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Tracing troposphere-to-stratosphere transport above a mid-latitude deep convective system

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Abstract. Within the project SPURT (trace gas measurements in the tropopause region) a variety of trace gases have been measured in situ in order to investigate the role of dynamical and chemical processes in the extra-tropical tropopause region. In this paper we report on a flight on 10 November 2001 leading from Hohn, Germany (52°N) to Faro, Portugal (37°N) through a strongly developed deep stratospheric intrusion. This streamer was associated with a large convective system over the western Mediterranean with potentially significant troposphere-to-stratosphere transport. Along major parts of the flight we measured unexpectedly high NOx mixing ratios. Also H2O mixing ratios were significantly higher than stratospheric background levels confirming the extraordinary chemical signature of the probed air masses in the interior of the streamer. Backward trajectories encompassing the streamer enable to analyze the origin and physical characteristics of the air masses and to trace troposphere-to-stratosphere transport. Near the western flank of the streamer features caused by long range transport, such as tropospheric filaments characterized by sudden drops in the O3 and NOx mixing ratios and enhanced CO and H2O can be reconstructed in great detail using the reverse domain filling technique. These filaments indicate a high potential for subsequent mixing with the stratospheric air. At the south-western edge of the streamer a strong gradient in the NOx and the O3 mixing ratios coincides very well with a sharp gradient in potential vorticity in the ECMWF fields. In contrast, in the interior of the streamer the observed highly elevated NOx and H2O mixing ratios up to a potential temperature level of 365 K and potential vorticity values of maximum 10 PVU cannot be explained in terms of resolved troposphere-to-stratosphere transport along the backward trajectories. Also mesoscale simulations with a High Resolution Model reveal no direct evidence for convective H2O injection up to this level. Elevated H2O mixing ratios in the ECMWF and HRM are seen only up to about tropopause height at 340 hPa and 270 hPa, respectively, well below flight altitude of about 200 hPa. However, forward tracing of the convective influence as identified by satellite brightness temperature measurements and counts of lightning strokes shows that during this part of the flight the aircraft was closely following the border of an air mass which was heavily impacted by convective activity over Spain and Algeria. This is evidence that deep convection at mid-latitudes may have a large impact on the tracer distribution of the lowermost stratosphere reaching well above the thunderstorms anvils as claimed by recent studies using cloud-resolving models.

1 Introduction

Reactive nitrogen (NOx) has a pivotal influence on the ozone chemistry of the troposphere and the stratosphere. While in the stratosphere (above 18 to 20 km height) nitrogen oxides (NOx=NO+NO2) destroy ozone in a catalytic cycle and therefore contribute to ozone depletion (IPCC, 1999), in the troposphere they act together with hydrocarbons and carbon monoxide (CO) as important precursors of ozone formation. Also for the lowermost stratosphere (LMS), the region between the tropopause and the 380 K potential temperature surface (Holton et al., 1995), numerical simulations show that nitrogen oxides tend to enhance O3 mixing ratios (IPCC, 1999). Furthermore, O3 represents a strong greenhouse gas in the tropopause region and an increase in O3 mixing ratios at this altitude may have significant influence on the radiative forcing and hence on surface temperatures (Lacis et al., 1990; Forster and Shine, 1997). Likewise water vapor in the upper troposphere and lower stratosphere has important radiative and chemical effects (Oltmans and Hofmann, 1995). Nevertheless, for both stratospheric O3 as well as for H2O
there is considerable uncertainty concerning their trends in
the tropopause region (IPCC, 2001).

The budget of NO\textsubscript{x} in the LMS is largely determined by its
main source, the downward transport from the lower strato-
sphere, where odd nitrogen is produced through the reac-
tion between N\textsubscript{2}O and O\textsuperscript{17}D (Brasseur and Solomon, 1986).
Other sources may be transport from the extratropical tropo-
osphere along isentropic surfaces (Hoerling et al., 1993; Dessler et al., 1995; Dethof et al., 2000) and convective in-
jection in overshooting thunderclouds (Poulida et al., 1996; Strörm et al., 1999). This tropospheric air may be enriched
in NO\textsubscript{x} through production of NO by lightning or upward
transport of polluted air from the planetary boundary layer.
Also, sedimentation of polar stratospheric cloud particles in
the polar vortex can lead to redistribution of NO\textsubscript{x} and a nitric-
fication of the LMS down to altitudes as low as 10–12.5 km
(Fischer et al., 1997; Koike et al., 2002). In addition, air-
craft may directly emit NO\textsubscript{x} into the LMS. Estimates suggest
that 20–40% of total current aircraft emissions are released in
the LMS (Hoinka et al., 1993; Schumann, 1997). While
emissions by aircraft are well quantified, large uncertainties
remain concerning the source strength of NO\textsubscript{x} by lightning
(Lee et al., 1997) or the importance of deep convective events
first reported on distinct plume features visible as cirrus
cloud layers above the anvil of convective storms in im-
gages from the Advanced Very High Resolution Radiometer
(AVHRR) on NOAA polar orbiting satellites. Recent model-
studies by Mullendore (personal communication, 2003) and
Wang (2003) show that the breaking of gravity waves at the
top of deep convective systems in mid-latitudes can cause
injection of water vapor and other tropospheric tracers into the
lowermost stratosphere in the form of plumes well above
the thunderstorm anvils. Experimental evidence for the in-
trduction of aerosols from forest fire smoke into the lower
stratosphere has been given by aircraft measurements dur-
ing the STREAM II campaign (Waibel et al., 1999) and by
POAM satellite observations (Fromm et al., 2000). Another
mechanism for cross-tropopause mixing is described by Zierl
and Wirth (1997) where radiation causes a sinking of the
tropopause in upper tropospheric anticyclones leading to a
net troposphere-to-stratosphere mass flux. This again may
lead to an enhancement of the NO\textsubscript{x} mixing ratio in the low-
nermost stratosphere as NO\textsubscript{x} mixing ratios in the upper tropo-
sphere at mid-latitudes are higher than in the tropics (Fischer
et al., 2000). Finally, NO\textsubscript{x} is a chemically long-lived tracer
in the stratosphere, and stratosphere-to-troposphere transport
(STT) represents the main sink of NO\textsubscript{x} from the stratosphere.

The composition of the lowermost stratosphere changes
with season and location and is mainly influenced by the
relative importance between residual-mean downward trans-
port from the stratospheric overworld above 380 K and up-
ward transport from the troposphere across the extratropi-
cal tropopause. Further investigations are needed to quan-
tify the amount of tropospheric air transported into the LMS
by convective activity as opposed to isentropic transport and
how this alters the chemical composition and in particular the
NO\textsubscript{x} mixing ratios of the LMS.

In this paper we present a case study providing experi-
mental evidence that troposphere-to-stratosphere transport
(TST) caused by mid-latitude intense thunderstorms is able
to strongly influence the chemical composition of the lower-
most stratosphere. These findings tie in with the results by
Fischer et al. (2003) showing injection of boundary layer air
into the LMS by deep convection during the MINOS cam-
paign and by Poulida et al. (1996) on the observations of ex-
tensive stratosphere-troposphere exchange (STE) within an
intense mid-latitude thunderstorm over North Dakota. In the
project SPURT (trace gas transport in the tropopause region)
high resolution in-situ measurements of a variety of chemi-
tral tracers were performed in order to investigate their dis-
tribution and seasonal variability and to gain better insight
into the role of dynamical and chemical processes shaping the
tropopause region. We focus on the measurements ob-
tained during the SPURT mission flight on 10 November
2001 leading through a spectacularly deep stratospheric in-
trusion which developed over south-western Europe on that
day. The whole episode from 4 to 19 November was very
active in terms of meteorological phenomena over the North
Atlantic and triggered a number of other investigations. For
instance Stohl et al. (2003) reported on fast transport of air
pollution from North America to Europe supported by an ex-
plosive cyclogenesis near Greenland on 7 to 11 November as
traced by GOME satellite observations of NO\textsubscript{x} and simulated
using a trajectory parcel model. Thomas et al. (2003) ana-
lyzed a severe weather event over Spain and Algeria and its
impact on the distributions on O\textsubscript{3} and NO\textsubscript{x} using combined
GOME NO\textsubscript{x} observations and simulations of a stratospheric
chemical transport model (ROSE), and concluded that about
two-third of the total atmospheric content of NO\textsubscript{x} was lo-
cated in the troposphere above cloud levels. Martius et al. (in
preparation, 2004) present a detailed analysis of the dy-
namical situation and the formation mechanism of the deep
stratospheric intrusion.

2 Chemical measurements

The data presented in this paper have been measured on
10 November 2001 on a flight from Hohn (52° N), Ger-
many, to Faro (37° N), Portugal (see Fig. 1). A Lear-
jet 35A which reaches altitudes of up to 13 km was
used as experimental platform. Total reactive nitro-
gen (NO\textsubscript{x} = NO + NO\textsubscript{2} + NO\textsubscript{3} + HNO\textsubscript{2} + HNO\textsubscript{4} + HONO
+ PAN + RONO\textsubscript{2} + ClONO\textsubscript{2} + 2N\textsubscript{2}O\textsubscript{5} + BrONO\textsubscript{2} + or-
ganic nitrate + particulate nitrate (<1 µm)) was measured
by chemiluminescence, by letting NO react with O\textsubscript{3} after
reducing NO\textsubscript{x} species to NO using an externally mounted

1Martius, O.: A case-study on the dynamical effects of a hurri-
catalytic gold converter with CO as reduction agent (Fahey et al., 1985). The converter samples in backward direction in order to exclude particles with diameter > 1 μm. For technical details of the measuring system see Lange et al. (2002). Previously the measurement system has been implemented successfully on several STREAM campaigns (Fischer et al., 2000).

The conversion efficiency of the analyzer was quantified by three in-flight calibrations with known amounts of NO2 and varied between 90 and 94%. Similarly, the sensitivity was calibrated twice during the flight by adding known amounts of NO and varied by less than 3.5%. The chemiluminescence detector was switched every 9 s to pre-chamber mode in order to determine the temporal evolution of the instrument background signal (Ridley and Howlett, 1974). In addition, zero calibrations with synthetic air were performed four times during the flight to determine the temporal evolution of the NO2 artifact signal which is a common feature in NO2 measurements. Its origin is not fully understood but is in part due to impurities of the CO reduction agent (Fahey et al., 1985). Note that the specific design of the converter allows adding all calibration gases upstream of the converter at the inlet tip. The precision of the NO2 measurement has been determined by the standard-deviation of the in-flight NO2 calibrations to ±8% (2σ). The accuracy includes the uncertainty of the calibration gas standards (±4% (2σ)), and the uncertainty in the conversion efficiency for different NOy species (±13% (2σ)). During the first flight hour of the flight on 10 November 2001, the overall accuracy was very low (±50% (2σ)) mainly due to an initial contamination of the converter but improved to the normal level of about ±16% (2σ) for the rest of the flight. The reason of the contamination and its effect on the background signal have been investigated in detail. A possible reason could have been pollution of the converter through sampling of air and aerosols in the airport environment. This is not likely as the converter is flushed during the ascent phase with synthetic air. The enhanced background was found to result primarily from a startup procedure during takeoff using for security reasons ambient air instead of pure O2 to generate the reaction agent O3. Its influence exhibits an exponential decrease and could be reproduced in laboratory studies. An additional influence on the background signal was also due to the calibration gas NO2 entering the converter accidentally during the ascent phase of the aircraft as well as during zero air and NO calibrations. The influence of this problem and its temporal evolution has been estimated in different tests on subsequent flights. Both contaminations were accounted for in the evaluation of the raw data. The artifact signal is normally determined by linear interpolation of the signals measured between subsequent zero air calibrations. Due to the accidental contaminations an additional not well characterized offset had to be subtracted. By subtracting this signal, modeled as an exponentially decreasing function of time, the zero air calibration values were reduced to the normal level observed on other flights. The uncertainty associated with this procedure was conservatively assumed to be 50% of the subtracted signal itself and sensitivity tests with other exponential fits showed that corrections larger or smaller than this range would result in an unrealistic background signal level (e.g. lower than pre-chamber mode signals) and/or negative NO2 concentrations.

O3 was measured by a UV absorption photometer (accuracy ±5%, precision ±2.5 ppbv, Mottaghy, 2001), and H2O by a Lyman-α-Hyrometer with an overall accuracy better than ±5% (Zöger et al., 1999). N2O and CO data were sampled with a Tunable Diode Laser Absorption Spectrometer (TDLAS) with total uncertainty of the measurements of less than ±2% (Hoor et al., 2002).

3 Models and diagnostic methods

To support the analysis of the observations extensive use was made of trajectory calculations and analyses of meteorological fields from a synoptic-scale global and a mesoscale regional model. The two models and the applied methods are briefly described here.

3.1 ECMWF data and trajectories

We used analyses of the European Centre for Medium-Range Weather Forecasts (ECMWF) available on 60 vertical levels from the surface up to 0.01 hPa for the description of the meteorological situation and analysis of the dynamical processes. The 6 hourly analyses were complemented by intermediate 3-h forecasts. Sequences of the three-dimensional wind field were used for the calculation of trajectories with the Lagrangian Analysis Tool (LAGRANTO)(Wernli and Davies, 1997). Several variables characterizing the physical state of the air parcels including temperature, specific humidity and potential vorticity were traced along the trajectory paths.

Four different types of trajectories have been calculated: (i) 10-day backward trajectories were started from the exact location of the aircraft in time and space every 10 sec along the flight path. (ii) For use with the Reverse Domain Filling technique we calculated a total of 1.5 Mio 10-day backward trajectories from a specified grid from three different starting times as explained in Sect. 3.2. (iii) Additional 1.5-day backward trajectories from three distinct levels of the same grid at 200, 250, and 270 hPa and starting times of 09:00 and 12:00 UTC to trace convective influence and (iv) 10-day forward trajectories started from the same grid for starting times of 09:00 and 12:00 UTC.

3.2 Reverse Domain Filling

The Reverse Domain Filling technique (RDF)(Schwoerbel and Newman, 1995) has been applied in this case study to investigate the three-dimensional fine-scale structure of the
into the LMS. The CHRM's parent model is the High Resolution Model (HRM) of the German Weather Service (DWD). It is a hydrostatic model using Kessler-type (Kessler, 1969) grid-scale cloud microphysics as refined by Lin et al. (1983) and the mass flux convection scheme after Tiedtke (1989). The model simulations have been initialized by the ECMWF analysis at 00:00 UTC on 9 November 2001 and have been run for 48 h. They were initialized on forty layers between 1010 hPa and 15 hPa in height, with an enhanced density of levels in the tropopause region. The grid-resolution was 0.25° between 26° N to 58° N and 21° W to 21° E.

4 Meteorological situation

Figure 1 shows the potential vorticity distribution on a hybrid model level corresponding to about 230 hPa on 10 November 2001 as analyzed by ECMWF and the flight track overlaid in black. The whole period from 4 to 19 November spanned a series of remarkable meteorological events. On 4 and 5 November two hurricanes, Michelle and Noel, were located over the western North Atlantic. Strong Rossby wave activity led to significant north-south excursions of air masses. Over the central Atlantic subtropical air masses were advected northwards forming a ridge which extended from the eastern coast of North America to northern Europe. The ridge was fed by diabatic processes associated with several cyclones and the aforementioned hurricanes, and was associated with a particularly strong anticyclonically curved jet stream. Lifting of the tropopause in the ridge and the northward advection of ozone-poor subtropical air (Koch et al., 2002) resulted in an ozone mini-hole on 9 November centered over Scotland with minimum ozone columns of only 190 DU as seen by the GOME instrument (http://www.esa.int/export/esaCP/ESAHFRQQSTC_index_0.html). To the east of the ridge a Rossby wave is observed to be breaking with both poleward and equatorward components very similar to the meteorological situation on 5 June 2000 described by Bradshaw et al. (2002). The poleward component is seen in Fig. 1 as a thin tropospheric intrusion of low PV extending from Scandinavia up to the northeastern tip of Greenland. The equatorward component is seen as an elongated and deep stratospheric intrusion in the area of the flight path consisting of Arctic air which had moved rapidly from Scandinavia to the Iberian peninsula during the previous two days. Maximum tropopause pressures in the intrusion were as high as 650 hPa. This streamer was associated with a surface low that caused severe flooding in Algeria (Thomas et al., 2003). On its forward flank the streamer triggered a series of deep convective storms across the western Mediterranean from the Algerian coast towards Sicily and the Apennines. For a detailed analysis of the dynamical situation and the physical mechanism related to the formation of the streamer see Martius et al. (in preparation, 2004). On the main part of the flight from

Hohn to Faro, the Learjet flew within the streamer reaching \(\Theta\)-levels up to 365 K and probing air masses with relatively high potential vorticity (maximum=10 PVU). In segment I of the flight (as indicated in Fig. 1) and again briefly before landing (including the air masses of flight segment III), and also on the return flight later on the same day, the aircraft probed the subtropical air masses described above. In flight segment II (between 09:36 and 11:15 UTC), the probed air masses originated from the polar region with maximum PV-values along the backward trajectories of 12 PVU (for more details see Sect. 5.3).

5 Results and discussion

In Fig. 2 time series of the measured trace gases and meteorological quantities during the flight on 10 November 2001 are depicted. Different flight segments are shaded in grey and labeled with numbers I to III. Their geographical position is indicated in Fig. 1. In the following sections we discuss the features in the trace gas measurements of the different flight segments. In Fig. 3a and b, showing the correlations between measured \(O_3\) and \(NO_y\) and between \(O_3\) and \(H_2O\), respectively, the different characteristics of the air masses of the three segments are clearly distinguishable.

5.1 Flight-segment I

The most striking feature in the tracer-measurements is found in the first segment at the entrance to the streamer (label I in Figs. 1, 2 and 3). While flying on a constant pressure level of 215 hPa, potential vorticity (Fig. 2d) and potential temperature (not shown) were steadily increasing. However, the concurrent increases in \(NO_y\) and \(O_3\) mixing ratios were interrupted three times by sudden drops. Mixing ratios of \(NO_y\) decreased from 1.3, 3.3 and 3.6 ppbv to 0.9, 1.4 and 1.1 ppbv, while \(O_3\) mixing ratios decreased from 150, 250, and 320 ppbv to values of 90, 120, and 130 ppbv, respectively. The second and third drop were accompanied by simultaneous increases in the CO mixing ratio by about \(\Delta[CO]=30 \text{ ppbv}\). As shown by the minimum PV values depicted in Fig. 2d, the 10-day backward trajectories started along the flight track and coinciding with the drops in \(NO_y\) and \(O_3\) partly originated in the troposphere or at least close to the tropopause, where we would expect to find smaller \(O_3\) and higher CO mixing ratios than in the stratosphere. In the upper troposphere \(NO_y\) mixing ratios are also expected to be low provided that the air masses experienced no recent influence of deep convection of polluted planetary boundary air or lightning activity. Values for upper tropospheric air lie between 500 and 1000 pptv as measured during subsequent SPURT winter and autumn campaigns or as reported by Weinheimer et al. (1994). The sharp features of the tracers imply that the mixing between the stratospheric and the tropospheric air masses has not been completed yet, but that rather the air masses still represent distinct filaments of air with tropospheric and stratospheric characteristics. We used the RDF technique as described above to reproduce the three dimensional structure of the tropospheric filaments similar to Beuermann et al. (2002).

Figure 4 shows that except for the first drop and in contrast to the standard ECMWF analysis (Fig. 4a), rPV fields reproduce the structures in the measurements in almost perfect agreement (Fig. 4b). The horizontal scale of the filaments

![Time series of trace gases and meteorological parameters during the SPURT mission flight on 10 November 2001. (a) Measured \(O_3\) (red), measured \(NO_y\) (black) and its 2\(\sigma\) uncertainty (blue shading) and \(NO_y\) as expected from a correlation with \(O_3\) (green) (see text). (b) Measured CO (black) and flight pressure altitude (red). (c) Measured total water (black) and water vapor mixing ratio (in ppmv) from ECMWF analyses interpolated in space and time onto flight track (red). (d) PV from ECMWF analyses interpolated onto flight track (black) and minimum and maximum PV along the 10-day backward trajectories started from the flight track (red lines). Grey shading highlights the three different flight segments I, II and III (see Fig. 1 and text for details).](https://www.atmos-chem-phys.org/acp/4/741/)

Atmos. Chem. Phys., 4, 741–756, 2004
was between 50 and 100 km in width and several 1000 km in length. Their height reached up to a potential temperature level of 370 K, which is normally well within the lowermost stratosphere at mid-latitudes. Evaluation of the backward trajectories shows, that the development of the filaments was initiated on 8 November 2001 by the wrapping of tropospheric air around an almost cut-off stratospheric trough over Newfoundland. This was followed on the next two days by a continuous zonal stretching and meridional compression of both the tropospheric air mass and the trough, now located to the south of the tropospheric air, during their rapid transit across the North Atlantic. Observations of similar yet less pronounced filaments along the western flank of a stratospheric streamer have been reported previously by Bradshaw et al. (2002). They showed that the layering was generated by the breaking of a Rossby wave in the presence of vertical gradients in the background wind fields. The successful reconstruction of the tropospheric filaments coinciding with the drops in the high-spatial resolution tracer measurements shows that the calculated backward trajectories are reliable and that the RDF technique produces an accurate picture of the small scale structures not seen in the PV-fields of the original ECMWF analyses. The filaments represent due to the enlarged contact surface between stratospheric and tropospheric air masses and the strong tracer gradients a high potential for subsequent mixing.

On this flight segment stratospheric NO$_y$ mixing ratios between and after the tropospheric filaments were much higher than values from the expected NO$_y$-O$_3$ correlation given by Murphy et al. (1993) or Weinheimer et al. (1993) with a slope dNO$_y$/dO$_3$ ~0.004. The correlation between NO$_y$ and O$_3$ is well established and supposed to show only little variation in the LMS besides a certain latitudinal dependence (Ziereis et al., 2000). The scatter plot of NO$_y$ versus O$_3$ (Fig. 3a) from the here presented measurements clearly shows the different relations between the two species in the different phases of the flight. The expected slope close to 0.004 was only observed later on the flight in segment III (see Sect. 5.2). In contrast to the here presented data, the NO$_y$/O$_3$ correlations obtained during two subsequent flights on the following day were well within the range of correlations found in former observations. In order to illustrate how unusual the observed NO$_y$ mixing ratios were during most of the time the NO$_y$ values calculated from the ozone data using the above correlation are indicated by the green line in Fig. 2a. Higher slopes of NO$_y$/O$_3$ correlations up to 0.01 were found by Bregman et al. (1995) during STREAM I and Fischer et al. (1997) during STREAM II and were attributed to renitrification which is unlikely to take place in November. Ray et al. (1999) however concluded from balloon measurements of different CFCs that the lowermost stratosphere in summer and autumn may be stronger influenced by isentropic transport from the tropical upper troposphere than in other seasons. Hoor et al. (2004) present in a more detailed study with broader latitudinal and seasonal coverage than the measurements of Ray et al. (1999) that isentropic transport indeed affects a distinct layer of Δθ=30 K above the tropopause in summer with decreasing tropospheric influence beyond this layer. Higher
NO$_y$ mixing ratios as obtained in the probed subtropical air masses during flight segment I therefore could result by isentropic mixing. Nevertheless, the mean NO$_y$ mixing ratios in the upper troposphere between 0.75 and 3 ppbv during summer and autumn (Fischer et al., 2000) may only partly attribute to the observed elevated NO$_y$ mixing ratios. Also, the measurements were taken in a $\Delta \Theta$ range between 35 K and 55 K above the tropopause and therefore we assume that the here presented measurements indicate a rather exceptional feature. It is important to note that the contamination problem mentioned in Sect. 2 can not be responsible for the unusually steep slope seen in flight segment I. The individual points on this line belong to the rapid transitions from stratospheric air into tropospheric air and back when the aircraft passed the flanks of the tropospheric filaments. These transitions occurred on a time scale of only 1–2 min which is much too short for significant changes in the background signal. A contamination effect still unaccounted for in the data would simply shift the correlation line up or down on the $y$-axis but would not affect its slope. The steep slope as well as the high absolute NO$_y$ mixing ratios at the upper end of the correlation line indicates that stratospheric air with unusually high NO$_y$ mixing ratios has mixed with tropospheric air. H$_2$O mixing ratios (Fig. 2c) were also higher than stratospheric mixing ratios of a few ppmv. However, these enhanced values are frequently observed in a mixing layer close to the troposphere at O$_3$ mixing ratios of 200–300 ppbv (Schiller et al., in preparation, 2004$^3$) and are further reproduced by the ECMWF model. The high NO$_y$ mixing ratios in the stratospheric air are most probably of tropospheric origin. Figure 5 shows that the majority of air parcels in the vicinity of the flight track in segment I has experienced troposphere-to-stratosphere transport between 6 and 10 days prior to the measurements (light to dark blue boxes). The tropospheric filaments described above, on the other hand, are significantly younger than this and are represented by the yellow to red boxes. The trajectory analysis thus suggests that the unusual tracer relation results from two subsequent mixing processes, the first one taking place between 1 and 4 November (or even before) and transporting tropospheric air with high NO$_y$ mixing ratios of about 4 to 5 ppbv into the stratosphere, and the second one taking place between 5 and 9 November transporting again tropospheric air into the stratosphere but this time with low NO$_y$ mixing ratios of less than 1 ppbv. It yet remains unclear what caused the high NO$_y$ mixing ratios in this first mixing process. Possible explanations remain that the tropospheric air masses introduced into the LMS during the first mixing process were impacted by NO$_y$ production by lightning or transport from polluted planetary boundary layer air to the upper troposphere through deep convection.

5.2 Flight-segment III

On the third flight-level, at the exit of the streamer, we observed a strong gradient in the measured trace gases though the aircraft remained at a constant pressure of 176 hPa and an almost constant potential temperature of about 365 K. NO$_y$ decreased within 10 min from 2.6 to 0.8 ppbv and at the same time O$_3$ decreased from about 600 to 100 ppbv. The correlation between the two tracers with a slope of 0.0034 is

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Fig. 5. Flight-track-following vertical cross section of the day an air parcel experienced TST. Air parcel trajectories were calculated 10 days backward from a fine 3-D grid and starting at three different times (06:00, 09:00 and 12:00 UTC) on 10 November as described in Sect. 3.2. The day when TST occurred was then projected onto the start points of the trajectories (i.e., onto the 3-D grid) and finally these values were interpolated in time and space onto vertical columns along the flight track. Only air parcels having a PV value greater than 3 PVU at the time of the flight may have experienced TST. The day of TST is defined as the last day the air parcels’ PV value was below 3 PVU. A value of 8, for example, indicates that TST occurred 8 days before the flight. The majority of air parcels has not experienced TST and is therefore left white. The lower border of the colored regions follows approximately the tropopause. Black circles represent the O\textsubscript{3} measurements, the radius being proportional to the mixing ratios.

5.3 Flight-segment II: standard analysis

While O\textsubscript{3} and NO\textsubscript{y} in the undisturbed stratosphere are expected to correlate positively with a slope between 0.003 and 0.006 (Murphy et al., 1993; Weinheimer et al., 1993), we observed a weak negative correlation in segment II (see Fig. 3a), suggesting that these air masses have been severely perturbed by air containing higher mixing ratios of NO\textsubscript{y} than expected. This is corroborated by the measurements of H\textsubscript{2}O which show remarkable differences from the mixing ratios normally found in the stratosphere. In contrast to flight segment I, H\textsubscript{2}O mixing ratios modeled by ECMWF are only between 5 and 8 ppmv as opposed to measured values between 20 and 30 ppmv (see Fig. 2c), which are among the highest H\textsubscript{2}O mixing ratios ever observed in air masses with O\textsubscript{3}>500 ppbv (Schiller et al., in preparation, 2004\textsuperscript{4}). This

results in a bulge in the O$_3$/H$_2$O relationship as shown in Fig. 3b which can not be simply explained by a mixing between air masses of segment I and III.

Figure 7 shows the different origin of the air parcel trajectories started from flight segment I (Fig. 7a) and II (Fig. 7b). The trajectories in segment II are clearly stratospheric with maximum PV values of 12 PVU and a minimum value of about 8 PVU (see Fig. 2d). They remained together during the previous 10 days following a path circulating about the north pole. In contrast, the trajectories of segment I originated in the subtropics and mid-latitudes with much lower PV maxima and minima. Also different from segment I, trajectories started from the RDF grid in the vicinity of the flight track show a clear stratospheric signature with no indication of TST during the previous 10 days (see Fig. 5). These findings suggest that in contrast to segment I the introduction of tropospheric air most likely did not happen through a process resolved by the ECMWF wind fields, at least not during the time period covered by the trajectories. However, the potential for transport and mixing at an earlier time, e.g. several months before sampling, is currently being studied with the 3-D version of the Chemical Lagrangian Model of the Stratosphere (CLaMS) (Günther et al., in preparation, 2004$^5$).

Denitrification and subsequent renitrification as a possible source for enhanced NO$_y$ mixing ratios (Koike et al., 2002) is not expected at this time of the year. Another possible explanation for both elevated NO$_y$ and H$_2$O mixing ratios is injection of tropospheric air through deep convection. The outflow of deep convective clouds may be strongly enhanced in NO$_x$ due to lightning (Brunner et al., 1998; Huntrieser et al., 1998; Lange et al., 2001). In Fig. 8 a Meteosat water vapor image is overlaid with the flight path (along orange circles) and with the 2 PVU contours at 200 and 350 hPa (dashed and solid blue line) marking the position of the streamer at two different altitudes. Widespread convection (bright areas) covers the Mediterranean with centers of activity located near the east coast of the Iberian peninsula and near Corsica, Sardinia and middle Italy. The area of convective activity is quite

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convective systems, we also analyzed the situation as simulated by the Climate High Resolution Model (CHRM). The cloud layers generated in the CHRM are highly consistent with the observed clouds over the Mediterranean. The corresponding convective activity has generated a lot of fine scale structure around the tropopause and has pushed the 2 PVU surface to about 270 hPa at 37° N (see Fig. 9b). This is significantly higher than in the ECMWF model but still well below flight altitude. Similar to the ECMWF model the CHRM shows no enhanced water vapor at flight level suggesting no connection between the observed elevated H₂O mixing ratios and the nearby convective activity. However, Fig. 8 only provides a picture of the convective activity for a single time, while numerous convective cells developed already during the previous day. The predictive capability of the CHRM for the location and particular timing of such an observed small-scale tracer anomaly is limited. This might be due to the particular model formulation for subgrid-scale or non-hydrostatic processes – and in fact, it is conceivable, that an element of both is present. For instance, processes associated with deep convection could be poorly represented by the parameterization, and subgrid-scale, non-hydrostatic or small-scale turbulent effects are inaccessible for investigation with the present model’s approximations.

5.4 Flight-segment II: convective influence analysis

As shown in the previous section, the analyses based on fields simulated by the ECMWF and CHRM are unable to explain the unusually high NOₓ and H₂O mixing ratios in flight segment II. In order to obtain an integral picture of the convective influence experienced by each air parcel of the RDF grid, we traced the convective activity observed by the Meteosat infrared channel along the backward trajectories over the course of the previous 1.5 days (from 9 November 00:00 UTC to 10 November 12:00 UTC). At each time step (30 min) the air parcel’s temperature was compared to the brightness temperature of the coinciding satellite pixel, and whenever the temperature was higher, the air parcel was said to have undergone convective influence, similar to the approach used by Jeker et al. (2000). Individual convective influences along a trajectory were then summed up, converted into an overall “convective influence time”, and projected forward onto the trajectory start points. Figure 10 shows the results of this analysis for backward trajectories started on the 200 hPa level (and therefore on flight level). In segment II the aircraft was closely following the northwestern edge of a region which experienced significant convective influence. Individual air parcels in this area remained up to 8 out of 36 h high inside or below high (cold) clouds. Closer inspection of the paths of these air parcels reveals that they have crossed areas of intense lightning activity over northern and southern Spain and over the north coast of Algeria (see Fig. 11). The encounter time corresponds to the time of the most vigorous lightning activity on the evening of 9 November (dark blue to bright blue spots in Fig. 11).
A comparison between ECMWF winds interpolated onto the flight track and the wind measured aboard the aircraft reveals notable differences for the time between 10:00 and 11:00 UTC (see Fig. 12). The ECMWF analysis shows northeasterly winds (between 55–65°) whereas the measurements show a much stronger easterly component (70–90°). The significant differences between measured and ECMWF analyzed wind directions, which was seen only in segment II, is probably due to the fact that the aircraft was flying close to a shear line marking the center of the streamer at 200 hPa. Across this shear line the wind changed dramatically from northern to southern directions and therefore small errors in the positioning of this line would result in large errors in the predicted wind direction. It appears that part of the outflow of the convective activity over the Mediterranean was directed more strongly towards the west than simulated by the ECMWF model, resulting in a more eastern wind component on the flight track than predicted by the model.

The above analyses suggest that convective injection of tropospheric air, enhanced in NO through lightning activity, may indeed serve as an explanation for the elevated H$_2$O and NO$_x$ mixing ratios. Most of the troposphere-to-stratosphere transport appears to have occurred more than 12 h earlier in convective systems over northern and southern Spain and over Algeria. Turbulent mixing close in the outflow to the deep convective cells may have smeared out individual plumes in the following hours resulting in a rather uniform enhancement in the tracers as observed from the aircraft.

In order to estimate the amount of NO$_x$ and H$_2$O initially contained in the injected tropospheric air, we performed a simple mass balance calculation following Ray et al. (1999). The measured CO mixing ratios along segment II (ranging from 25 to 35 ppbv) were used to quantify the fraction of tropospheric air within the probed air masses, based...
on estimates of CO mixing ratios in the undisturbed stratosphere (15–20 ppbv) (Hoor et al., 2002) and in the troposphere (70 ppbv as measured during a second flight on the same day in tropospheric and subtropical air masses or as taken from a classification by Fischer et al., 2002). From this we inferred that between 5 and 20% tropospheric air has been mixed into the lowermost stratosphere and that this tropospheric air originally contained between 6 and 10 ppbv NO$_y$ and between 100 and 400 ppmv H$_2$O. The mixing ratios for NO$_y$ in the tropospheric air calculated in this way are rather high, but previous studies already reported on mixing ratios around 3 ppbv NO$_y$ (Lange et al., 2001) or even up to 5 ppbv NO$_x$ (Brunner et al., 1998) measured in regions of convective outflow. The air masses probably influenced by the lightning and convective activity over northern and southern Spain and Algeria end up in that part of the colored area in Fig. 10 which lies to the west of 5$^\circ$ E. This area has a size of approximately 3.5 $\times$ 10$^5$ km$^2$ and if we assume that the convective outflow occurred in a layer of about 3 km thickness we can calculate a total volume of the outflow air of 1.05 $\times$ 10$^6$ km$^3$. The total number of lightning strokes registered by the Arrival Time Detection System of the UK Meteorological Office between 9 November 00:00 UTC and 10 November 12:00 UTC in the convective systems over the considered region sums up to about 11 200. Assuming a NO production of 4 $\times$ 10$^{26}$ molecules per flash (Drapcho et al., 1983) and assuming that up to 75% of the produced NO$_x$ ends up in the upper troposphere above 8 km (Pickering et al., 1998) we can estimate an average NO$_y$ mixing ratio of 0.7 ppbv due to lightning activity alone. Part of this NO$_y$-rich air is eventually mixed into the LMS and increased the NO$_y$ mixing ratios along the flight path. However, the 0.7 ppbv are roughly one order of magnitude less than the 6 to 10 ppbv required to explain the observations and we can only speculate on possible reasons for this. Either the assumption of 4 $\times$ 10$^{26}$ molecules per lightning flash is much too low, or the assumptions on the total volume of convective outflow is overestimated, or other sources unaccounted for in our estimates such as aircraft emissions must have contributed substantially. Note that correlations with CO and CO$_2$ do not indicate a significant influence on NO$_y$ through transport of polluted planetary boundary air. However, elevated tropospheric NO$_2$ columns in the area of the flight track on 10 November have also been observed from space by the GOME instrument and were attributed to the lightning activity over the Mediterranean (Thomas et al., 2003). Thomas et al. (2003) calculated a maximum tropospheric column content of 5.5 $\times$ 10$^{15}$ molecules cm$^{-2}$ on 11 November and attributed the enhancement to a layer located above cloud levels. Assuming a column-height between clouds and tropopause of 3 km a NO$_2$ mixing ratio of 1.5 ppbv results. With a typical ratio NO/NO$_2$ in photochemical equilibrium...
of 4:1 the expected NO mixing ratio would be around 6 ppbv and would thus be able to explain our findings.

Finally, we calculated 10-day forward trajectories to estimate the stratospheric residence time of this strongly affected air mass in order to judge whether the perturbation to the LMS was long-lasting and hence significant following the arguments for detecting significant TST events presented by Wernli and Bourqui (2002). The results show, that 97% of the air parcels, which started on the 200 hPa-level and experienced strong convective influence as seen in Fig. 10, remain in the lowermost stratosphere and therefore may have a considerable impact on the chemical and radiative properties of this region.

6 Conclusions

We have reported on airborne measurements of NOy and other tracers in the lowermost stratosphere during an extraordinary meteorological episode featuring a deep stratospheric intrusion over south-western Europe.

At the south-western edge of the streamer, a strong gradient in the NOy and the O3 mixing ratios has been encountered while flying on a constant pressure (176 hPa) and on an almost constant potential temperature level (365 K). In agreement with the tracer observations the ECMWF analysis shows a strong gradient in PV along this flight segment demonstrating how well the model located the position of this part of the streamer.

Sudden drops in NOy and O3 at the western flank of the streamer could be well understood by reconstruction of the fine scale structures in potential vorticity using the RDF technique, which showed the presence of a sequence of three narrow tropospheric filaments embedded into the stratospheric streamer. The steep tracer gradients at the borders of these filaments indicate that only little mixing with the surrounding stratospheric air had occurred at that stage. Mixing is most likely taking place later on as the tropospheric air masses are stretched into ever longer and narrower filaments. The air with stratospheric characteristics located between the filaments showed strongly enhanced NOy mixing ratios, which probably was introduced by TST about 10 days ago, long before the formation of the tropospheric filaments occurred. However, given the uncertainty in trajectory calculations over 10 days backward we were not able to find a fully conclusive explanation for these elevated NOy mixing ratios.

During the second flight segment in the interior of the streamer, however, strongly enhanced mixing ratios of NOy by up to 100% of the expected values are shown to be most probably caused by recent mixing with tropospheric air through injection by deep convection on the forward (eastern) flank of the streamer. The data are corroborated by observations of H2O showing enhancements of up to 200%. Neither the ECMWF model nor the mesoscale CHRM are able to reproduce the observed enhancement in H2O in the lowermost stratosphere at the altitude of the aircraft. Convective activity in the two models extends only to about 340 hPa and 270 hPa, respectively, well below flight altitude of about 200 hPa. The impact of the convective system with its continuously developing cells during the previous two days was potentially large and the outflow region stretched along the flight path over a distance of about 700 km.

We traced convective influence along backward trajectories started from a large area surrounding the flight track at 200 hPa using the observations of cold (high) clouds observed by Meteosat at 30 min time intervals during the previous 1.5 days. As a measure for the convective influence we used the total amount of time an air parcel’s temperature traced along its back trajectory was higher than the brightness temperature of the corresponding satellite pixels. The results show that the aircraft was flying along the border of a region with strong convective influence confirming our hypothesis of mixing with tropospheric air. The fraction of the introduced tropospheric air has been calculated by a mass balance calculation based on the CO measurements to lie between 5 and 20% and the expected NOy mixing ratios in the tropospheric air between about 6 and 10 ppbv. As the CO mixing ratios were hardly enhanced, the source of the elevated NOy is assumed to be production by lightning. In fact, high lightning activity has been registered in the convective cells over the western Mediterranean. However, the observed enhancement in NOy is approximately one order of magnitude higher than expected from a rough estimate based on the total number of lightning strokes, the total mass of air affected by the lightning, and typical values for NO-production per lightning stroke taken from the literature. This reflects the high uncertainty range in the made assumptions as well as in the used NO-production rate per lightning stroke.

An analysis of forward trajectories started from an area inside the stratospheric streamer for which a strong convective influence was calculated shows that 97% of these air parcels remain in the lowermost stratosphere during the following 10 days. This suggests that the convective injection of air with elevated mixing ratios of NOy and H2O may have had a considerable impact on the chemistry and radiative properties of this region. These findings support recent model studies by Wang (2003) and Mullendore (personal communication, 2003) showing that the breaking of gravity waves at the top of deep convective systems in mid-latitudes can cause injection of water vapor and other tropospheric tracers into the lowermost stratosphere in the form of plumes well above the thunderstorm anvils. Airborne measurement campaigns with high reaching aircraft especially designed to investigate the environment well above the anvils of the convective systems will be needed to validate the impact on tracer distributions predicted by these models.
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