



HAL
open science

Out of Africa: High aerosol concentrations in the upper troposphere over Africa

J. Heintzenberg, M. Hermann, D. Theiss

► **To cite this version:**

J. Heintzenberg, M. Hermann, D. Theiss. Out of Africa: High aerosol concentrations in the upper troposphere over Africa. *Atmospheric Chemistry and Physics Discussions*, 2003, 3 (3), pp.2661-2689. hal-00301073

HAL Id: hal-00301073

<https://hal.science/hal-00301073>

Submitted on 18 Jun 2008

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

**High aerosol
concentrations in the
upper troposphere
over Africa**

J. Heintzenberg et al.

Out of Africa: High aerosol concentrations in the upper troposphere over Africa

J. Heintzenberg, M. Hermann, and D. Theiss

Leibniz-Institute for Tropospheric Research, Permoserstr. 15, 04318 Leipzig, Germany

Received: 22 February 2003 – Accepted: 7 May 2003 – Published: 19 May 2003

Correspondence to: J. Heintzenberg (jost@mail.tropos.de)

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

⏪

⏩

◀

▶

Back

Close

Full Screen / Esc

Print Version

Interactive Discussion

Abstract

In the year 2000, six flights (three southbound and three northbound) of the CARIBIC project were conducted between Germany and two destinations in the southern hemisphere (Windhoek, Namibia and Cape Town, South Africa). In the present report, results on particle number concentrations are discussed in three size ranges (> 4 nm, > 12 nm, and > 18 nm particle diameter) during the unique transequatorial Africa flights. The flights covered a total of about 80 h in May, July, and December. Thus, no claim can be made for long-term representativeness of the aerosol data. Nevertheless, they are the first upper systematic tropospheric transequatorial aerosol profiles over Africa. The average aerosol results show a broad maximum, roughly symmetrical to the equator, which compares well in latitudinal extent to a maximum of CO concentrations measured on the same flights. This export of continental surface aerosol to the upper troposphere will be dispersed on a global scale both with the easterly flow near the equator and with the westerlies in the adjacent subtropical regions. There was strong evidence of recent new particle formation before aerosol arrival at flight level, in particular during the time periods between 9:00 and 13:00 local time over Africa. Direct and indirect climate effects of the respective particulate matter remain to be investigated by future flights with the ongoing extension of the CARIBIC payload towards size-resolved measurements above 100 nm particle diameter. At the same time global chemical transport models and aerosol dynamics models need to be extended to be able to reproduce the CARIBIC findings over Africa.

1. Introduction

Aerosol forcing of climate is an important issue in the discussion of climate change (Charlson and Heintzenberg, 1995). Despite its importance, the influence of aerosol particles on the Earth's radiation budget is still highly uncertain (Houghton et al., 2001). In particular, particles in the upper troposphere (UT) and lower stratosphere (LS) play

High aerosol concentrations in the upper troposphere over Africa

J. Heintzenberg et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

⏪

⏩

◀

▶

Back

Close

Full Screen / Esc

Print Version

Interactive Discussion

an important role in heterogeneous chemistry and radiative forcing, e.g. by cirrus clouds (Laaksonen et al., 2000), and there is some evidence of their affecting tropical cumulus clouds (Sherwood, 2002). However, the knowledge on UT/LS aerosol particles is limited today (Raes et al., 2000).

5 The CARIBIC research project (Civil Aircraft for Regular Investigation of the atmosphere Based on an Instrument Container, Brenninkmeijer et al., 1999) was designed to increase the very limited knowledge of trace gases and aerosol particles in the upper troposphere and lower stratosphere. Beginning in 1997 on long-range commercial flights continuous measurements of particle number concentrations in three submicrometer size ranges have been taken as part of this project. Initially, these flights
10 operated between Germany and the Indic. The southernmost latitudes of these flights reached about 5° N. Until the year 2000, 45 measurement flights had been conducted on that route. A statistical discussion of the particle number concentrations presented in Hermann et al. (2003) showed that in the tropics high number concentrations of
15 ultrafine particles smaller than about 10 nm in diameter frequently were encountered at flight altitudes of about ten kilometers. Trajectory analyses and trace gas information strongly indicated that the some of the related air masses originated from low altitudes over Africa.

20 In 2000, six flights (three outbound and three homebound) were conducted between Germany and two destinations in the southern hemisphere (Windhoek, Namibia and Cape Town, South Africa). In May 2002, the CARIBIC aircraft has been decommissioned and the payload is being rebuilt for deployment on another aircraft starting in late 2003. In the present report, the results on particle number concentrations during the unique transequatorial Africa flights are discussed.

25 2. Instrumental

Particle number concentrations were measured with a combination of three Condensation Particle Counters (CPCs, TSI Model 7610, St. Paul, MN) operated in paral-

High aerosol concentrations in the upper troposphere over Africa

J. Heintzenberg et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

⏪

⏩

◀

▶

Back

Close

Full Screen / Esc

Print Version

Interactive Discussion

High aerosol concentrations in the upper troposphere over Africa

J. Heintzenberg et al.

[Title Page](#)[Abstract](#)[Introduction](#)[Conclusions](#)[References](#)[Tables](#)[Figures](#)[◀](#)[▶](#)[◀](#)[▶](#)[Back](#)[Close](#)[Full Screen / Esc](#)[Print Version](#)[Interactive Discussion](#)

lel. These instruments were modified for low-pressure conditions and aviation requirements, and were extensively tested with respect to their counting efficiency at operating pressures of 160–1000 hPa (Hermann and Wiedensohler, 2001). The lower threshold diameters of the CPCs (i.e. the particle diameter with 50% detection efficiency) were set to 4 nm (N_4), 12 nm (N_{12}), and 18 nm (N_{18}), respectively, by increasing or decreasing the temperature difference between the saturator and condenser block inside the counters. The upper threshold diameter was determined by the aerosol inlet system to 1.3 μm (50% sampling efficiency), which had been characterized in detail in terms of aspiration efficiency and transmission by Hermann (2000). Subtracting the readings of the first two counters yields the number concentration of ultrafine particles (N_{4-12}).

Number concentrations were recorded with a time resolution of two seconds on flight levels corresponding to pressures below 500 hPa. As all single-particle counters, the CARIBIC CPCs suffer from coincidence errors due to finite probabilities of more than one particle being present concurrently in the sensing volume. Up to true concentrations of 40 000 cm^{-3} (at working pressure) this error was corrected for after an empirical coincidence calibration (Hermann and Wiedensohler, 2001). The coincidence-corrected counting results are reduced to STP-conditions (1013.25 hPa, 273.15 K). Consequently, the upper limit of acceptable CPC data increases to about 120 000 cm^{-3} (STP) and was set to 100 000 cm^{-3} (STP) in all results presented below.

Besides correction for particle coincidence, number concentrations were corrected for losses in the inlet and in the sampling line ($\approx 27\%$ for N_{4-12} and 10% for N_{12} and N_{18}), for pressure-dependent CPC flow rates, and CPC counting efficiencies (Hermann and Wiedensohler, 2001). Together, these corrections account for a particle-size and number-concentration-dependent increase in raw particle concentrations by a factor of 1.2 to 2.

For data analysis, the CARIBIC system provided additional information on ozone and carbon monoxide mixing ratios along the flight track (cf. Brenninkmeijer et al. 1999). For each CARIBIC flight, aircraft position and altitude information were used together with ECMWF data to calculate three-dimensional 5-day back trajectories every

five minutes along the CARIBIC flight paths. This work was done by the Royal Dutch Meteorological Service, KNMI, de Bilt, Netherlands.

3. Results and discussion

3.1. Overview

5 General parameters of the six Africa flights are listed in Table 1. Figure 1 shows the two transequatorial routes taken on these flights. Because of different African flight destinations, they differ mainly over southern Africa whereas the rest of Africa was crossed on very similar routes on each flight. Small differences are due to larger tropical convective systems that were circumvented by the pilots. Two sources of meteorological information were used to delineate the extent of the Intertropical Convergence Zone (ITCZ) along the CARIBIC flight tracks. For the planetary boundary layer the 700 hPa, potential temperatures from ECWMF fields were used. Strong temperature gradients in the climatologically most probable regions were taken as delineators. ECWMF high cloud fields and METEOSAT images in the visible channel were used as ITCZ delin-

10 eators in flight level. As expected, the ITCZ divided into two branches diverging in easterly direction over the African continent. The estimated northern and southern limits of the ITCZ are also listed in Table 1.

In Fig. 2a, N_{12} for all flights is drawn as a function of latitude in order to visualize similarities and differences between the six flights. As on all other CARIBIC flights the European region stands out with short periods of high concentrations that can be explained by high air traffic densities with ensuing emissions of exhaust gases and particles. During the day-flight in boreal summer, (cf. subplot 000728 in Fig. 2a), convective activity over Europe added to the upper tropospheric aerosol burden.

20 On most flights, northern subtropical latitudes are characterized by low and rather stable particle concentrations, confirming the findings of Hermann et al. (2003) for CARIBIC flights to the Indic. Except for the last two flights, the same result holds for

High aerosol concentrations in the upper troposphere over Africa

J. Heintzenberg et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Print Version

Interactive Discussion

High aerosol concentrations in the upper troposphere over AfricaJ. Heintzenberg et al.

[Title Page](#)[Abstract](#)[Introduction](#)[Conclusions](#)[References](#)[Tables](#)[Figures](#)[◀](#)[▶](#)[◀](#)[▶](#)[Back](#)[Close](#)[Full Screen / Esc](#)[Print Version](#)[Interactive Discussion](#)

southern subtropical latitudes as well. Very different results were found in the tropics. In boreal spring and summer there is a broad latitudinal region of roughly $\pm 15^\circ$ around the equator in which evening and night flights (000518, 000727, 001202) yielded N_{12} values, which were elevated by a factor of 3–5 compared to the remainder of the flights.

5 In austral summer, (cf. subplot 001202 in Fig. 2a), this region is shifted to the south and exhibits a gap near the equator. The region of elevated particle concentrations roughly compares with the extent of the ITCZ along the flight tracks (cf. Table 1). The dip in average particle concentrations close to the equator can be explained by the bifurcation in the ITCZ over the African continent. According to the satellite images in
10 the visible this bifurcation became more pronounced towards the end of the series of flights (cf. flights 001202/03). The three day-flights (000519, 000728, 001203) likewise show the broad maximum in the particle number concentration. However, there are high concentration peaks superposed, exceeding $100\,000$ particles cm^{-3} STP. These are the highest concentrations ever measured during CARIBIC. The general shape of
15 the increase in particle concentrations over the African continent is quite similar within each of the three pairs of flights, each of which covered a total period of about 24 h.

Average ratios of northern to southern hemisphere concentrations can be formed by combining the data of the northern subtropics with the rest of the respective flights data north of these latitudes (cf. Table 2). Averaging over all flights yields concentration
20 ratios of northern-to-southern hemisphere data of 1.8, 1.4, and 1.3 in the size channels N_4 , N_{12} , and N_{18} , respectively. These values are smaller as similar hemispheric ratios given in Minikin et al. (2003) for the INCA project who report values between two and three. The CARIBIC data, however, only reach down to 33° S, whereas the INCA data reach 53° S, Furthermore, the present data set does not encompass data
25 from the heavily trafficked north Atlantic flight corridor, as did the INCA data. We also note that on all of the first four flights a decrease in N_{12} was encountered between the southernmost end of the flights and the onset of the concentration bulge near the equator (cf. Fig. 2a). The last two flights, on the other hand, did not exhibit this feature.

3.2. Ultrafine particles over tropical Africa

During flights, which passed over tropical Africa around local noon, much higher ratios of tropical to subtropical number concentrations were measured, reaching average values up to ten for the ratio of N_4 (tropical to northern subtropics, cf. Table 2). Often, extremely high and extremely variable particle concentrations were recorded. According to the coincidence limits of the CPCs (cf. Sect. 2), values larger than $120\,000\text{ cm}^{-3}$ (STP) were excluded from the latitudinal averages presented in Table 2 and from the graphs.

The difference between day flights and night flights becomes much more obvious in the difference N_{4-12} , also termed ultrafine particles (UFP), as plotted in Fig. 2b in the same way as N_{12} is shown in Fig. 2a. This difference emphasizes the smallest particles, which most recently have been formed near the flight path or have been transported up rapidly from their source region in the lower troposphere. We note that N_{4-12} may become insignificant or even negative for short periods, at low values of N_{4-12} because of statistical fluctuations in the two counters. Therefore, only values of N_{4-12} are shown that are larger than the Poisson counting error of N_4 .

N_{4-12} of the day flights is presented as a function of local time in Fig. 3 in order to emphasize their connection with the sunlit part of the day. Strong maxima on all three daytime flights appeared between 09:30 and 12:30 LT, the period during which many continental UFP studies reported new particle formation due to mixing and/or photochemical activity in the planetary boundary layer (e.g. Birmili et al., 2001). Low particle concentrations occurred in between, e.g. 10:30 and 11:30 LT (cf. Fig. 3). The corresponding trajectories (not shown) indicated upper tropospheric and lower stratospheric air and the satellite images indicate a cloud-free region in the bifurcated continental ITCZ.

If these high concentrations were related to recent particle nucleation events it can be expected that these events should be strongest with a minimum of preexisting particulate surface that could compete for the condensable vapors that otherwise nucleate.

High aerosol concentrations in the upper troposphere over Africa

J. Heintzenberg et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

⏪

⏩

◀

▶

Back

Close

Full Screen / Esc

Print Version

Interactive Discussion

High aerosol concentrations in the upper troposphere over AfricaJ. Heintzenberg et al.

[Title Page](#)[Abstract](#)[Introduction](#)[Conclusions](#)[References](#)[Tables](#)[Figures](#)[⏪](#)[⏩](#)[◀](#)[▶](#)[Back](#)[Close](#)[Full Screen / Esc](#)[Print Version](#)[Interactive Discussion](#)

To date, no particle surface-related parameter is measured in the CARIBIC payload. As a compromise, the largest particle channel N_{18} was related to N_{4-12} in Fig. 4 for the three daytime flights. Consistently on all three day-flights, there is a general trend towards highest values of N_{4-12} with lowest N_{18} in accord with atmospheric nucleation concepts.

From the available information it is difficult to assess where the measured UFPs had been produced. From local ozone concentrations, the origin of the air masses can be gleaned. During the periods of the strong local maxima of N_{4-12} , mostly values below 80 ppb were recorded, during the December flight often below 50 ppb (cf. Fig. 5). Average values for the period 09:30 and 12:30 LT were 67, 63, and 49 ppb for the flights 000519, 000728, and 001203, respectively.

Low ozone values are typical for tropical surface air. Over the tropical Atlantic (20° S–30° N, 1977–1996), average surface values for the period 1977–1996 as measured by the German Weather Service (P. Winkler, private communication), were 33, 31, and 21 ppb for the months May, July, and December, respectively. For the Mt. Kenya GAW station the corresponding average values for the period 1999–2002 were 26, 19, 16 ppb. Whereas average absolute ozone values at the surface apparently were somewhat lower than during the UFP-events on the CARIBIC flights, the seasonal trend is in qualitative agreement, with the lowest values during the austral summer and the highest values in May.

For the flight 000728 Mühle et al. (2002) have evaluated the trace gas analyses together with the particle data. Their CO enhancement ratios over tropical Africa for non-methane hydrocarbons and methyl halides are consistent with those reported from fresh biomass burning plumes, thus corroborating the results of the present study.

3.3. The origin of sampled air masses

The CARIBIC trajectories were explored for their potential information content concerning convective transport of polluted continental surface air to the flight paths. We focused our attention on the latitude band $\pm 25^\circ$ about the equator which covers both

High aerosol concentrations in the upper troposphere over AfricaJ. Heintzenberg et al.

[Title Page](#)[Abstract](#)[Introduction](#)[Conclusions](#)[References](#)[Tables](#)[Figures](#)[◀](#)[▶](#)[◀](#)[▶](#)[Back](#)[Close](#)[Full Screen / Esc](#)[Print Version](#)[Interactive Discussion](#)

major areas of biomass burning and strong convective activity (Herman et al., 1997; Sherwood, 2002). During periods of high number concentrations only few trajectories reached down into the African planetary boundary layer allowing to explain periods with high particle number concentrations. This limitation is due to the fact that the meteorological models on which the trajectory calculations are based are unable to resolve the processes and scales which control rapid vertical mixing of polluted surface air to the flight level. As a simplistic first indirect approach, the horizontal trajectory positions were merged with a GISS-based map of the national boundaries of African states and the lengths of the trajectories over each country relative to the total territorial boundary of each country was calculated. These relative lengths were summed over all African states within $\pm 25^\circ$ latitude about the equator and then correlated with the aerosol information after interpolating the GISS results linearly in between the five-minute intervals of trajectory information with the timing information of the CPC data. For individual flights correlation coefficients up to 0.63 were reached (flight 000727). However, the average correlation coefficient over all six flights was only 0.38 with a minimum of -0.01 for flight 000728.

In two ways this simplistic trajectory evaluation was augmented. First the three-dimensional information was exploited by weighing the land passages of the trajectories with the pressure changes along the trajectories by the following procedure. A cubic spline approximation developed by Reinsch (1967) was used for this purpose. The pressure differences were calculated as first derivative of the spline-interpolated trajectory at the center of each interval of three one-hour time steps of a trajectory. A factor was determined to normalize this derivative and transform it into a range of values between zero on one. The normalization was by difference of estimated pressure at this point to a fixed maximum pressure. This weighing is based on the fact that large-scale subsidence as indicated by corresponding pressure changes suppresses convective upward transport of surface air and that the influence of surface air increases at trajectory levels near the surface. Additionally, the relative lengths of land passages were weighted with the sum of GEIA emission inventories for black carbon emissions

from biomass and fossil fuel burning over Africa (<http://weather.engin.umich.edu/geia/>). Whereas some improvements of trajectory/CPC correlations in subsidence regions south of the ITCZ and close to the equator (within the divided ITCZ over the African continent), the overall correlation coefficients did not improve in comparison with those from the unweighted relative land passages. Hence the refined trajectory approach did not provide more conclusive information on the continental origin of the particles.

3.4. The question of artifacts

All flights crossed the ITZC that is characterized by strong convective activity with cumulus towers reaching beyond the average ten-kilometer flight altitudes. There are many reports of artifact signals created at aerosol inlets on aircraft when operating in clouds (e.g. Clarke et al., 1997; Hudson, 1993; Hudson and Frisbie, 1991; Weber et al., 1998). Cloud drops shattering and splashing at the edges of the inlets are supposed to create large numbers of fragments that are aspirated, dried, and recorded as small residues in aerosol sensors. The presumed processes include the formation of water films on the inlets that tear off at downstream edges of the inlet, again creating fragment drops that cause artifact aerosol signals.

To date the CARIBIC payload does not comprise any cloud sensor. Thus, there are no direct measurements available to decide if in-cloud artifact formation occurred. In the following, we are collecting arguments supporting our assumption that the present aerosol data were not affected significantly by cloud artifacts.

Guided by on-board radar commercial aircraft specifically avoid large cumulus towers, trying to pass the area 100–200 km upwind of the core by changing course some ten minutes ahead of the cloud passage. Therefore, cloud encounters can be expected to be infrequent.

Due to the nature of the artifact processes, the distribution of drop fragment sizes and numbers is expected to be chaotic in nature. Consequently, the size distribution of residue aerosol particles should be chaotic in nature as well. As a first test of the regularity of the number size distributions in the region of potential cloud passages the

High aerosol concentrations in the upper troposphere over Africa

J. Heintzenberg et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

⏪

⏩

◀

▶

Back

Close

Full Screen / Esc

Print Version

Interactive Discussion

High aerosol concentrations in the upper troposphere over Africa

J. Heintzenberg et al.

[Title Page](#)[Abstract](#)[Introduction](#)[Conclusions](#)[References](#)[Tables](#)[Figures](#)[⏪](#)[⏩](#)[◀](#)[▶](#)[Back](#)[Close](#)[Full Screen / Esc](#)[Print Version](#)[Interactive Discussion](#)

shape of the measured size distributions was checked by comparing ratios of concentration increases in the three size channels. As a sudden concentration step, a factor of two increase within two seconds in N_4 relative to the average concentration of the past ten seconds was defined. Whenever this occurred, the respective increases in N_{12} and N_{18} were calculated. In only 22 of 92497 recordings did the concentration ratios as defined above increase monotonically from N_4 to N_{18} . This very small number increased to 63 when only comparing N_{12} to N_4 , i.e. only in 63 cases did N_{12} increase more strongly than N_4 according to the above definition. In the vast majority of cases concentration increases were characterized by the smallest particles increasing most strongly, which is typical for recently nucleated atmospheric aerosols.

To our knowledge, there are no reports of artifact aerosol signals in ice clouds. On the opposite, even reduced interstitial aerosol signals have been reported in cirrus (Ström et al., 1999). Arguments have been given for the fragmentation processes being less important in these clouds (Weber et al., 1998). We took this as the starting point of a second test for potential artifacts in our data. Air temperatures of -40°C are usually taken as a limit between water clouds and ice clouds. For the flight starting 001203 the total number of data points with temperatures below -40°C was roughly equal to that above -40°C . We used an FFT analysis in order to compare the frequency (f) distributions of the two populations (three size channels each). Over the range of 0.001 to 0.2 Hz the data smoothly followed an $f^{-5/3}$ power law, indicating a well-mixed underlying atmosphere without indications of chaotic processes as would have been expected for chaotic cloud drop fragmentation. Furthermore, the frequency distributions are very similar in all three size-channels, corroborating the results of the first test.

4. Conclusions

The CARIBIC flights over Africa covered a total of about 80 h in May, July, and December 2000. Thus, no claim can be made for long-term representativeness of these

High aerosol concentrations in the upper troposphere over AfricaJ. Heintzenberg et al.

[Title Page](#)[Abstract](#)[Introduction](#)[Conclusions](#)[References](#)[Tables](#)[Figures](#)[◀](#)[▶](#)[◀](#)[▶](#)[Back](#)[Close](#)[Full Screen / Esc](#)[Print Version](#)[Interactive Discussion](#)

particle data. Nevertheless, they are the first systematic upper tropospheric transequatorial aerosol profiles over Africa, which global chemical transport models need for their validation. Thus, we plotted average meridional distributions of the three aerosol channels in Fig. 6 in 2.5° latitudinal bins to demonstrate the effect of the African continent on the upper tropospheric submicrometer aerosol. The average particle results show a broad maximum, roughly symmetrical to the equator, which compares well in latitudinal extent to a maximum of CO concentrations measured on the same flights. Ozone, on average, on the other hand does not show a corresponding minimum around the equator.

This export of continental surface aerosol to the upper troposphere will be dispersed on a global scale both with the easterly flow near the equator and with the westerlies in the adjacent subtropical regions. Direct and indirect climate effects of the respective particulate matter remain to be investigated on future flights with the ongoing extension of the CARIBIC payload towards size-resolved measurements above 100 nm particle diameter. At the same time global chemical transport models and aerosol dynamics models need to be extended to be able to reproduce the CARIBIC findings over Africa.

Acknowledgement. Part of this work has been funded by the European Commission through contract number ENV-CT95-0006 and the Ministry for Research and Education of the Federal Republic of Germany under project number 07ATF17 (CARIBIC) within the AFO2000 research program. We gratefully acknowledge the support by the carrier of the CARIBIC payload and industry partner of the project, LTU International Airways. We thank P. van Velthoven and H. Cuijpers, KNMI, de Bilt, Netherlands for calculating backward trajectories, and EUMETSAT (Darmstadt, Germany) for providing the Meteosat-5 satellite images. Furthermore, we thank the German Weather Service, represented by P. Winkler, and the Kenya Meteorological Department, represented by J. K. Rotich for their respective ozone data. H. Siebert of IfT kindly did the FFT analysis.

References

- Charlson, R. J. and Heintzenberg, J.: Aerosol Forcing of Climate (Ed.), John Wiley & Sons, New York, 416, 1995.
- 5 Birmili, W., Wiedensohler, A., Heintzenberg, J., and Lehmann, K.: Atmospheric particle number size distribution in Central Europe: Statistical relations to air masses and meteorology, J. Geophys. Res., 106, 32 005–32 018, 2001.
- 10 Brenninkmeijer, C. A. M., Crutzen, P. J., Fischer, H., Güsten, H., Hans, W., Heinrich, G., Heintzenberg, J., Hermann, M., Immelmann, T., Kersting, D., Maiss, M., Nolle, M., Pitscheider, A., Pohlkamp, H., Scharffe, D., Specht, K., and Wiedensohler, A.: CARIBIC – civil aircraft for global measurements of trace gases and aerosols in the tropopause region, J. Atmos. Oceanic Technol., 16, 1373-1383, 1999.
- Clarke, A. D., Uehara, T., and Porter, J. N.: Atmospheric nuclei and related aerosol fields over the Atlantic: Clean subsiding air and continental pollution during ASTEX, J. Geophys. Res., 102, 25 281–25 292, 1997.
- 15 Herman, J. R., Herman, Barthia, P. K., Torres, O., Hsu, C., Seftor, C., and Celarier, E.: Global distribution of UV-absorbing aerosols from Nimbus 7/TOMS data, J. Geophys. Res., 102: 16 911-16 922, 1997.
- Hermann, M.: Development and application of an aerosol measurement system for use on commercial aircraft, Ph.D. Thesis, University of Leipzig, Leipzig, 114, 2000.
- 20 Hermann, M., Heintzenberg, J., Wiedensohler, A., Brenninkmeijer, C. A. M., Heinrich, G., and Zahn, A.: Meridional distributions of aerosol particle number concentrations in the upper troposphere and lower stratosphere obtained by CARIBIC flights, J. Geophys. Res., 108, 4114, doi:10.1029/2001JD0001077, 2003.
- Hermann, M. and Wiedensohler, A.: Counting efficiency of condensation particle counters at low-pressures with illustrative data from the upper troposphere, J. Aerosol Sci., 32, 975–991, 2001.
- 25 Houghton, J. T., Ding, Y., Griggs, D. J., Noguer, M., Linden, P. J. v. d., and Xiaosu, D.: Climate Change 2001: The Scientific Basis, J. T. Houghton (Ed.), Cambridge University Press, Cambridge UK: 896, 2001.
- 30 Hudson, J. G.: Cloud condensation nuclei near marine cumulus, J. Geophys. Res. 98, 2693–2702, 1993.
- Hudson, J. G. and Frisbie, P. R.: Cloud condensation nuclei near marine stratus, J. Geophys.

High aerosol concentrations in the upper troposphere over Africa

J. Heintzenberg et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Print Version

Interactive Discussion

High aerosol concentrations in the upper troposphere over AfricaJ. Heintzenberg et al.

[Title Page](#)[Abstract](#)[Introduction](#)[Conclusions](#)[References](#)[Tables](#)[Figures](#)[◀](#)[▶](#)[◀](#)[▶](#)[Back](#)[Close](#)[Full Screen / Esc](#)[Print Version](#)[Interactive Discussion](#)

Res., 96, 20 795–20 808, 1991.

Laaksonen, A., Pirjola, L., Kulmala, M., Wohlfrom, K.-H., Arnold, F., and Raes, F.: Upper tropospheric SO₂ conversion into sulfuric acid aerosols and cloud condensation nuclei, *J. Glaciol.* 105, 1459–1469, 2000.

5 Minikin, A., Petzold, A., Ström, J., Krejci, R., Seifert, M., Velthoven, P. v., Schlager, H., and Schumann, U.: Interhemispheric differences in the upper tropospheric fine particle aerosol at midlatitudes, *Geophys. Res. Lett.*, (in press), 2003.

Mühle, J., Brenninkmeijer, C. A. M., Rhee, T. S., Slemr, F., Oram, D. E., Penkett, S. A., and Zahn, A.: Biomass burning and fossil fuel signatures in the upper troposphere observed during a CARIBIC flight from Namibia to Germany, *Geophys. Res. Lett.*, 29, 19 16-1–16-4, 2002.

10 Raes, F., Van Dingenen, R., Vignati, E., Wilson, J., Putaud, J.-P., Seinfeld, J. H., and Adams, P.: Formation and cycling of aerosols in the global troposphere, *Atmos. Environ.*, 34, 4215–4240, 2000.

15 Reinsch, C. H.: Smoothing by Spline Functions, *Numer. Math.*, 10, 177–183, 1967.

Sherwood, S. C.: A microphysical connection among biomass burning, cumulus clouds, and stratospheric moisture, *Science*, 295, 1272-1275, 2002.

Ström, J., Fischer, H., Lelieveld, J., and Schröder, F.: In situ measurements of microphysical properties and trace gases in two cumulonimbus anvils over western Europe, *J. Geophys. Res.*, 104, 12 221–12 226, 1999.

20 Weber, R. J., Clarke, A. D., Litchy, M., Li, J., Kok, G., Schillawski, R. D., and McMurry, P. H.: Spurious aerosol measurements when sampling from aircraft in the vicinity of clouds, *J. Geophys. Res.*, 103, 28 337–28 346, 1998.

Table 1. Parameters of the six Africa flights of project CARIBIC in 2000. ITCZ-North = Northern latitudinal boundary of the Intertropical Convergence Zone (ITCZ) along the CARIBIC flight track, delineated from high clouds in ECWMF analyses; in parentheses the corresponding values from strong gradients in the modeled 700 hPa temperature fields are given. ITCZ-South = as ITCZ-North but for the southern boundary of the ITCZ

#	From	Departure Date, UTC ¹	To	Arrival Date, UTC ¹	ITCZ-North	ITCZ-South
1	Munich	000518 19:40	Windhoek	000519 03:43	13° N (36° N)	4° S (7° S)
2	Windhoek	000519 06:32	Munich	000519 15:27	7° N (36° N)	0° N (7° S)
3	Munich	000727 19:00	Windhoek	000728 03:45	12° N (38° N)	4° S (4° S)
4	Windhoek	000728 06:31	Munich	000728 15:17	8° N (38° N)	3° S (5° S)
5	Munich	001202 11:30	Cape Town	001202 21:30	10° N (12° N)	22° S (29° S)
6	Cape Town	001203 06:44	Munich	001203 17:13	13° N (12° N)	20° S (29° S)

¹ start/stop of measurements

High aerosol concentrations in the upper troposphere over Africa

J. Heintzenberg et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Print Version

Interactive Discussion

High aerosol concentrations in the upper troposphere over Africa

J. Heintzenberg et al.

Table 2. Latitudinal arithmetic average number concentrations (cm^{-3} , STP) of the six Africa flights of project CARIBIC in 2000. Latitude ranges are given for each average as minimum and maximum latitudes (Lat_{\min} , Lat_{\max}). $N_{4,\text{av}}$ = diameters = 4 nm, $N_{12,\text{av}}$ = diameters = 12 nm, $N_{18,\text{av}}$ = diameters = 18 nm. Averages over all flights of regional number concentrations and average ratios of concentrations in the tropics to those in northern and southern subtropics are given in the last two lines, respectively

Subtropical southern hemisphere						Tropics					Subtropical northern hemisphere				
#	Lat_{\min}	Lat_{\max}	$N_{4,\text{av}}$	$N_{12,\text{av}}$	$N_{18,\text{av}}$	Lat_{\min}	Lat_{\max}	$N_{4,\text{av}}$	$N_{12,\text{av}}$	$N_{18,\text{av}}$	Lat_{\min}	Lat_{\max}	$N_{4,\text{av}}$	$N_{12,\text{av}}$	$N_{18,\text{av}}$
1	-18	-6.7	2200	2100	1780	-6.7	14	9940	9380	8900	14	30	2150	2110	1970
2	-18	-3.6	2730	2670	2380	-3.6	17	19000	11800	9810	17	30	2170	2120	1890
3	-21	-12	2680	2650	2480	-12	21	11700	11000	10400	21	30	923	918	919
4	-18	-5.5	1830	1830	1730	-5.5	19	24800	12200	8260	19	30	1610	1580	1280
5	-32	-25	1700	1710	1530	-25	13	8170	7130	5880	13	26	1470	1420	1260
6	-28	-23	1560	1620	1450	-23	13	30000	16400	11300	17	26	1580	1590	1350
Average			2120	2100	1890	17300			11300	9090	1650			1620	1450
Ratio tropical /subtropical			8.2	5.4	4.8						10			7.0	6.3

[Title Page](#)
[Abstract](#)
[Introduction](#)
[Conclusions](#)
[References](#)
[Tables](#)
[Figures](#)
[⏪](#)
[⏩](#)
[◀](#)
[▶](#)
[Back](#)
[Close](#)
[Full Screen / Esc](#)
[Print Version](#)
[Interactive Discussion](#)

High aerosol concentrations in the upper troposphere over AfricaJ. Heintzenberg et al.

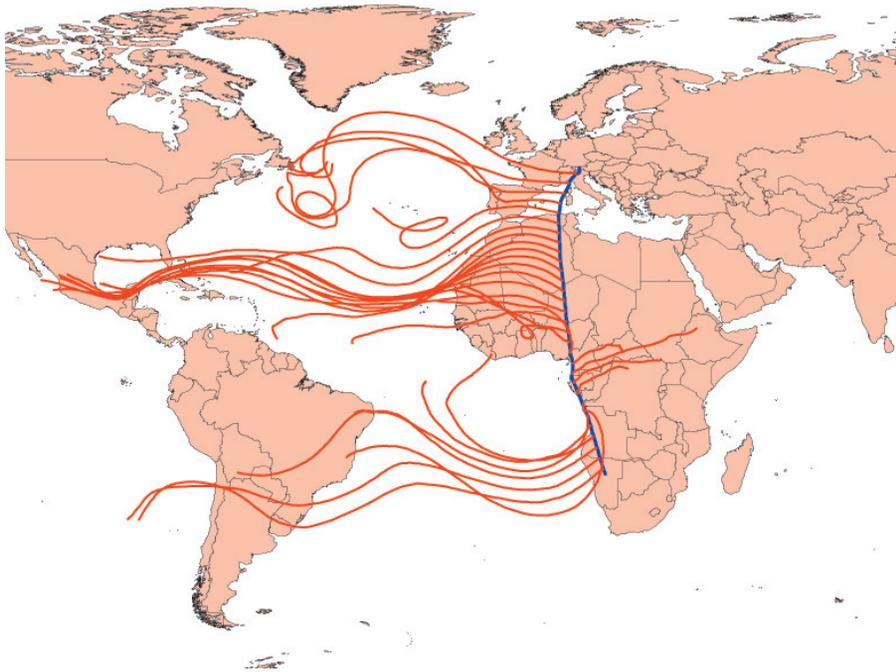


Fig. 1. (a) Flight route and horizontal projections of 3-dimensional 5-day back trajectories plotted from the flight positions every 30 min for flight 000519 from Windhoek, Namibia to Munich, Germany.

[Title Page](#)[Abstract](#)[Introduction](#)[Conclusions](#)[References](#)[Tables](#)[Figures](#)[⏪](#)[⏩](#)[◀](#)[▶](#)[Back](#)[Close](#)[Full Screen / Esc](#)[Print Version](#)[Interactive Discussion](#)

High aerosol concentrations in the upper troposphere over Africa

J. Heintzenberg et al.

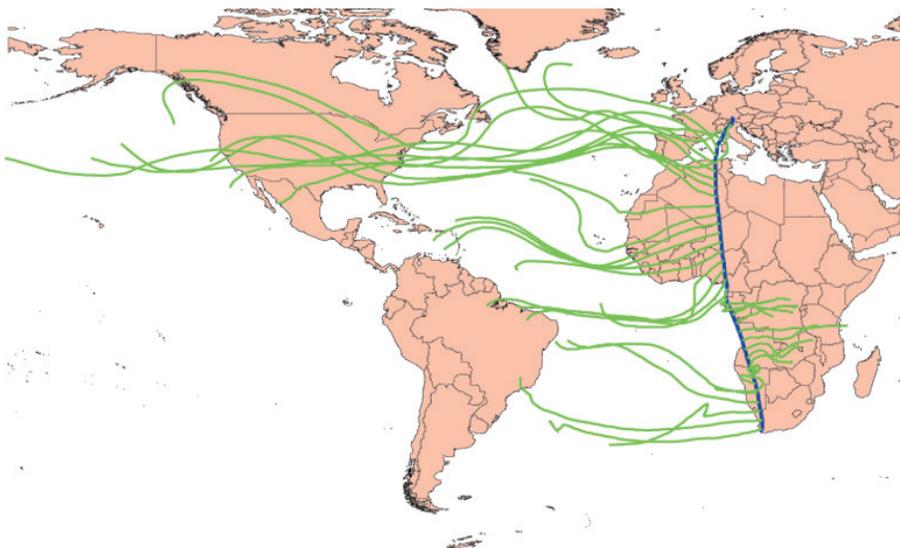


Fig. 1. (b) As (a) but for flight 001203 from Cape Town, South Africa to Munich, Germany.

[Title Page](#)[Abstract](#)[Introduction](#)[Conclusions](#)[References](#)[Tables](#)[Figures](#)[◀](#)[▶](#)[◀](#)[▶](#)[Back](#)[Close](#)[Full Screen / Esc](#)[Print Version](#)[Interactive Discussion](#)

High aerosol concentrations in the upper troposphere over AfricaJ. Heintzenberg et al.

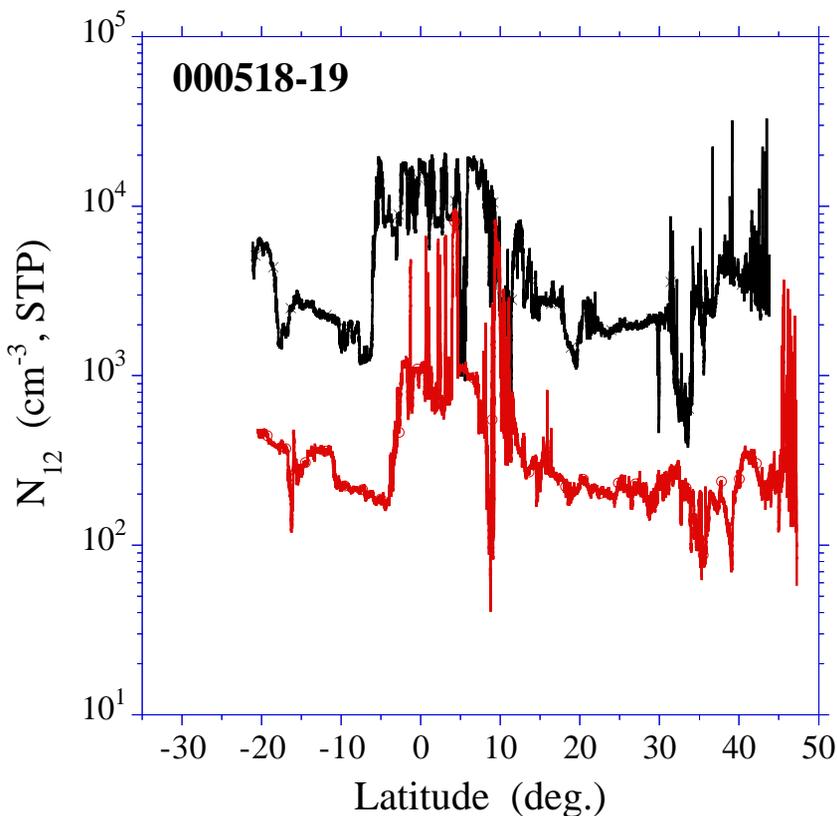
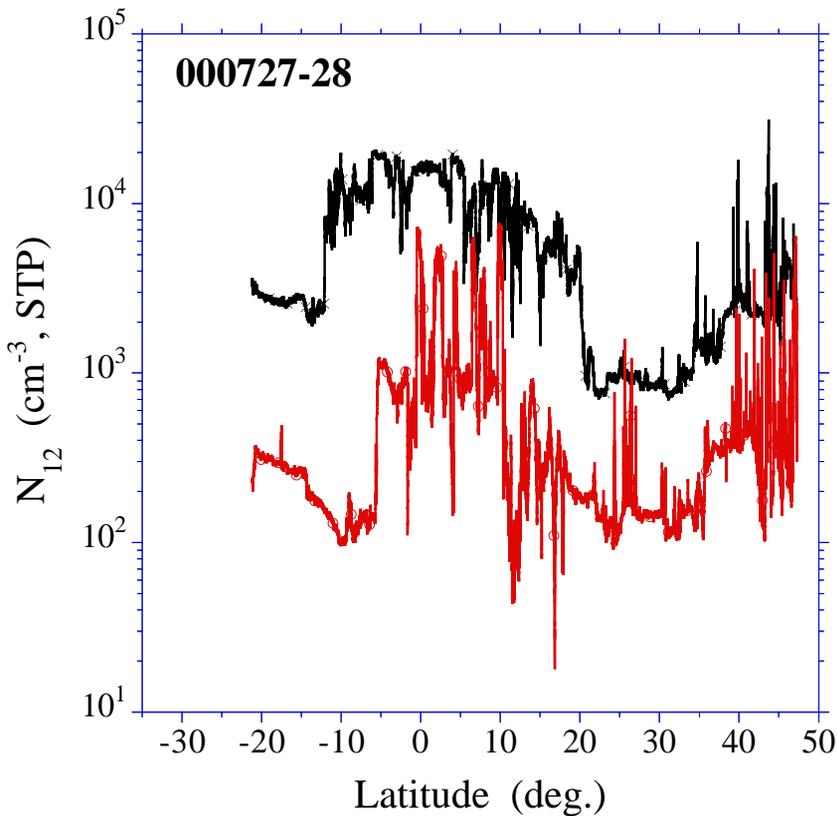


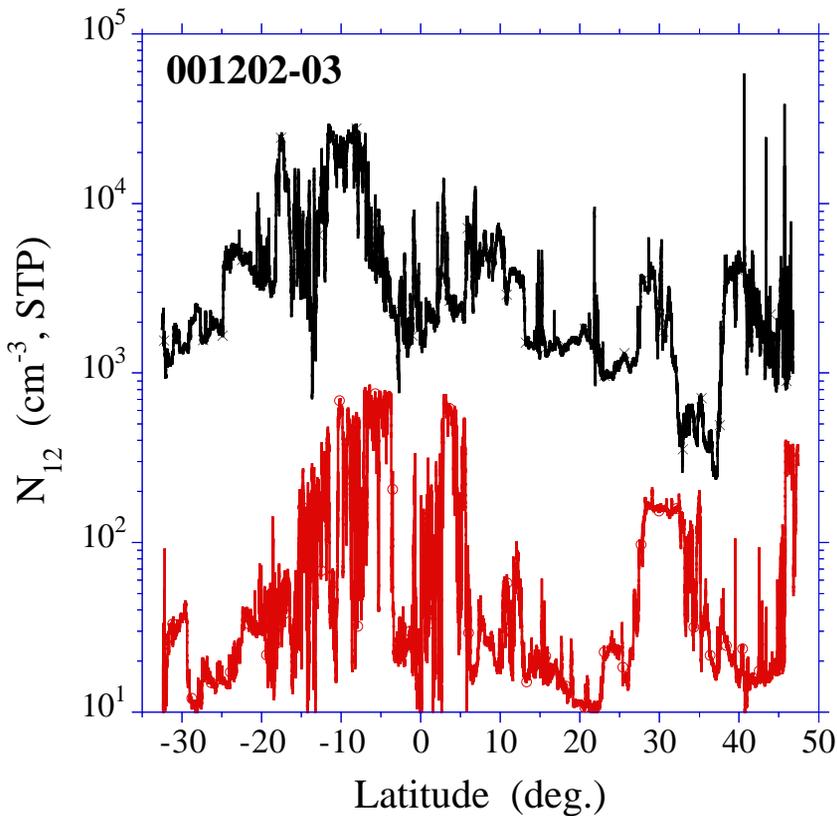
Fig. 2. (a) Latitudinal distributions of particle number concentrations larger than 12 nm in diameter (N_{12} , cm^{-3} , STP) for the six Africa flights of project CARIBIC. The flights are identified by their starting dates (cf. Table 1). Respective outbound and return flights are collected in one graph each. Data of return flights are shifted down by one order of magnitude, except for flight 001203, which is shifted down by two orders of magnitude to facilitate intercomparisons.

[Title Page](#)[Abstract](#)[Introduction](#)[Conclusions](#)[References](#)[Tables](#)[Figures](#)[◀](#)[▶](#)[◀](#)[▶](#)[Back](#)[Close](#)[Full Screen / Esc](#)[Print Version](#)[Interactive Discussion](#)

High aerosol concentrations in the upper troposphere over AfricaJ. Heintzenberg et al.

**Fig. 2. (a)** Continued.[Title Page](#)[Abstract](#)[Introduction](#)[Conclusions](#)[References](#)[Tables](#)[Figures](#)[◀](#)[▶](#)[◀](#)[▶](#)[Back](#)[Close](#)[Full Screen / Esc](#)[Print Version](#)[Interactive Discussion](#)

High aerosol concentrations in the upper troposphere over AfricaJ. Heintzenberg et al.

**Fig. 2. (a)** Continued.[Title Page](#)[Abstract](#)[Introduction](#)[Conclusions](#)[References](#)[Tables](#)[Figures](#)[◀](#)[▶](#)[◀](#)[▶](#)[Back](#)[Close](#)[Full Screen / Esc](#)[Print Version](#)[Interactive Discussion](#)

High aerosol concentrations in the upper troposphere over Africa

J. Heintzenberg et al.

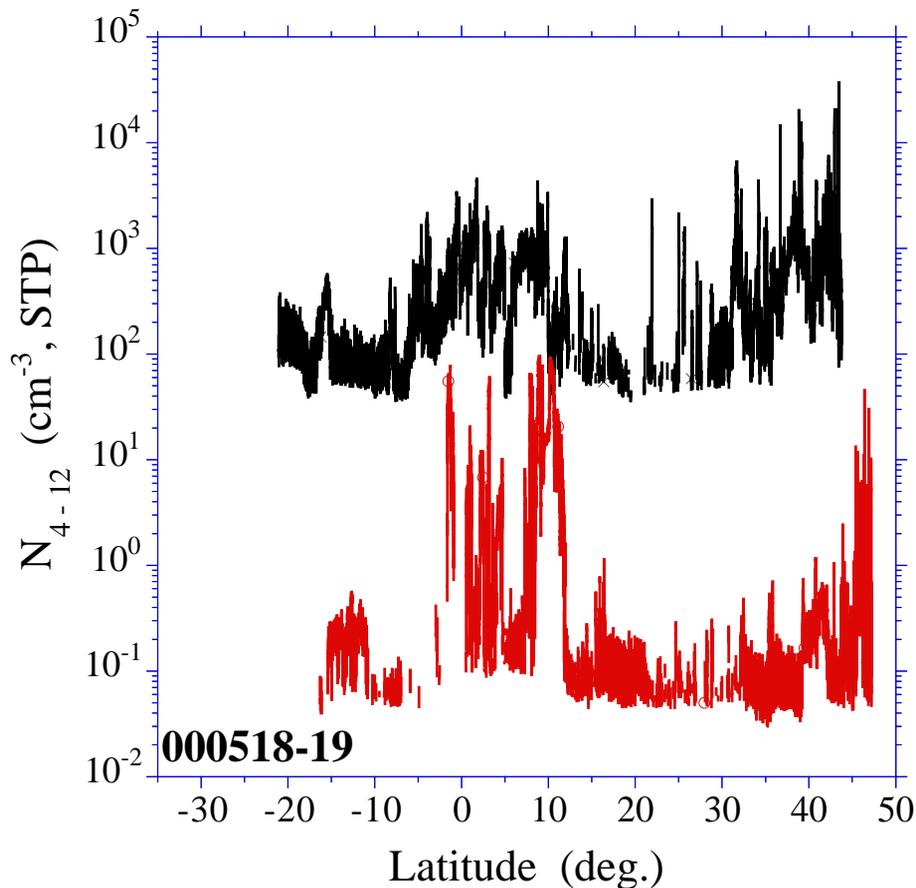
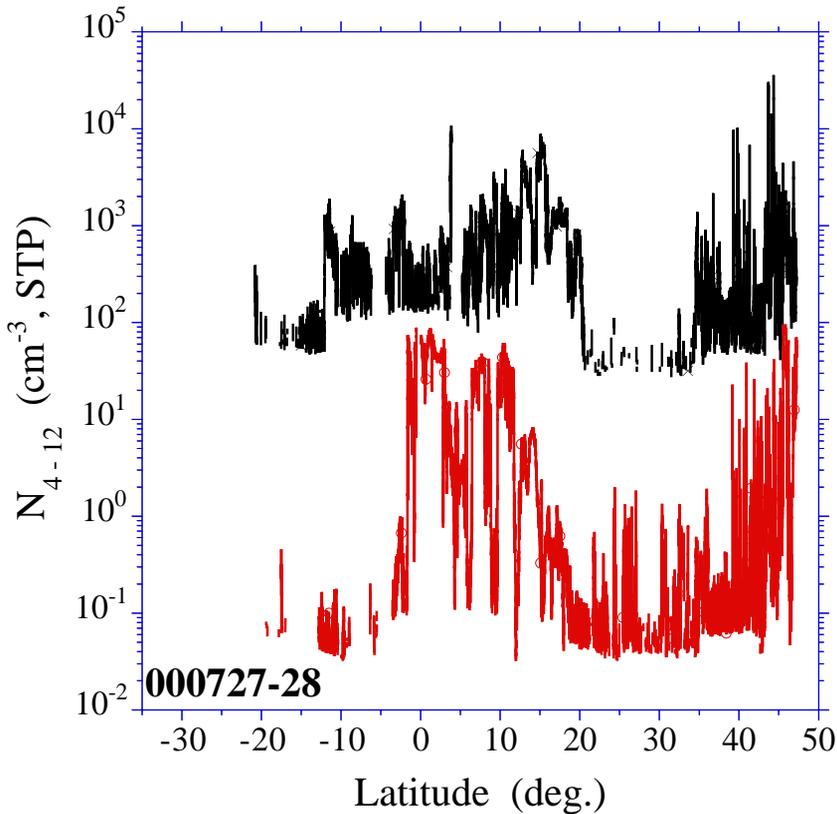


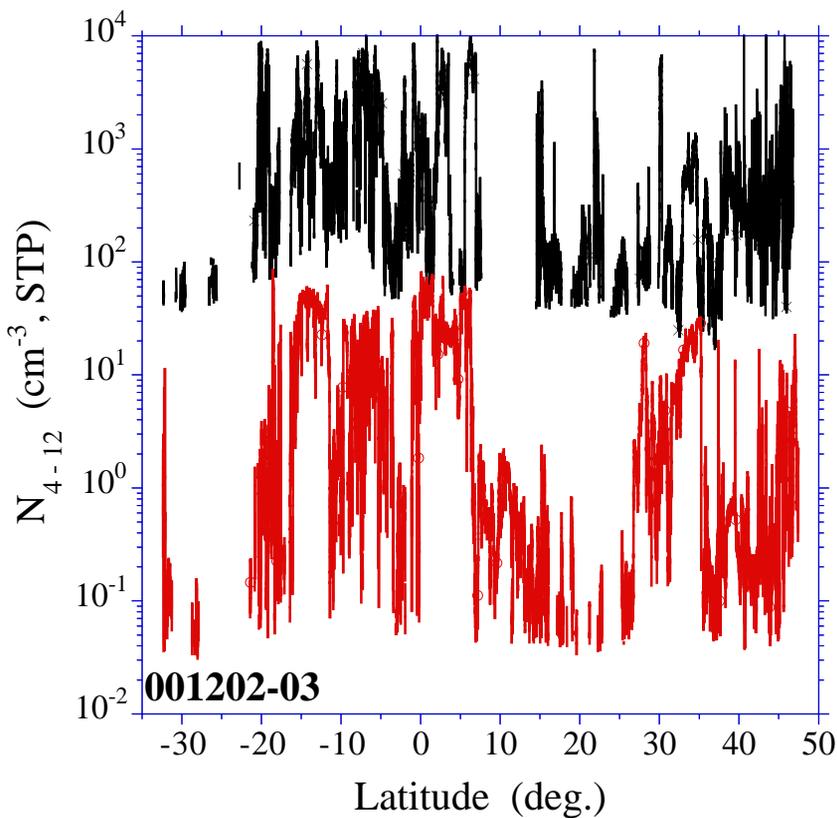
Fig. 2. (b) Latitudinal distributions of particle number concentrations between 4 and 12 nm in diameter (N_{4-12} , cm^{-3} , STP) for the six Africa flights of project CARIBIC. The flights are identified by their starting dates (cf. Table 1). Respective outbound and return flights are collected in one graph each. Data of return flights are shifted down by two orders of magnitude to facilitate intercomparisons.

[Title Page](#)[Abstract](#)[Introduction](#)[Conclusions](#)[References](#)[Tables](#)[Figures](#)[◀](#)[▶](#)[◀](#)[▶](#)[Back](#)[Close](#)[Full Screen / Esc](#)[Print Version](#)[Interactive Discussion](#)

High aerosol concentrations in the upper troposphere over AfricaJ. Heintzenberg et al.

**Fig. 2. (b)** Continued.[Title Page](#)[Abstract](#)[Introduction](#)[Conclusions](#)[References](#)[Tables](#)[Figures](#)[◀](#)[▶](#)[◀](#)[▶](#)[Back](#)[Close](#)[Full Screen / Esc](#)[Print Version](#)[Interactive Discussion](#)

High aerosol concentrations in the upper troposphere over AfricaJ. Heintzenberg et al.

**Fig. 2. (b)** Continued.[Title Page](#)[Abstract](#)[Introduction](#)[Conclusions](#)[References](#)[Tables](#)[Figures](#)[◀](#)[▶](#)[◀](#)[▶](#)[Back](#)[Close](#)[Full Screen / Esc](#)[Print Version](#)[Interactive Discussion](#)

**High aerosol
concentrations in the
upper troposphere
over Africa**

J. Heintzenberg et al.

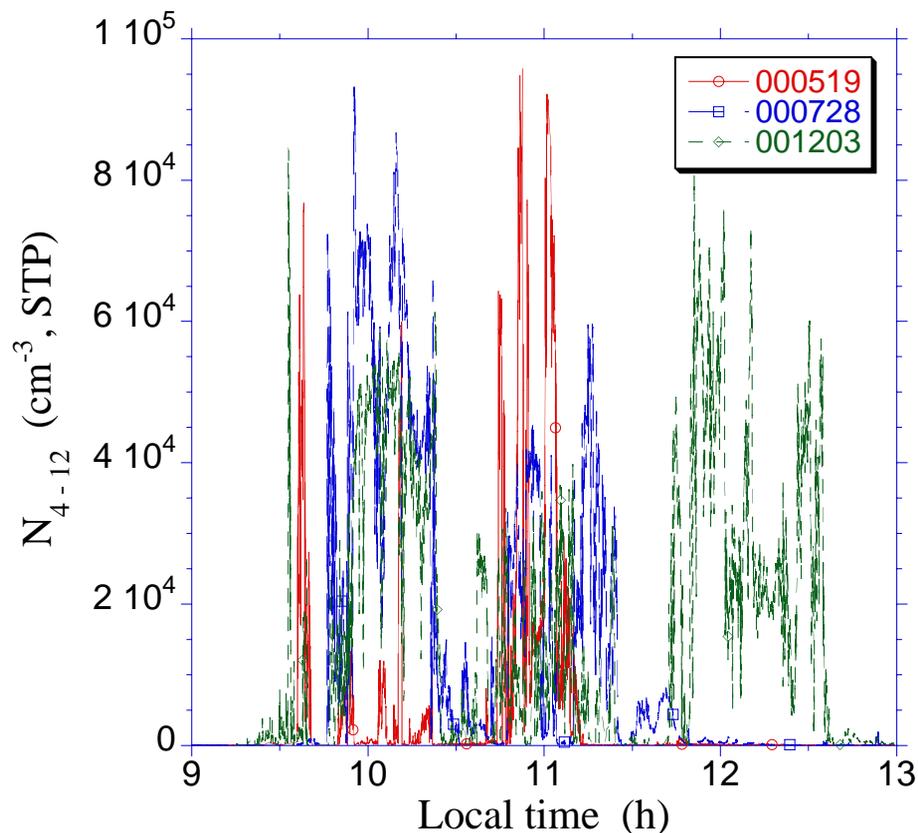


Fig. 3. Number concentrations of aerosol particles between 4 and 12 nm diameter (N_{4-12} , cm^{-3} , STP) as a function of local time during the period 09:00–13:00 on the flights 000519, 000728, and 001203.

[Title Page](#)[Abstract](#)[Introduction](#)[Conclusions](#)[References](#)[Tables](#)[Figures](#)[◀](#)[▶](#)[◀](#)[▶](#)[Back](#)[Close](#)[Full Screen / Esc](#)[Print Version](#)[Interactive Discussion](#)

High aerosol concentrations in the upper troposphere over AfricaJ. Heintzenberg et al.

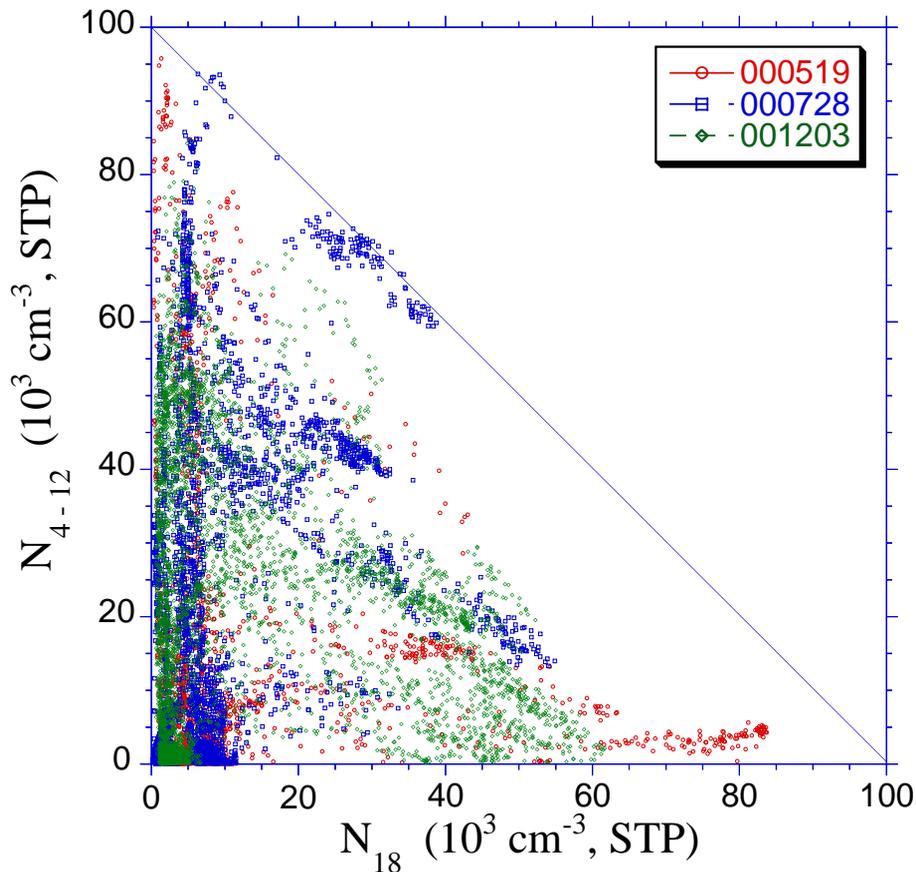


Fig. 4. Number concentrations of aerosol particles between 4 and 12 nm diameter (N_{4-12} , cm^{-3} , STP) as a function of number concentrations of aerosol particles larger than 18 nm diameter (N_{18}) cm^{-3} STP during local times between 09:00 and 13:00 on the flights 000519, 000728, and 001203.

[Title Page](#)[Abstract](#)[Introduction](#)[Conclusions](#)[References](#)[Tables](#)[Figures](#)[◀](#)[▶](#)[◀](#)[▶](#)[Back](#)[Close](#)[Full Screen / Esc](#)[Print Version](#)[Interactive Discussion](#)

High aerosol concentrations in the upper troposphere over Africa

J. Heintzenberg et al.

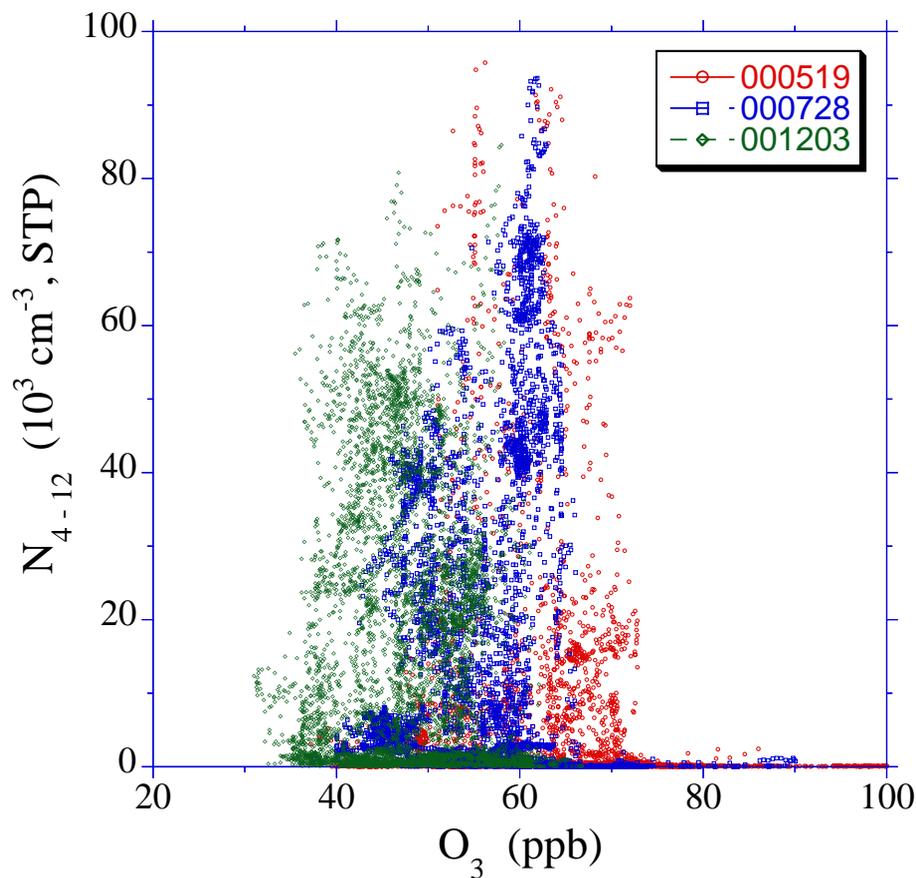


Fig. 5. Number concentrations of aerosol particles between 4 and 12 nm diameter (N_{4-12} , cm^{-3} , STP) as a function of local ozone concentrations in ppb during local times between 09:00 and 13:00 on the flights 000519, 000728, and 001203.

[Title Page](#)[Abstract](#)[Introduction](#)[Conclusions](#)[References](#)[Tables](#)[Figures](#)[◀](#)[▶](#)[◀](#)[▶](#)[Back](#)[Close](#)[Full Screen / Esc](#)[Print Version](#)[Interactive Discussion](#)

High aerosol concentrations in the upper troposphere over AfricaJ. Heintzenberg et al.

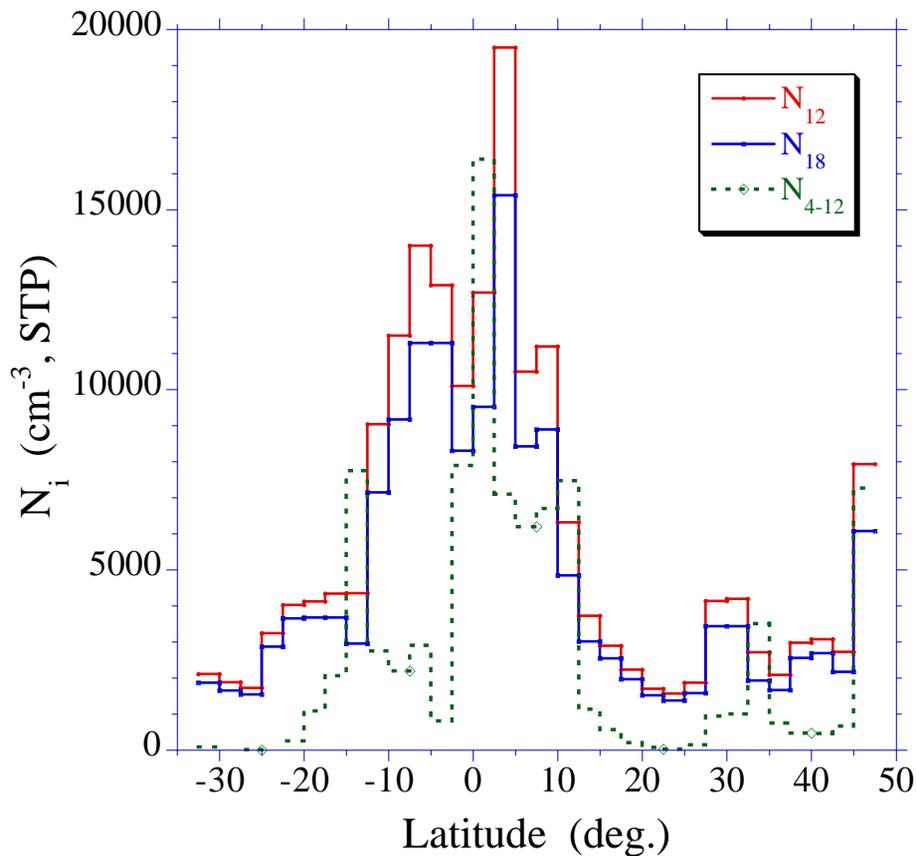


Fig. 6. (a) Latitudinal averages of number concentrations of aerosol particles larger than 4, 12, and 18 nm diameter (N_i , cm^{-3} , STP) taken on 6 flights in May, July, and December 2000.

[Title Page](#)[Abstract](#)[Introduction](#)[Conclusions](#)[References](#)[Tables](#)[Figures](#)[◀](#)[▶](#)[◀](#)[▶](#)[Back](#)[Close](#)[Full Screen / Esc](#)[Print Version](#)[Interactive Discussion](#)

High aerosol concentrations in the upper troposphere over AfricaJ. Heintzenberg et al.

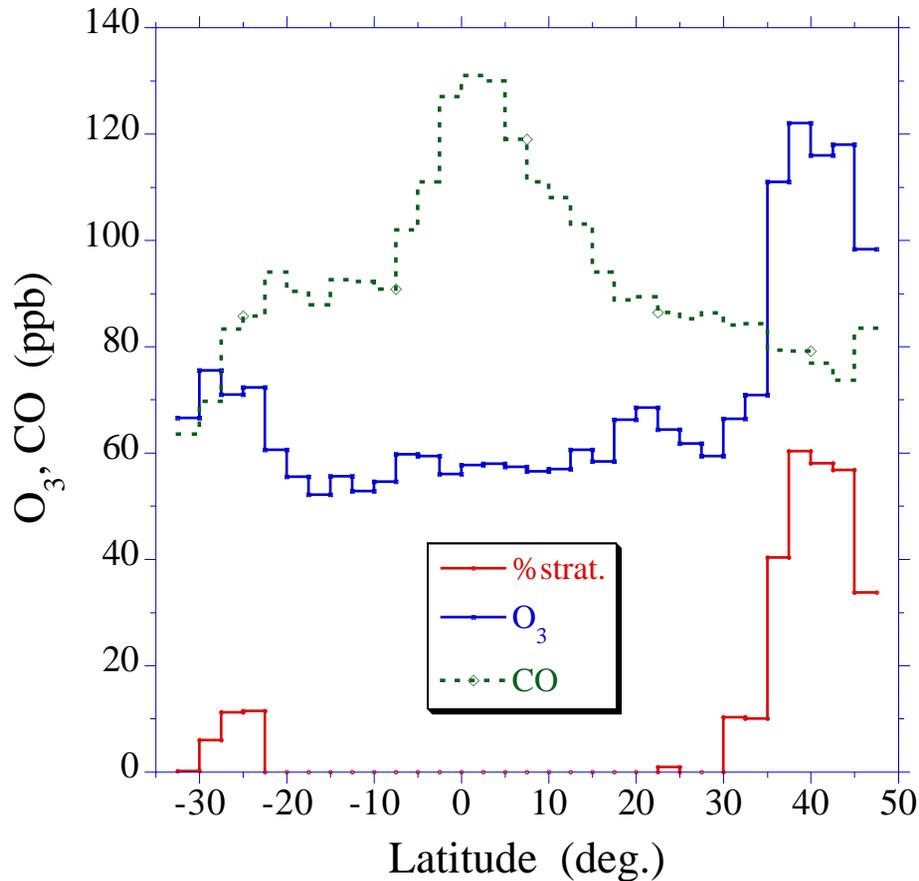


Fig. 6. (b) Latitudinal averages of CO, ozone, and percentage of flight time in the stratosphere (%strat) measured on 6 flights in May, July, and December 2000.

[Title Page](#)[Abstract](#)[Introduction](#)[Conclusions](#)[References](#)[Tables](#)[Figures](#)[◀](#)[▶](#)[◀](#)[▶](#)[Back](#)[Close](#)[Full Screen / Esc](#)[Print Version](#)[Interactive Discussion](#)