RESULTS OF OZONE MEASUREMENTS IN NORTHERN GERMANY
- A CASE STUDY -

Manfred Schmidt
Max-Planck-Institut für Aeronomie
D-3411 Katlenburg-Lindau, Germany

ABSTRACT

At most of the German ozone recording stations which have records over a sufficiently long period, the results of the summer months 1989 showed the highest values since the beginning of the measurements. One of the reasons for this phenomenon was the high duration of sunshine in that summer; for example, in Potsdam near Berlin in May 1989 the sunshine duration was the highest in May since the beginning of the records in 1893. For that reason we selected this summer for a case study.

The basis for the study was mainly the ozone measuring stations of the network of Lower Saxony and the Federal Office of Environment (Umweltbundesamt). The results of these summer measurements point to intense sources of ozone, probably in form of gaseous precursors, in the Middle German industrial areas near Leipzig and Halle and in Northwestern Czechoslovakia, with coal-mining, chemical and petrochemical industries, coking plants and others. The maps of average ozone concentrations, number of days with high ozone maxima, ozone-windroses of the stations etc., suggest that these areas could be a main source of precursors and of photochemical ozone production in summer smog episodes in Central Europe.

Stations on the North Sea coast, at which early ozone measurements were made by our institute in 1973/74 are compared with similarly located stations of the Lower Saxon network in 1989 and the results show a reversal of the ozone-windroses. In 1973/74, the highest ozone concentrations were correlated with wind directions from the sea while in 1989 these concentrations were correlated with directions from the continent. In the recent years, photochemical ozone production on the continent is probably predominant, while in former years the higher ozone content of the maritime subpolar air masses has been explained by stratospheric-tropospheric exchange.

1. INTRODUCTION

Figure 1 shows the summer ozone concentration averages (May-August) of some stations in Germany with longest duration of records; the longest series are at Arkona on the coast of the Baltic Sea (since 1856) and at Hohenpeilpenberg in southern Germany (since 1971) [e.g. Low et al., 1990]. The geographical positions of these stations, and others considered in this paper, are plotted in Fig. 2. The most striking features in Fig. 1 are the very high O₃ values in the summer of 1989, at all stations without exception. These are the highest average concentrations since beginning of the records. The 1989 summer in Germany was dry and the recorded sunshine durations and global radiations were far above normal (up to 120-140% sunshine duration). If we consider sufficient concentrations of gaseous precursors, NOₓ, CO, CH₄ and non-methane hydrocarbons (NMHC), an increased photochemical produc-
tion of ozone during these enhanced radiation conditions with simultaneously increased UV radiation is to be expected [Schmidt, 1989]. A further reason for choosing the 1989 summer months for a case study was the unusually high ozone concentration at all stations in the southern part of Lower Saxony, the district which contains the city of Göttingen and the location of our institute. This is the main point of discussion in this paper.

We have, therefore, concentrated on air quality measurement data from the 17 stations in the Federal State Lower Saxony, in addition to the 15 stations of the Federal Office of Environment, and the two WMO Background Air Pollution Monitoring stations of the former GDR (names and positions in Fig. 2). Wind direction measurements have been taken from suitable stations of the German Weather Service.

2. RESULTS

In Fig. 3, we have plotted the average ozone concentrations for May–August 1989 from all stations mentioned and indicated in Fig. 2. To a careful first approximation, we have also included isolines. Unexpectedly, "a priori", the highest ozone concentrations > 100 µg/m³ appeared in districts north and southeast of Göttingen, > 90 and > 80 µg/m³ in the adjacent regions, all situated near the border to the former GDR. Similar high concentrations were
found in the northernmost parts of Germany, at and near the North Sea and the Baltic Sea coasts, and again at elevated places in Southern Germany. The highest average concentration of 130 μg/m³ was measured at the peak of Schauinsland in the Black Forest, 1284 m a.s.l. At this mountain station, the O₃ concentrations are normally the highest of all stations, also during other periods – months and seasons –, a fact, which should be investigated further on. Very surprising and impressive are supporting results, shown in Fig. 4, the number of days with O₃ maxima > 180 μg/m³ at the stations considered. This value is an upper limit during summer smog episodes, from which the authorities disseminate warnings to patients and persons with diseases affected by high ozone concentrations and other poisonous substances in the photochemical smog. In the southeastern part of Lower Saxony and probably to the east we find the highest number of such days, more than 20 to 40, in the summer of 1989. Similar high values, 31 and 35 days, appeared at two further stations, Schauinsland (1284 m) and Deuselbach (460 m a.s.l.), the latter probably influenced by two other industrial centers (Fig. 5 and section 3).

In Fig. 5, we have plotted the directions and amounts of the maximum and 2nd maximum of the ozone windroses of nearly all of the stations. These ozone wind vectors are the key to an explanation of the results in Figs. 3 and 4.

3. DISCUSSIONS

The results given in Figs. 3, 4 and 5, notably in Fig. 4, suggest that the sources of these unusually high ozone concentrations may be in the strongly polluted industrial areas of the former GDR, in the district Halle–Leipzig–Bitterfeld (at the time of these measurements, 1989, HL, L, Bi in Figs. 3–5). Nearly all maximum ozone-wind vectors of nearby stations in Fig. 5 point in this direction. Similar strongly polluted industrial areas are also located in the northwestern part of CSFR (Fig. 2), indicated by an extension of the vectors above. Even the vectors of the station Brotjackriegel in the Bavarian Forest and one vector of the station Hof point in this direction.

The industrial areas just mentioned are, admittedly, the highest polluted areas in Central Europe, particularly with regard to the SO₂ emission. One of the reasons for this is the almost exclusive use of the pit-brown coal for energy production, raw material for chemical and petrochemical industries and for coke production. The most important chemical factories of the former GDR are located in the district Halle–Bitterfeld. One of these factories was the largest factory of all kinds in the GDR. These factories mainly produced fuels of all kinds, acids, nitrogen fertilizers, technical gases, chemicals, medicaments and plastics. One of these factories was the greatest producer of calcium carbide CaC₂ in Europe, made from brown coal and lime, with water transformed into acetylene C₂H₂, and finally used as a basic substance via other NMHC’s for the production of plastics. From extended artificial mud pools containing the remaining Ca(OH)₂, large amounts of residual C₂H₂ must surely have escaped into the atmosphere, to be converted there into further NMHC’s. Sources of precursors are to be expected from parts of the gas, escaping from coking plants operated there. Coke-oven gas mainly contains tar vapour, H₂, CH₄, NMHC’s, N₂, CO, CO₂ and others, most of them being precursors for photochemical O₃ production. Unfortunately, no direct measurements of trace gases including ozone are available from these areas before the time of unification of the German states. Our results are supported by some other observations. Kourtidis [1991] published results of PAN measurements, made in the spring of 1989 at our institute in Katlenburg-Lindau near Göttingen. His PAN windroses point with the maxima in the same directions, East to South, like our windrose vectors in Fig. 5. In the 3 summers prior to 1989, 1986 – 1988, the numbers of half-hour averages of O₃ concentrations > 120 μg/m³ and > 180 μg/m³ were highest at the two Lower Saxon stations Dassel and Oker, compared with 4 further stations in the North, Hanover, Braunschweig, Wilhelmshaven and Emden. All stations attained maxima in 1989. Even the annual O₃ means of Dassel and Oker in the years 1986–1988 show higher values than those of Hanover and Emden. In these years, the number of stations was still reduced.
ney increased ozone concentrations were found in all sea-

cent Lower Saxon network (Fig. 2). At station Norder-

ozone measuring station at Norderney, in the same coastal

seventies

ted in Fig. 6. During project "Tropospheric Ozone" in the

also should be the subject of further investigation.

wind

from

of ozone, transported by subpolar air masses and coming

dle European sea shores are, at least partially,

the Northwest in Arkona. These facts suggest that the

maximum in Fig. 5. The vectors of HohenpeiBenberg to the similarly populated dis-

and coal

thermore to the South, in the direction of the

Mannheim-Ludwigshafen-Frankfort are located and, fur-

directions where the

vectors of Helmshaven

point to the densely populated area of Stuttgart, the

vectors of Hohenpeissenberg to the similarly populated dis-

trict of Munich [like Wege and Vandersee, 1991]. In the

northermost part of Germany, the ozone concentrations in

Fig. 3 increase towards the North, up to 100 \( \mu g/m^3 \) at

station Westerland. The vectors of the 2nd ozone max-

imum in Fig. 5 point to the West in Westerland and to

the Northwest in Arkona. These facts suggest that the

ozone concentrations at such rural stations near the Mid-

tle European sea shores are, at least partially, composed

of ozone, transported by subpolar air masses and coming

from stratospheric-tropospheric exchange processes, if the

wind is blowing from the North towards the stations. This

also should be the subject of further investigation.

An interesting result, regarding this question, is plotted

in Fig. 6. During project "Tropospheric Ozone" in the

seventies [e.g. Schmidt, 1981], our institute operated an

ozone measuring station at Norderney, in the same coastal

area as the stations Emden and Wilhelmshaven of the re-

cent Lower Saxon network (Fig. 2). At station Norder-

ney increased ozone concentrations were found in all sea-

ons during wind directions from the Southwest via North
to the Northeast, i.e. from the North Sea and Baltic Sea, 

and decreased concentrations from all other directions, i.e.

from the European continent. Fig. 6 shows two examples

in the summers of 1973 and 1974 (N73,74). On the other

hand, the nearby located stations Emden (E89) and Wil-

helmshaven (W89) show the highest \( O_3 \) concentrations

15 years later in the summer of 1989 during wind directions

from the continent, with slightly lower concentrations when

the wind was from the North Sea, W to NNW. The total

sunshine durations during each of these 3 summers were

nearly the same, as indicated in Fig. 6 below. An explana-
tion of these contrary results could be that, in recent years,

the photochemical ozone production on the European con-

tinent was predominant, especially in the summer months,

and caused by a distinct increase of anthropogenic gaseous

precursors; this is also suggested by measurements. In for-

mer years, the higher ozone content of the subpolar air

masses correlated with wind directions from the northern

sector; this was explained by stratospheric-tropospheric ex-

change and demonstrated in a series of papers [e.g. Fabian

and Pruchniewicz, 1977; Schmidt and Fabian, 1980].

4. CONCLUSIONS

The assumption is justified, the industrial areas in

Eastern Germany and CSFR have been important sources

of precursor gases for the ozone production in summers

such as 1989 with its relatively strong UV radiation. Since

some of these factories are now obsolete and have since been

closed (for renewal), further investigations of ozone during

sunny summers in the future should consider, whether or

not the ozone concentrations will now decrease in these

locations.

REFERENCES

Fabian, P. and P.G. Pruchniewicz: Meridional Distribution of Ozone

in the Troposphere and Its Seasonal Variations. J.G.R. 82,

2063-2073, 1977

Kourtidia, K.: Analysis of PAN (Peroxyacetyl nitrate) in the Atmos-

tphere. PhD Thesis, Aristoteles-University of Thessaloniki,

Greece, 1991

Low, P.S., T.D. Davies, P.M. Kelly and G. Farmer: Trends in Surface

Ozone at Hohenpeissenberg and Arkona. J.G.R. 95, 22441-

22453, 1990

Schmidt, M. and P. Fabian: Relationships Between Tropospheric

Ozone Concentration and the General Weather Situation.


Schmidt, M.: Further Results from the MPAE-Project "Tropospheric


Schmidt, M.: Relationships Between Tropospheric Ozone and Global

Solar Radiation in Germany. In: Bojkov R.D. and P. Fabian

(Eds): Ozone in the Atmosphere, 451-454, Deepak Publ., Hamp-

ton, VA, 1989

Wege, K. und W. Vandersee: Ozonbeobachtungen am Nordrand der


173