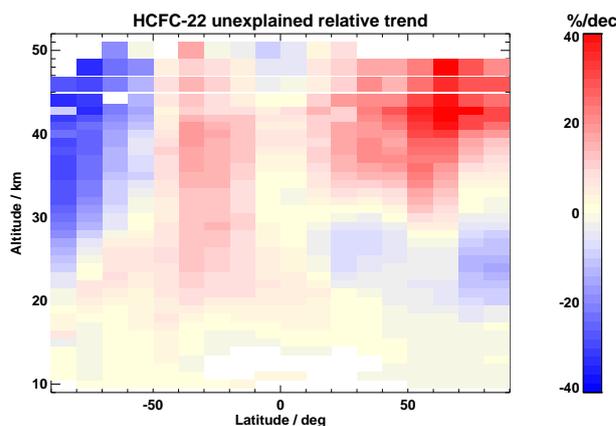


Figure 20. Differences between MIPAS stratospheric relative trends (referenced to the HCFC-22 background distribution in 2005, i.e. the constant term of the regression analysis (see Eq. 1) and relative trends at ground measured by the NOAA/GMD program (global means), the latter corrected for the time lag between the air parcels' start in the troposphere and its arrival in the stratosphere.

olution MIPAS measurements (nominal mode) of the period from 27 January 2005 to 8 April 2012 were inverted using the MIPAS IMK/IAA scientific data processor. The profile retrieval was performed by Tikhonov-constrained non-linear least squares fitting of measured limb spectral radiances. The total error of the retrieval is 7% on 20 km height and 9% on 30 km height, and the error budget is dominated by noise. A typical retrieved profile represents approximately 5 degrees of freedom, corresponding to an altitude resolution of typically between 3 km at 10 km height and 7 km at 30 km height, further increasing with height. The percentage of non-converged profiles is about 0.01 to 0.02%.

The profiles obtained were compared with ACE-FTS satellite data (v3.5), as well as with MkIV balloon profiles and in situ measurements performed by the University of Frankfurt. The comparisons are ambiguous with respect to a bias of MIPAS measurements; in general we can state that MIPAS agrees within ± 15 pptv (1σ) with ACE-FTS and MkIV the reference measurements. Between 13 km and 22 km, good agreement with MkIV and ACE-FTS profiles is demonstrated. A high bias of 30–50 pptv relative to cryosampler measurements was found below 24 km but no bias was found for higher altitudes.

The global distribution of HCFC-22 vmr reflects the mean circulation in the stratosphere and reveals also seasonal oscillations. The HCFC-22 annual cycle in the SH is more pronounced than in the NH. The HCFC-22 volume mixing ratio in the NH troposphere is generally higher than in the SH, due to main emission sources residing there. A volume mixing ratio maximum is situated at about 16 km height at low latitudes, which exceeds mixing ratios measured at all remote surface sites in the NH or SH. We attribute this to advection to low latitudes of HCFC-22-rich air uplifted in the Asian

monsoon area. The source of this HCFC-22-rich air could be South-East Asia. United Nations Environment Programme, 2012 reports that China has been the largest global producer and consumer of HCFCs for a number of years.

A multi-variate regression analysis was performed for 10°-latitude/1-2 km altitude bins, with terms for linear variations, sinusoidal variations with various periods, and a proxy for the QBO variation. We find positive linear trends for all latitude/altitude bins, ranging from 83 pptv/decade in the northern subtropical lower stratosphere to 18 pptv/decade in the southern polar upper stratosphere. The absolute trends are in the range of previous analyses. Percentage trends are highest in the southern mid-latitudinal stratosphere (50-70%/decade) and in the northern polar upper stratosphere (70 to 120 %/decade) and lowest in the northern lower stratosphere (around 30%/decade). The highest seasonal amplitudes are observed in the 60° regions; in the tropical lower stratosphere the amplitudes are low.

Global NOAA/GMD, AGAGE and MIPAS tropospheric values show good absolute agreement and similar trends. Based on the absolute values of HCFC-22 from MIPAS and on the MIPAS derived HCFC-22-growth rate, one can conclude that the HCFC-22 global volume mixing ratio in the lower stratosphere has risen by 49 pptv in 7 years. A pronounced seasonality has been detected in the upper troposphere with minima in spring in the NH and in local summer in the SH. The latter is attributed to the seasonality of the main tropical uplift and outflow regions. The seasonality in the NH is attributed to the intrusion of HCFC-22-poor stratospheric air at the end of the Arctic winter. Inconsistencies in percentage trends between ground-based and age-corrected MIPAS stratospheric trends hint at recent changes in stratospheric circulation. Similar indication has been found by analysis of trends of the mean AoA (Stiller et al., 2012), ozone (Eckert et al., 2014) and CFC-11 as well as CFC-12 (Kellmann et al., 2012). A more detailed analysis of these circulation changes is currently under investigation.

MIPAS HCFC-22 data presented here can be downloaded after registration from <http://www.imk-asf.kit.edu/english/308.php>. The HCFC-22 trends shown in Figs. 18 and 20 are provided as numerical values in the Supplement.

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