Abstract. HALOE observations of O₃, CH₄, HF, H₂O, NO, NO₂, and HCF collected during the October 1991 Antarctic spring period are reported. The data show a constant CH₄ mixing ratio of about 0.25 ppbv for the altitude range from 65 km down to about 25 km at the position of minimum wind speed in the vortex: i.e., the vortex center pressure versus latitude contours of NO, NO₂, HF, and HCF in this same region. Water vapor, HF, and HCF enhancement are also observed in the vortex center region above ~25 km. Between 10 and 20 km, the expected mixing ratio signatures exist within the vortex, but low values and deviations larger than 25 km pressure increase by 50%, and the ozone level doubled inside the vortex between October 11 and 24 in the 15 to 20 km layer. These changes predict a time constant for recovery from ozone hole conditions of 19 and 30 days for O₃ and H₂O, respectively. The data further show the presence of air inside the vortex between 3 and 30 mb which has mixing ratios characteristic of mid latitudes.

Introduction

The Halogen Occultation Experiment (HALOE) was launched September 12, 1991, by the Space Shuttle Discovery onboard the Upper Atmosphere Research Satellite (UARS) into a 57° inclination, 565 km, near-circular orbit. After allowing for a period of outgassing, the experiment started routine science observations on October 11, 1991. HALOE uses the experiment principle of satellite solar occultation and the instrument techniques of gas filter and broadband radiometry to sound the middle atmosphere temperatur, pressure versus, O₃, HCF, HF, CH₄, NO, NO₂, and H₂O vertical profiles [see Russell et al., 1993]. The solar occultation experiment coverage in the UARS orbit extends from ~80°S to 80°N over the course of a year. From October 11 through November 1, 1991, HALOE observed over the range from 50°N to 80°S, providing a large number of profiles of the parameters measured. These data were collected in the early-to-middle stage of the Antarctic ozone hole recovery phase. Since the UARS spacecraft conducts a yaw maneuver about every 32 days, other UARS experiments (i.e., the limb emission sounders which view perpendicular to the solar direction) were not observing the Antarctic region during this time. Because of the uniqueness of the HALOE observations and their implications for understanding photochemical and dynamical processes which are occurring. A concise and thorough review of theories explaining the ozone hole phenomena has been presented by Solomon [1990]. In addition, a wealth of data and a collection of recent theories on polar phenomena are contained in three JGR Special Issues on Polar Ozone and the Airborne Antarctic Ozone Experiment [Volume 97, D8, and Volume 94, D9 and D14, respectively]. It is now clear that for Antarctic ozone destruction to occur at the observed rate (~1 to 2% per day) in late September and early October, there must be low temperatures and concentrations of ozone and other gases

Observations

Figure 1 shows four ozone vertical profiles at 77°S on October 22, 1991. Note that the profiles vary longitudinally in an orderly way with the solid curve observed at 248°E giving the lowest mixing ratio. The total column ozone above 100 mb is 138 D.U. for this profile and 258 D.U. for the profile which occurs at 175°E. The 30 mb analyzed wind field and θ=700K potential vorticity (PV) map on this day provided by the European Centre for Medium-range Weather Forecasts (ECMWF) are shown in Figures 2a and b, respectively. Note that all HALOE measurements are made well inside the vortex (as defined by the maximum winds or as judged by quasi-horizontal [isentropic] PV gradients). The smallest column ozone occurs at the center of the vortex (set 7), while the largest value occurs nearer the vortex edge (set 13). Vertical profiles of CH₄ and HF on this day are shown in Figures 3 and 4. There is a longitudinal ordering of the profiles as it is for ozone. We will expand here on the definition of "vortex center." Note that the geographical position of minimum wind velocity (Figure 2a), located around (76S, 300E), is coincident with the methane profile, having a constant mixing ratio from ~65 km (0.1 mb) to ~25 km (25 mb) (solid curve in Figure 3). The point of zero
The occurrence of a constant CH₄ mixing ratio over such a broad altitude range (65 km to 25 km) is highly unusual and, to our knowledge, has not been observed previously. Like ozone, the smallest values occur at the vortex center, and the largest mixing ratios occur nearer the edge at 175°E. The same can be said of NO₂. The HF ordering is reversed, however, with the largest mixing ratio occurring at the vortex center. Methane and HF pressure versus longitude cross sections on October 22 are shown in Figures 5 and 6, respectively. The region of low CH₄ mixing ratio extends over the 2 mb to ~30 mb range and from ~240°E to ~360°E. There is also a local maximum in the 90°E to 240°E range centered at ~20 mb. HF, on the other hand, shows its largest stratospheric values over the 240°E to 360°E range and a local minimum in the region where CH₄ is a maximum. Note the close correspondence of features in the two cross sections, illustrating the excellent suitability of the HALOE CH₄ and HF measurements as tracers. Nitric oxide, NO₂, and O₃ exhibit the same sense of variation as CH₄, while H₂O varies in a similar way as HF. HCl shows a maximum in the 240°E to 360°E range, but no local features are apparent in the 90°E to 240°E region. This general picture persisted from October 18 through October 26 and for October 11-17 although, during the earlier period, the longitudinal distribution was different. After October 26, the HALOE measurement sample, which is determined by the spacecraft-earth-sun geometry, moved equatorward of 70°S. The data show that the CH₄ and HF patterns indicated in Figures 3 and 4 persisted up to 62°S at least through October 31, 1991. These patterns were gone by November 3 at 56°S, which is essentially outside the vortex.

Data Validation

An intensive data validation effort is being conducted by the HALOE Science Team. Comparisons have been made with past observations, model calculations, and measurements by other
UARS experiments. In addition, a few comparisons have been performed for balloon and aircraft observations. In all cases, for all HALOE parameters being measured, the agreement is good (on the order of ~20% or less differences). A paper by Tuck et al. [1993] compares HALOE CH4 and H2O data with results from the 2nd Airborne Arctic Aircraft Expedition (AASE-II). These results agree to within 13% or better. Based on these comparisons and our error studies, we conclude that the state of validation is sufficient to establish the correctness of the variations we are reporting.

Aerosol effects which can contaminate signals and cause increased mixing ratio errors are small or negligible in the Antarctic region over most of the time period discussed here. The data show that some high altitude (~25 km) aerosol layers are present, but the absorption occurs in well-defined layers and the data have been screened sufficiently so that any artificial retrieval effects due to aerosols will not change the character of the features we have discussed.

Discussion

The constant CH4 mixing ratio profile with a low value (~0.25 ppmv) implies that unmixed vertical descent at the vortex center is occurring, and this was invariably observed during the entire month of October. The only region in the vortex where the low values occur (240°E - 360°E) is in the vortex center as defined by the geographical position of zero winds where unmixed descent could be expected. The other variations reported are consistent with this explanation. HF-, HCIF-, and H2O-enriched air and NO- and NO2-poor air brought down from above would cause the patterns that are shown by the data for the 240°E to 360°E region (i.e., enhanced HF, HCIF, and H2O, but depressed NO and NO2). The concave upward curvature of the HF contours, as well as those of HCIF, NO, and NO2 not shown in this paper, also support the idea of horizontally unmixed vertical descent. The timescale for this vertical descent to occur is of interest; even if it had taken place over the whole period since vortex spin-up in March and April, the descent rate implied is about a scale height per month, and this rate is a lower limit assuming no horizontal mixing occurred over a half a year, a rather unlikely state of affairs. These descent rates are very large and are in line with the observationally based predictions of Danielsen and Houben [1988] and the deduced rates of Tuck [1989] and Proffitt et al. [1989b]. Recently, O'Neill et al. [personal communication, 1992] have corroborated this descent of mesospheric air into the stratosphere over a time scale of a few months using a numerical primitive equation model. The observed radial gradients of CH4 increase outward from the position of zero winds (the vortex center). The balance between descent and meridional motion must be such as to maintain unmixed air from aloft at the vortex center with increasing mixing ratio radially outwards towards the jet. To our knowledge, no model has reproduced this structure. The inhomogeneity with longitude and altitude inside the vortex during October also undermines attempts to treat polar ozone loss by applying the law of mass action to the vortex as though it was a single, well-stirred air mass.

The ozone hole is generally evident in the HALOE data between 12 and 25 km, and the expected signatures, e.g. dehydration, low O3, low NO2, are indeed observed on the days for which observations exist within the vortex, namely October 11 and 12 (sunrises) and October 17-18, and 21-24 (sunsets). Any time trend during this period is of interest, and the lower stratosphere time history of the zonal average H2O mixing ratio between 13 and 20 km is accordingly shown in Figure 7. Both water and ozone (not shown because of space limitations) show substantial recovery from ozone hole conditions during the 13 days. During the 13 days, the area of the vortex bounded by the -24 PV units contour at \( \theta = 425K \) (approximately at the center of the 15-20 km region) decreased by 7.7% from 29.9 million square kilometers on October 11 to 27.6 on October 24 (the PV map was provided by the ECMWF). Since all the observations on these dates were at latitudes and PV levels which occur within the vortex, the recovery in H2O must have a substantial component arising from descent of
undehydrated air from aloft, in addition to the horizontal component arising from the 7.7% erosion of the vortex. The time behavior of the average water vapor mixing ratios, increasing from 2.7 ppmv to 4.0 ppmv between October 11 and October 24, clearly demonstrates that air is being resupplied to the ozone hole with a dynamical time constant of approximately 30 days. The same number is obtained by using (2CH4 + H2O) a conservative quantity. The great inhomogeneity within the vortex basically makes the time constant approximate, but it is of the same magnitude as the time constant of the mixing ratio, which destroyed the ozone during late August and September. The ozone in the same region shows an increase from about 0.7 ppmv to 1.4 ppmv and is due to a combination of chemical regeneration and poleward air transport (i.e., resupply of undehydrated relatively ozone-rich air). This gives a shorter recovery time constant of 19 days, which has a 30-day dynamical resupply component (as indicated by the H2O changes), with the remainder presumably due to chemistry.

The observed mixing ratios of CH4, NO2, and O3 and local minima in HF and H2O in the 90°E to 240°E range and at pressures between 3 and 30 mb is an interesting problem. The vortex profile features over the same altitude range. The same is possible to do this comparison on October 22 using HALOE data and H2O and CH4 observations collected during the Second Airborne Arctic Stratospheric Experiment (AASE-II), Geophys. Res. Lett., In press, 1993. Tuck, A. F. et al., Polar stratospheric cloud processed air and potential vorticity in the northern hemisphere lower stratosphere at mid latitudes during winter, J. Geophys. Res., 97, No. D8, 7883-7904, 1992.


Summary

The HALOE data described in this paper show some unusual phenomena in the early-to-late October Antarctic spring period. The data are internally consistent with the observed feature being the same, but not the same altitude range. The same is also true for the other tracer, HF, thereby lending support to the explanation that mid-latitude air has moved horizontally into the vortex in the middle stratosphere.

Conducted to date verify the correctness of the variations reported to at least the 20% level. Methane and H2O have been validated to 13% or better by comparison with ER-2 observations. The time constant for the resupply of air to the ozone hole in the lower stratosphere 15-20 km region during October 1991 was about 30 days. The H2O recovery must surely occur dynamically, while the ozone restoration, since its time constant is shorter, presumably is driven by both dynamics and chemistry. More extensive time series studies are underway to determine time constants for O3, NOx, H2O, HFC, and HF during October. These studies should further elucidate the mechanisms operating during this phase of the ozone-hole recovery.

Acknowledgments. The authors are grateful to T. Davies of the ECMWF for providing the wind and potential vorticity fields.

References


R. J. Cicerone, Geosciences Department, 220 Physical Sciences Bldg., University of CA/Irvine, Irvine, CA 92717

P. J. Crutzen, Max-Planck-Institute fur Chemie, Postfach 3060, D-6500 Mainz, Germany.

S. R. Drayson, Space Research Building, 2455 Hayward, University of Michigan, Ann Arbor, MI 48105

L. L. Gordley, G&A Technical Software, 28 Research Drive, Hampton, VA 23665

J. E. Harries, SERC, Rutherford Appleton Laboratory, Chilton, Didcot, Oxon, OX11 0QX, United Kingdom

J. H. Park, Mail Stop 401A, NASA Langley Res. Center, Hampton, VA 23681-0001

J. M. Russell III, Mail Stop 401B, NASA Langley Research Center, Hampton, VA 23681-0001

D. Tuck, NASA Aeronautic Laboratory, Code R/E/AL6, 325 S. Broadway, Boulder, CO 80303

Received December 14, 1992
Revised February 4, 1993
Accepted March 2, 1993