

NOAA Technical Memorandum ERL ARL-155



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THE USE OF BACK AIR TRAJECTORIES IN INTERPRETING ATMOSPHERIC  
CHEMISTRY DATA: A REVIEW AND BIBLIOGRAPHY

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# THE USE OF BACK AIR TRAJECTORIES IN INTERPRETING ATMOSPHERIC CHEMISTRY DATA: A REVIEW AND BIBLIOGRAPHY

John M. Miller

**ABSTRACT.** One of the most important aspects of atmospheric chemistry is interpreting the measurements of different chemical species in terms of their cycles in the earth-atmosphere system. One method--the calculation of single, back trajectories from the point of measurement--has been widely and successfully employed to evaluate transport of chemical substances in the atmosphere. This paper reviews how the single-trajectory analysis has been applied in atmospheric chemistry. It surveys the different models used, the methods of application, and the forms in which the results are presented. Important new analysis techniques have been developed in the last few years, such as flow climatology, automatic computer categorization, the use of meteorological forecast model output, and others. Despite its limitations, the single, back-trajectory analysis method has proved very useful in the interpretation of atmospheric chemistry data.

## 1. INTRODUCTION

Attempting to describe the path that both natural and anthropogenic substances follow in the atmosphere has become an important area of meteorological research. For the last several decades, scientists have tried to design ways of understanding and measuring these atmospheric pathways. Analyzing such transport requires that the measurements of the structure of the atmosphere be organized in some fashion. This paper is a review of one approach that has proved particularly useful, that is, calculation of single, back trajectories and evaluation of atmospheric chemistry data using these calculations.

Use of the concept of a trajectory has been a standard meteorological technique for over 30 years. In essence, a trajectory is the path of an imaginary air parcel as it is acted on by the winds (Huschke, 1959). Usually this idea has been applied to forward motion; that is, the parcel starts at some point and is pushed by the winds at a given level or surface of interest. One application of single, forward trajectories is to follow a parcel from the source of a given substance and to see where the material could be transported. When continuous or intermittent atmospheric chemistry measurements are made at some point, either on the ground or in the air, then the origin of the path of travel of that sampled air is of interest in interpreting the data. Thus for a chemical measurement at a given time and place, a single, back trajectory can be calculated, giving the path that the air parcel traveled previous to the measurement. An example of several single, back trajectories for one day is shown in Fig. 1, where the time refers to when the imaginary parcel whose path is described by the trajectory passed over the point of interest (the western Mediterranean). In this case, 10-day, 850 mb and 700-mb trajectories are shown. The number refers to the day back in time; that is, 1 is the position of the parcel 24 hours earlier, 2 is 48 hours earlier, etc.

MED TRAJECTORIES ARRIVING 81007 - 1/ 7/81

A: 0Z 850 MB

C: 12Z 850 MB

B: 0Z 700 MB

D: 12Z 700 MB



Figure 1. Example of 10-day back trajectories for January 7, 1981, from a point in the western Mediterranean. The 850-mb trajectories are A (00Z) and C (12Z); the 700-mb trajectories are B (00Z) and D (12Z). The numbers (1, 2, etc.) refer to days back in time.

Though the parcel is conceptionally helpful in describing the movement at a given level of the atmosphere, its use in understanding the transport of trace substances in the atmosphere can be very complicated for any single atmospheric event. Most of these materials are released at or near the earth's surface and may be eventually mixed through several kilometers of the atmosphere. If the parcel is restricted to a given level, then such widely dispersed trace materials could be transported by a group of parcels moving in different directions in both the vertical and the horizontal. Naturally, if the flow were unidirectional at several levels with a small vertical component, then describing the transport would be relatively simple. Under chaotic, discontinuous conditions, common in frontal discontinuities, parcels would follow different paths at different levels, and transport would be very complicated. This limitation of applying the parcel method must always be kept in mind when using the back-trajectory technique.

## 2. APPLICATION OF BACK-TRAJECTORY ANALYSIS TO ATMOSPHERIC CHEMISTRY

### 2.1 Method

Over a decade ago, objective computer methods (Heffter et al., 1975; Pack et al., 1978) were developed to take different parameters such as pressure, wind or temperature fields and, using interpolation schemes, to calculate trajectories on a given level. One advantage of the computer approach is the ability to make consistent calculations over long periods of time. Also, different fields can be mixed so that, for example, measured winds can be interpolated to an isentropic surface and trajectories constructed (Merrill et al., 1985). Thus the computer allows a broad range of options that can be applied to trajectory calculations and their interpretation.

Before reviewing the application of trajectories to atmospheric chemistry, it is appropriate to discuss the different approaches meteorologists have developed to calculate trajectories from the data, i.e., the twice-a-day vertical soundings. There are two basic kinds of trajectories---dynamic and kinematic. Dynamic calculations are made with such fields as pressure and temperature to draw isobaric or isentropic trajectories, whereas kinematic calculations are made with measured wind fields. A breakdown is shown in Table 1 in which the advantages and disadvantages of the different models are outlined. Though the isentropic dynamic model is more satisfying from a theoretical viewpoint, its added sophistication may not always improve the results because of a variety of factors (Artz et al., 1985; Merrill et al., 1986). When the flow is simple, the isentropic dynamic and the kinematic models track the same paths very well. In more complex meteorological situations, one can assume but not prove that the added calculation used in the isentropic approach gives better, more accurate trajectories.

On the basis of the above discussions, the application of the back-trajectory method is determined by the following factors: (1) the choice of the type of model used (dynamic, kinematic, mixed); (2) the level at which the calculation is made (whether it is a pressure surface, potential temperature field, layer in the atmosphere, etc.); (3) the grid size, which determines the area of interest and the distance back a trajectory can go; (4) the number of days backward; (5) the availability of data; and (6) the computer time required to make necessary calculations. Decisions about these and other practical problems must be made before the first trajectory can be drawn.

Table 1. Single back-trajectory methods

Kind	Type	Advantages	Disadvantages
Dynamic	Isobaric	Easy to calculate; large data bank; operational	Geostrophic winds not representative; no vertical motion
	Isentropic	Vertical motions taken into account	Large amount of computer time; good only for case studies; not operational
Kinematic	Wind field (real or calculated)	Easy to run; operational	No vertical motion
Mixed	Wind fields on isentropic surfaces	Vertical motion; easier to run than pure isentropic	Not completely operational; misses vertical motion due to adiabatic processes



## 2.2 Application

After the above decisions are made, the researcher is ready to use trajectories in data analysis. In the single, back-trajectory analysis the usual approach is to compare directly a 5- to 10-day back trajectory with an atmospheric chemistry measurement made over a 12- to 24-h period at the point of origin of the trajectory. If there are only a few measurements at the point, then each can be compared directly with its trajectory, as discussed by Byrd and Andreae (1986), Colbeck and Harrison (1985b), Galloway et al. (1983), Halter and Peterson (1981), and many others. If, however, there are a large set of measurements, a more structured approach must be made, and therefore a method of trajectory classification is required. The two elements in any classification scheme are to account for the direction from which the trajectory came and the distance it took to travel in a given time, i.e., the speed of transport. Direction is usually categorized by some kind of sectoring arrangement. The schemes are either a type of mechanical division, such as 45° quadrants, or are based on a hypothesis that a given sector represents a special transport zone. Figure 2 shows examples of these two kinds of classification schemes. Each sector may also be divided into distances from the origin so that the probable speed of flow can be determined.

How one does the classification must also be considered. To date, the most usual method is inspection by the individual researcher. This may lead to differences in interpretation, as is the case in analyzing weather maps. However, the approach ensures that each individual trajectory will be handled separately. For large data sets visual classification can be a laborious process. Examples of this method are given, for example, in Kurtz et al. (1984), Merrill et al. (1985), and Munn et al. (1984). An alternate approach is to design a scheme that allows automatic analysis by the computer. Though this saves considerable time, it may also lead to spurious results during complicated weather situations.

## 3. NEW APPROACHES

Though the methods discussed in Sec. 2 for applying back-trajectory analysis have been useful, researchers have been looking for other ways of extracting further information using this technique. Four methods seem promising: (1) automated long-term analysis of trajectories, i.e., flow climatology; (2) detailed studies of individual trajectories such as including diurnal changes, following layers, and adding extra fields such as rainfall amount along the trajectory path; (3) use of the fields provided by forecast models, to account for vertical motions; and (4) field studies to verify back trajectory calculations. Application of these new approaches could be an aid in establishing further understanding of transport and atmospheric chemistry that was not possible using earlier analysis techniques.

### 3.1 Flow Climatologies

From daily trajectories, a flow climatology can be constructed for the location of interest. Thus the transport potential of a given substance can be evaluated assuming that the analysis is tailored to the possible source areas. This is illustrated by an example shown in Fig. 3 (Miller et al., 1987). The ARL trajectory model was used to calculate 10 days back from a point in the

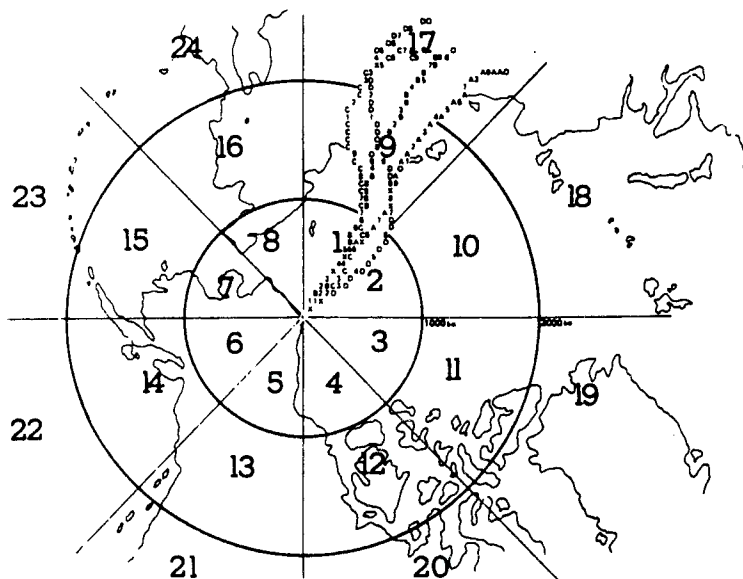


Figure 2(a). A scheme in which distances and speed of flow can be determined for each regularly spaced sector (Miller 1981a).

### CARIBOU TRAJECTORY CLASSIFICATION

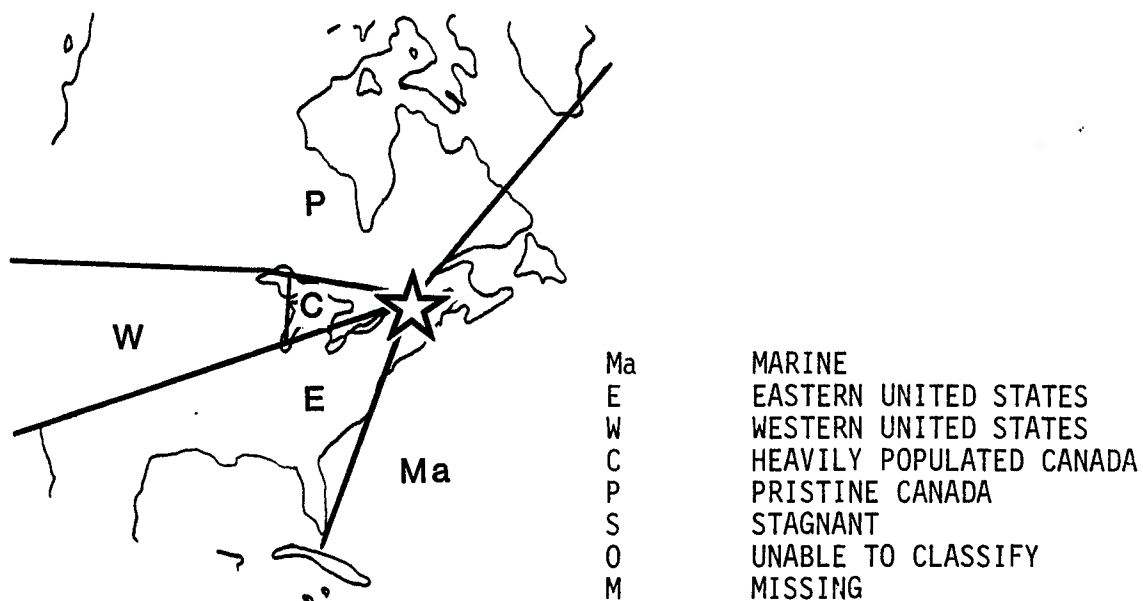


Figure 2(b). A scheme in which categorization is according to pollution sources (Artz and Dayan, 1986).

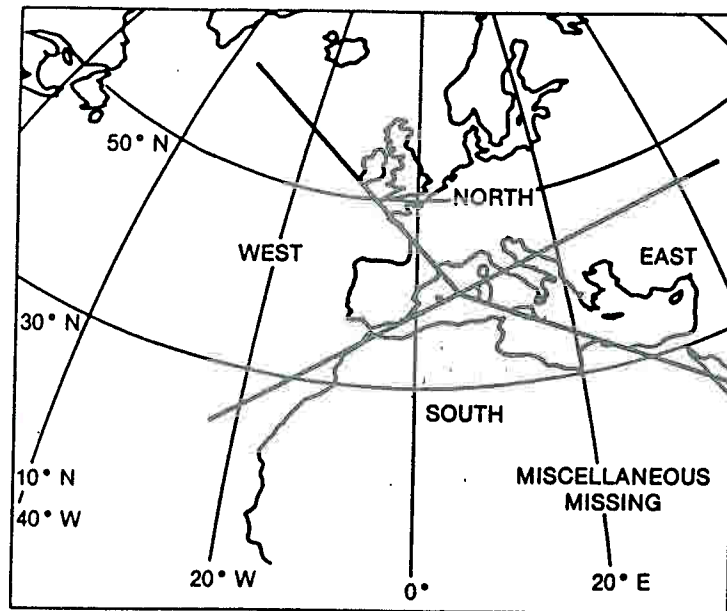


Figure 3(a). The typing scheme used to categorize 10 years (January 1975–December 1984) of 850-mb back trajectories from a point in the western Mediterranean.

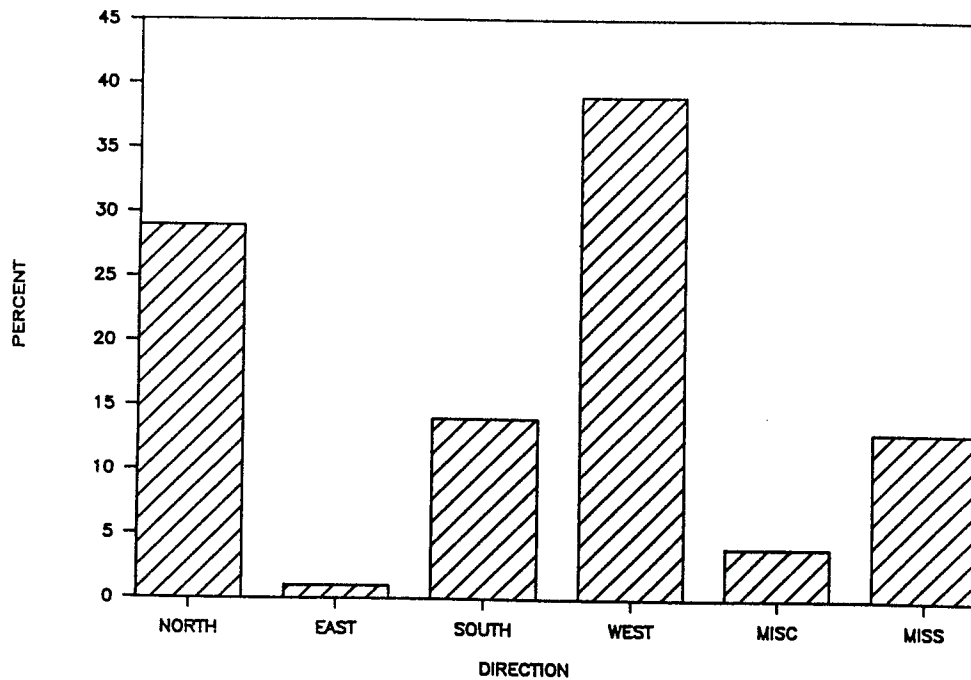


Figure 3(b). The number of trajectories for each type (Miller et al., 1987).

western Mediterranean (40°N, 6°E) for the period from January 1975 to December 1984. This procedure was performed twice a day for the 850-mb level. The trajectories were categorized according to the following scheme (see Fig. 3). There are six different categories: (1) North--trajectories coming from this area would conceivably bring more pollution with them; (2) East--trajectories rarely come from this direction; (3) South--this flow pattern brings air from the Sahara with accompanying desert dust; (4) West--trajectories from this sector could be expected to be the most clean; (5) Miscellaneous--this case includes both the times when the trajectories show strong cyclonic motion and the times when a categorization is impossible because of weak flow; and (6) Missing. The summary for the 10-yr period is shown in Fig. 3, which demonstrates the prevailing flow patterns to the area. Similar graphs can be made showing seasonal variation and year-to-year differences. Publications illustrating this approach are, for example, Pacyna et al. (1984a), Parekh and Husain (1982), Miller (1981a, 1981b), and Pitchford and Pitchford (1985).

A second example of flow climatology is presented in Lehmhaus et al. (1985). A trajectory rose was constructed for each station in the European monitoring network for the period of January 1978 to October 1982 (Fig. 4). Four 96-h 850-mb back trajectories were calculated to arrive at the station each day. Eight transport sectors were established based on 45° sections such as North, Northeast, etc. The position of the trajectories was identified every second hour. If more than half of the positions of the four trajectories in a given day were within a certain sector, then the day was allotted to that sector. A ninth sector was defined when this was not the case. Only trajectories between 150 and 1500 km were considered. An example of the results is shown for one station, F1 at Vert-le-Petit, France. Not only are the number of cases shown for each sector, but they are also compared with the measured and calculated values of SO<sub>2</sub>. This approach allows quick computer analysis of the trajectory data but could present uncertainties, because not all trajectories can be correctly identified because of the mechanical typing scheme.

A third approach to flow climatology and its relationship to atmospheric chemistry was shown by Munn et al. (1984). These authors developed a method of counting the number of trajectories that crossed grid squares of a 50 x 50 km<sup>2</sup> area. Thus they were able to draw isopleths of trajectory crossings for the entire period of study. A similar method was developed by Henmi and Bresch (1985). Figure 5 shows an example of this calculation, in which the percentages of times the trajectories of a given period were in a given grid square are displayed for the uppermost 10% of sulfate concentrations.

### 3.2 Multifield Analysis

The use of back trajectories can be further enhanced by the inclusion of other fields in the trajectory evaluation, such as temperature, relative humidity, and surface winds. Measurements such as gas and aerosol concentration fields could also be incorporated in the analysis upwind of the location of interest. Draxler (1983), for example, used precipitation, relative humidity, and temperature fields to evaluate the transport of regional sources of pollution. A difficulty with the multifield approach is to express these other fields in a form that can be used in the calculations. In addition, the manner in which each field interacts with the other is hard to establish and thus a number of assumptions must be made. Though this approach does present promise, establishing the fields needed for the calculations is a major handicap.

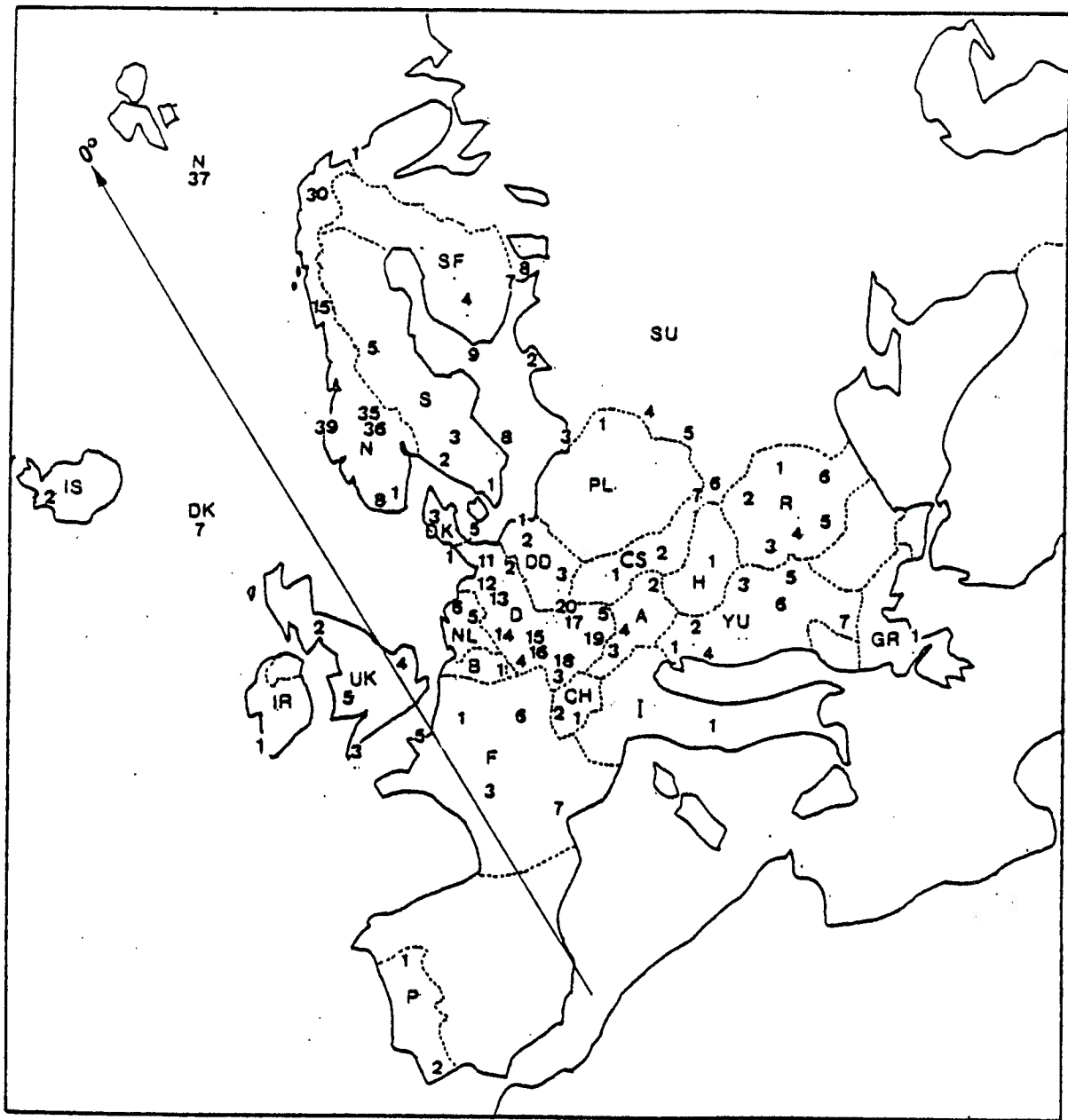


Figure 4(a). Locations of EMEP stations.

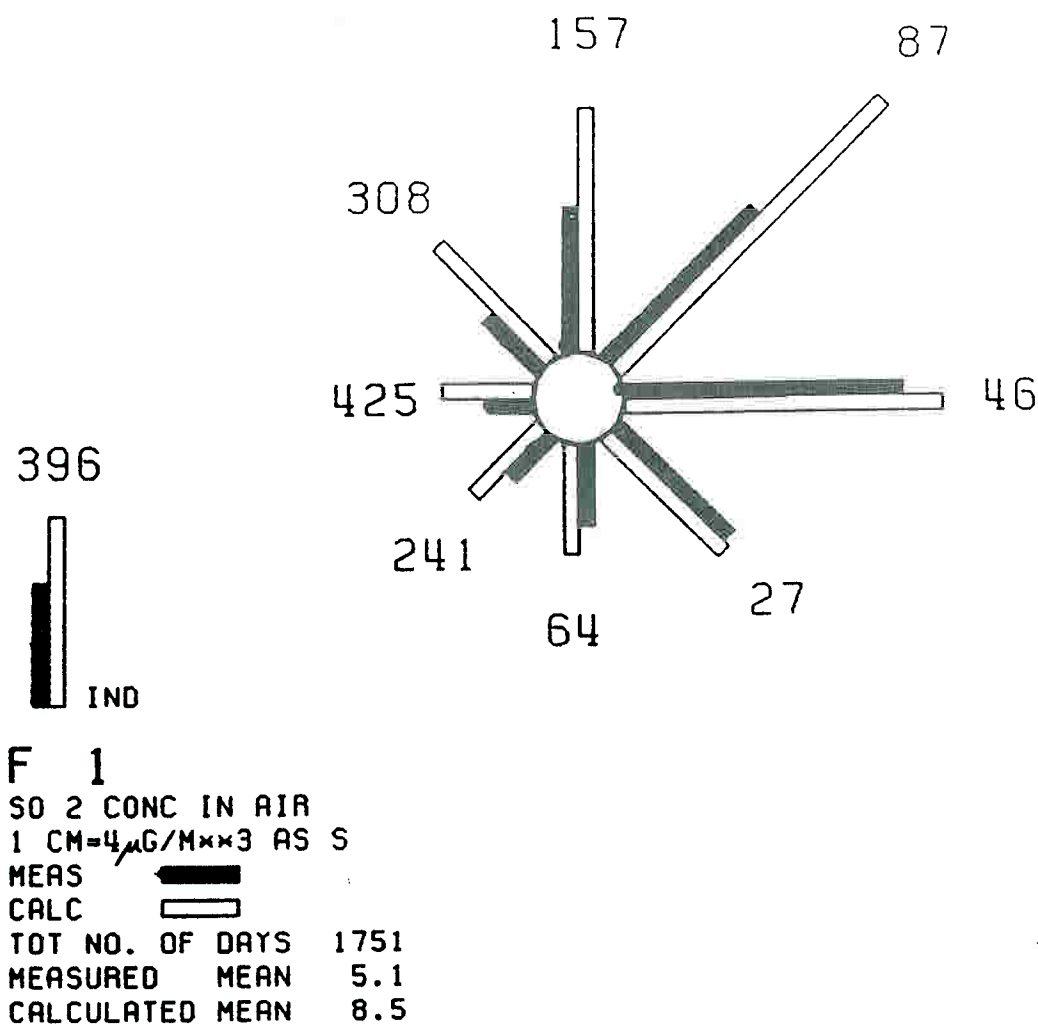


Figure 4(b). The number of trajectories coming from each of eight sectors (numerals), and the measured and calculated SO<sub>2</sub> concentrations for each sector, for station F1. The ninth "sector" marked IND represents the trajectories that were unclassifiable (Lehmhaus et al., 1985).

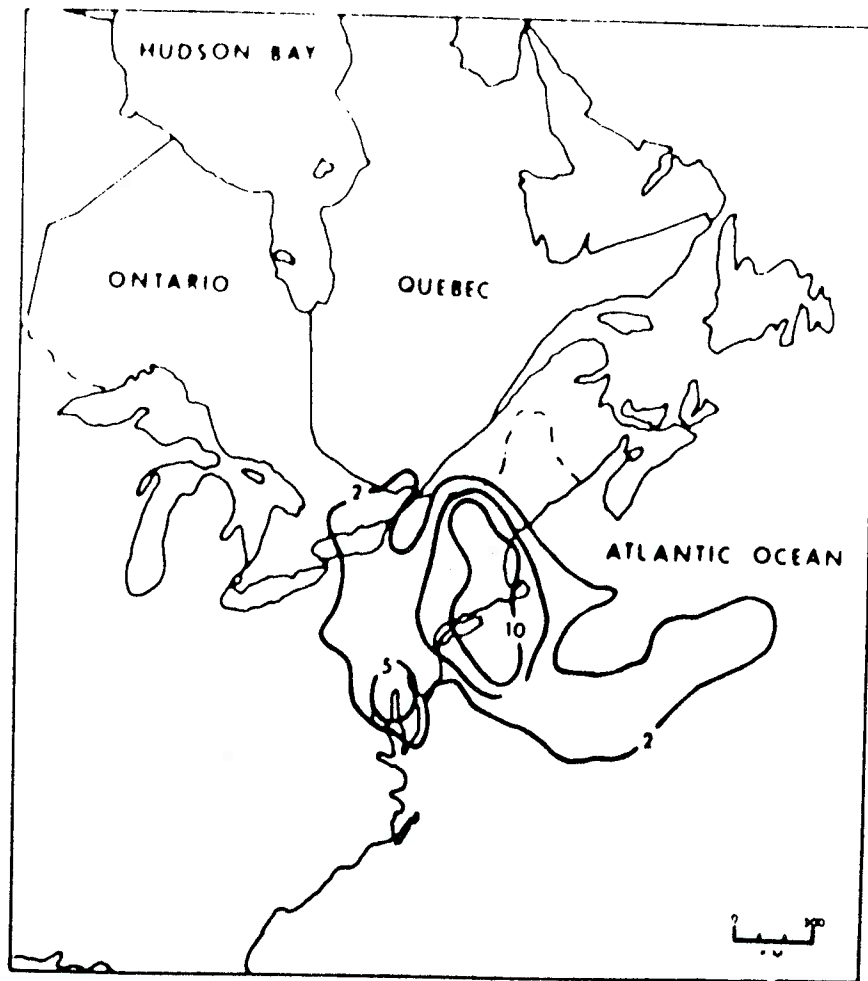


Figure 5. Isopleths of the percentage of times that back trajectories from Hubbard Brook, New Hampshire, crossed grid squares of 50 x 50 km<sup>2</sup> area. Percentages are for the uppermost 10% of the population of sulfate concentrations (Munn et al., 1984).

### 3.3 Vertical Motion

One of the problems that was discussed earlier was the lack of information on the vertical motion in the trajectory models used. To circumvent the problem, the output of forecast models that produce the vertical velocities is employed. These velocities come from the calculation of the convergence/divergence fields when the dynamic model begins its forecast in three dimensions. The possibility of using this information has only recently been recognized (Martin et al., 1984).

### 3.4 Field Experiments

The true path of a particle in the atmosphere can only be determined by in-situ measurements of atmospheric parameters, since measurement of all the needed values is impossible because of physical and financial reasons. However, special field experiments with wide spatial and temporal coverage could be useful instead in evaluating back trajectories. One such project is the Across North American Tracer Experiment (ANATEX), which is aimed at an evaluation of the meteorological transport over distances of 3000 km using an inert tracer. Figure 6 shows the configuration of the sampling array. The data collected at any of the sites could be used to verify back trajectories calculated from a given point.

## 4. SUMMARY

Since the advent of the computer over a decade ago, the single, back-trajectory analysis has proved useful in evaluating atmospheric chemistry data. This is witnessed by the large bibliography attached to this report. New techniques are slowly being evolved that will further the usefulness of this approach.

## 5. ACKNOWLEDGMENTS

Much of the collection of references was done during my stay at the Centre des Faibles Radioactivites. These references were supplemented, and a more extensive review was presented at the International Symposium on Acidic Precipitation, Muskoka, Canada, in September 1985.



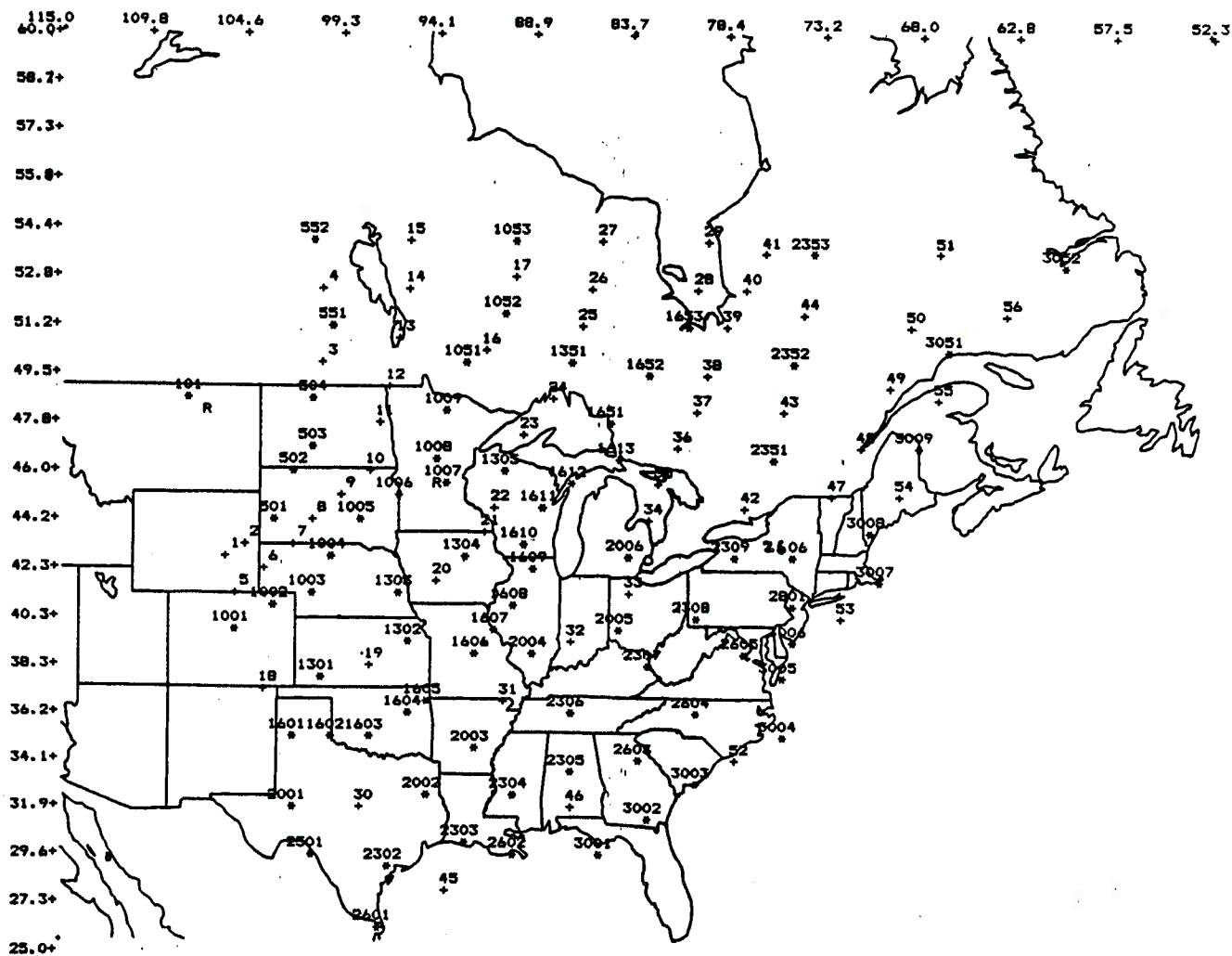


Figure 6. ANATEX source site locations at Glasgow, Montana (GGW), and at St. Cloud, Minnesota (STC), and the ground-level sampling network.

## 6. BIBLIOGRAPHY

February 27, 1987

The papers listed, with a few exceptions, deal only with single, back trajectories and their use in interpreting atmospheric chemistry data.

- Anlauf, K.G., M. Olson, H.A. Wiebe, and M.A. Lusi, 1980: Atmospheric transport of particulate sulphate and ozone into the Toronto region and its correlation with visibility. In Atmospheric Pollution 1980. M.M. Benarie (Ed.), Elsevier Scientific Publ., Amsterdam, 153-158.
- Anlauf, K.G., P. Fellin, H.A. Wiebe, and O.T. Melo, 1982: The Nanticoke Shoreline Diffusion Experiment, June 1978--IV. A. Oxidation of sulphur dioxide in a power plant plume. B. Ambient concentrations and transport of sulphur dioxide, particulate sulphate and nitrate, and ozone. Atmos. Environ. 16(3):455-466.
- Anlauf, K.G., J.W. Bottenheim, K.A. Brice, P. Fellin, H.A. Wiebe, H.I. Schiff, G.I. Mackay, R.S. Braman, and R. Gilbert, 1985: Measurement of atmospheric aerosols and photochemical products at a rural site in SW Ontario. Atmos. Environ. 19(11):1859-1870.
- Artz, R.A., and U. Dayan, 1986: Analysis and assessment of precipitation chemistry at Caribou, Maine. NOAA Tech. Memo. ERL ARL-143, NOAA Environmental Research Laboratories, Boulder, CO, 62 pp.
- Artz, R., R.A. Pielke, and J. Galloway, 1985: Comparison of the ARL/ATAD constant level and the NCAR isentropic trajectory analyses for selected case studies. Atmos. Environ. 19(1):47-63.
- Ashbaugh, L.L., 1983: A statistical trajectory technique for determining air pollution source regions. J. Air Pollut. Control Assoc. 33(11): 1096-1098.
- Asman, W.A.H., 1981: Meteorological interpretation of the chemical composition of rain-water at one measuring site. Water, Air, Soil Pollut. 16:159-175.
- Asman, W.A.H., and J. Slanina, 1980: Meteorological interpretation of the chemical composition of precipitation and some results of sequential rain sampling. Proceedings of the International Conference on the Ecological Impact of Acid Precipitation, March 11-14, 1980, Sandefjord, Norway, 140-141.
- Bamber, D.J., P.G.W. Healey, B.M.R. Jones, S.A. Penkett, A.F. Tuck, and G. Vaughan, 1984: Vertical profiles of tropospheric gases: Chemical consequences of stratospheric intrusions. Atmos. Environ. 18(9):1759-1766.
- Barrie, L.A., and R.M. Hoff, 1985: Five years of air chemistry observations in the Canadian Arctic. Atmos. Environ. 19(12):1995-2010.

- Barrie, L.A., H.A. Wiebe, K. Anlauf, and P. Fellin, 1980: The Canadian Air and Precipitation Monitoring Network APN. In Atmospheric Pollution 1980. M.M. Benarie (Ed.), Elsevier Scientific Publ., Amsterdam, 355-360.
- Barrie, L.A., R.M. Hoff, and S.M. Daggupaty, 1981: The Influence of mid-latitudinal pollution sources on haze in the Canadian Arctic. Atmos. Environ. 15(8):1407-1419.
- Barrie, L.A., K. Anlauf, H.A. Wiebe, and P. Fellin, 1984: Acidic pollutants in air and precipitation at selected rural locations in Canada. In Deposition Both Wet and Dry. B. Hicks (Ed.), Butterworth Publ., Boston, MA, 15-35.
- Billman Stunder, B.J., J.L. Heffter, and U. Dayan, 1986: Trajectory analysis of wet deposition at Whiteface Mountain: A sensitivity study. Atmos. Environ. 20(9):1691-1695.
- Bodhaine, B.A., J.J. DeLuise, J.M. Harris, P. Houmère, and S. Bauman, 1986: Aerosol measurements at the South Pole. Tellus 38(B):223-235.
- Bollinger, M.J., C.J. Hahn, D.D. Parrish, P.C. Murphy, D.L. Albritton, and F.C. Fehsenfeld, 1984: NO<sub>x</sub> measurements in clean continental air and analysis of the contributing meteorology. J. Geophys. Res. 89(D6):9623-9631.
- Brice, K.A., S.A. Penkett, D.H.F. Atkins, F.J. Sandalls, D.J. Bamber, A.F. Tuck, and G. Vaughan, 1984: Atmospheric measurements of Peroxyacetyl-nitrate (PAN) in rural south-east England: Seasonal variations winter photochemistry and long range transport. Atmos. Environ. 18(12):2691-2702.
- Brimblecombe, P., T. Davies, and M. Tranter, 1986: Nineteenth century black Scottish showers. Atmos. Environ. 20(5):1053-1057.
- Brosset, C., 1978: Water-soluble sulfur compounds in aerosols. Atmos. Environ. 12:25-38.
- Brosset, C., 1980: Types of transport episodes in northern Europe. In Aerosols: Anthropogenic and Natural, Sources and Transport. T.J. Kneip and P.J. Liroy (Eds.), Annals of New York Academy of Sciences, Vol. 338, 389-398.
- Buch, H., 1982: Meteorological indications of transport of air pollutants from the European continent to southern Scandinavia. Symposium on Meteorological Aspects of Air Pollution, Leningrad, USSR, March 1977, Vol I. Air Pollution Control Association, Philadelphia, I45-I59.
- Budd, W. W. 1986: Trajectory analysis of acid deposition data from the New Jersey pine barrons. Atmos. Environ. 20(12):2301-2306.
- Byrd, J.T., and M.O. Andreae, 1986: Concentrations and fluxes of tin in aerosols and rain. Atmos. Environ. 20(5):931-939.
- Castillo, R.A., J.E. Jiusto, and E. McLaren, 1983: The pH and ionic composition of stratiform cloud water. Atmos. Environ. 17(8):1497-1505.

- Chen, L., and R.A. Duce, 1983: The sources of sulfate, vanadium, and mineral matter in aerosol particles over Bermuda. Atmos. Environ. 17(10):2055-2064.
- Chester, R., E.J. Sharples, G.S. Sanders, and A.C. Saydam, 1984: Saharan dust incursion over the Tyrrhenian Sea. Atmos. Environ. 18(5):929-935.
- Chung, Y.-S., and H.V. Le, 1984: Detection of forest-fire smoke plumes by satellite imagery. Atmos. Environ. 18(10):2143-2151.
- Church, T.M., J.M. Tramontano, J.R. Scudlark, T.D. Jickells, J.J. Tokas, Jr., A.H. Knap, and J.N. Galloway, 1984: The wet deposition of trace metals to the Western Atlantic Ocean at the mid-Atlantic Coast and on Bermuda. Atmos. Environ. 18(12):2657-2664.
- Clark, T.L., and T.R. Karl, 1982: Application of prognostic meteorological variables to forecasts of daily maximum one-hour ozone concentrations in the Northeastern United States. J. Appl. Meteorol. 21(11):1662-1671.
- Clarke, J.F., and J.K.S. Ching, 1983: Aircraft observations of regional transport of ozone in the northeastern United States. Atmos. Environ. 17(9):1703-1712.
- Colbeck, I., and R.M. Harrison, 1985a: The frequency and causes of elevated concentrations of ozone at ground level at rural sites in north-west England. Atmos. Environ. 19(10):1577-1587.
- Colbeck, I., and R.M. Harrison, 1985b: The photochemical pollution episode of 5-16 July 1983 in north-west England. Atmos. Environ. 19(11):1921-1929.
- Covey, C., and P.L. Haagenson, 1984: A model of oxygen isotope composition of precipitation: Implications for paleoclimate data. J. Geophys. Res. 89(D3):4647-4655.
- Cox, R.A., A.E.J. Eggleton, R.G. Derwent, J.E. Lovelock, and D.H. Pack, 1975: Long-range transport of photochemical ozone in north western Europe. Nature 255(5504):118-121.
- Crabtree, J., and M. Kitchen, 1984: The long-range travel and dispersion of the plume from the Mount St. Helens Volcano. Atmos. Environ. 18(6):1073-1079.
- Danielsen, E.F., 1961: Trajectories: Isobaric, isentropic, and actual. J. Meteorol. 18:479-486.
- Danielsen, E.F., 1974: Review of trajectory methods. In Turbulent Diffusion in Environmental Pollution. F.N. Frenkiel and R.E. Munn (Eds.). Advance in Geophysics, Vol. 18B, Academic Press, New York, 73-94.
- Davidson, C.I., S. Santhanam, R.C. Fortmann, and M.P. Olson, 1985: Atmospheric transport and deposition of trace elements onto the Greenland ice sheet. Atmos. Environ. 19(12):2065-2081.
- Dayan, U., J.M. Miller, W.C. Keene, and J.N. Galloway, 1985: An analysis of precipitation chemistry data from Alaska. Atmos. Environ. 19(4):651-657.

- De Geer, L., 1977: Airborne short-lived radionuclides of unknown origin in Sweden in 1976. Science 198:925-927.
- De Pena, R.G., T.N. Carlson, J.F. Takacs, and J.O. Holian, 1984: Analysis of precipitation collected on a sequential basis. Atmos. Environ. 18(12): 2665-2670.
- De Pena, R.G., G.D. Rolph, J.F. Takacs, and J.O. Holian, 1986: Application of trajectory analysis to the assessment of local and long-range contributions to acidic deposition. Water, Air, Soil Pollut. 30:885-896.
- Dickerson, R.R., 1984: Measurements of reactive nitrogen compounds in the free troposphere. Atmos. Environ. 18(12):2585-2593.
- Dittenhoefer, A.C., and A.F. Ferullo, 1983: A dual-mode regional air back-trajectory model. In The Meteorology of Acid Deposition. Transactions, APCA Speciality Conference, Hartford, CT, Oct. 16-19, 1983, P.J. Samson (Ed.), Air Pollution Control Association, Philadelphia, 312-323.
- Dittenhoefer, A.C., and A.F. Ferullo, 1984: A comparison of Lagrangian precipitation statistics computed with two regional-scale atmospheric transport models. APCA Annual Meeting, 77th, San Francisco, CA, June 24-29, 1984. Air Pollution Control Association, Philadelphia, 1-12.
- Dittenhoefer, A.C., and A.F. Ferullo, 1985: A comparison of predicted and measured sulfate concentrations for precipitation events at Whiteface Mountain. APCA Annual Meeting, 78th, Detroit, MI, June 16-21, 1985. Air Pollution Control Association, Philadelphia, 1-15.
- Dollard, G.J., and M.H. Unsworth, 1983: Pollutant deposition as a result of interception of wind-driven cloud measurements made at a site in Northern England. In Precipitation Scavenging, Dry Deposition and Resuspension. H.R. Pruppacher et al. (Eds.), Elsevier, New York, 161-169.
- Dovland, H., and A. Semb, 1980: Atmospheric transport of pollutants. Proceedings of the International Conference on the Ecological Impact of Acid Precipitation, SNSF Project, Norway, 14-21.
- Draxler, R.R., 1983: Lagrangian meteorology and measurements of acidic precipitation at Washington, D.C. Atmos. Environ. 17(12):2525-2531.
- Duce, R.A., C.K. Unni, B.J. Ray, J.M. Prospero, and J.T. Merrill, 1980: Long-range atmospheric transport of soil dust from Asia to the tropical North Pacific: Temporal variability. Science 209:1522-1524.
- Dutkiewicz, V.A., J.A. Halstead, P.P. Parekh, A. Khan, and L. Husain, 1983: Anatomy of an episode of high sulfate concentration at Whiteface Mountain, New York. Atmos. Environ. 17(8):1475-1482.
- Eliassen, A., 1982: Aspects of Lagrangian air pollution modeling. In Air Pollution Modeling and Its Application III. Proceedings of the 13th International Technical Meeting, Ile des Embiez, France, Sept. 14-17, C. de Wispelleere (Ed.), Plenum Press, New York, 1-17.

- Eliassen, A., and J. Saltbones, 1983: Modeling of long range transport of sulphur over Europe: A two-year model run and some model experiments. Atmos. Environ. 17(8):1447-1473.
- Ellestad, T.G., 1980: Aerosol composition of urban plumes passing over a rural monitoring site. In Aerosols: Anthropogenic and Natural, Sources and Transport. T.J. Kneip and P.J. Liroy (Eds.), Annals of New York Academy of Sciences, Vol. 338, 202-218.
- Evans, G., P. Finkelstein, B. Martin, N. Possiel, and M. Graves, 1983: Ozone measurements from a network of remote sites. J. Air Pollut. Control Assoc. 33(4):291-296.
- Ferek, R.J., A.L. Lazrus, and J.W. Winchester, 1983: Elemental composition of aerosols collected with airborne cascade impactors. Atmos. Environ. 17(8):1563-1572.
- Ferm, M., U. Samuelsson, A. Sjodin, and P. Grennfelt, 1984: Long-range transport of gaseous and particulate oxidized nitrogen compounds. Atmos. Environ. 18(9):1731-1735.
- Fisher, B.E.A., 1983: A review of the processes and models of long-range transport of air pollutants. Atmos. Environ. 17(10):1865-1880.
- Fisher, B.E.A., 1984: The long-range transport of air pollutants--Some thoughts on the state of modelling. Atmos. Environ. 18(3):553-562.
- Fisher, B.E.A., and B.A. Callander, 1984: Mass balances of sulphur and nitrogen oxides over Great Britain. Atmos. Environ. 18(9):1751-1757.
- Fowler, D., and J.N. Cope, 1984: On the episodic nature of wet deposited sulphate and acidity. Atmos. Environ. 18(9):1859-1866.
- Fox, T.D., and J.D. Ludwick, 1976: Lead (Pb) concentrations associated with 1000 mb geostrophic back trajectories at Quillayute, Washington. Atmos. Environ. 10:799-803.
- Fuzzi, S., R.A. Castillo, J.E. Jiusto, and G.G. Lala, 1984: Chemical composition of radiation fog water at Albany, New York, and its relationship to fog microphysics. J. Geophys. Res. 89(D5):7159-7164.
- Galloway, J.N., A.H. Knap, and T.M. Church, 1983: The composition of Western Atlantic precipitation using shipboard collectors. J. Geophys. Res. 88(C15):10859-10864.
- Galvin, P.J., P.J. Samson, P.E. Coffey, and D. Romano, 1978: Transport of sulfate to New York State. Environ. Sci. Technol. 12(5):580-584.
- Gibson, T.L., P.E. Korosog, and G.T. Wolff, 1986: Evidence for the transformation of polycyclic organic matter in the atmosphere. Atmos. Environ. 20(8):1575-1578.
- Gjessing, Y.T., 1977: Episodic variations of snow contamination of an Arctic snowfield. Atmos. Environ. 11:643-647.



- Gordon, G.E., W.R. Piersen, J.M. Daisey, P.J. Troy, J.A. Cooper, J.G. Watson, and G.R. Cass, 1984: Consideration for design of source apportionment studies. Atmos. Environ. 18(8):1567-1582.
- Gotaas, Y., 1980: OECD program on long range transport of air pollutants-- Measurement from aircraft. In Aerosols: Anthropogenic and Natural, Sources and Transport. T.J. Kneip and P.J. Liroy (Eds.), Annals of New York Academy of Sciences, Vol. 338, 453-462.
- Gotaas, Y., 1982: Vertical distribution of sulphur in the atmosphere in a case of long range transport and the rate of transformation to sulphate. Atmos. Environ. 16(5):1043-1046.
- Graham, W.F., and R.A. Duce, 1982: The atmospheric transport of phosphorus to the Western North Atlantic. Atmos. Environ. 16(5):1089-1097.
- Haagensohn, P.L., A.L. Lazrus, Y.-H. Kuo, and G.A. Caldwell, 1985: A relationship between acid precipitation and three-dimensional transport associated with synoptic-scale cyclones. J. Clim. Appl. Meteorol. 24(9):967-976.
- Halter, B.C., and J.T. Peterson, 1981: On the variability of atmospheric carbon dioxide concentration at Barrow, Alaska during summer. Atmos. Environ. 15(8):1391-1399.
- Halter, B.C., J.M. Harris, and K.A. Rahn, 1985: A study of winter variability in carbon dioxide and Arctic haze aerosols at Barrow, Alaska. Atmos. Environ. 19(12):2033-2037.
- Hamrud, M., 1984: Lagrangian time scales connected with clouds and precipitation. Report CM-65, Dept. of Meteorology, Univ. of Stockholm, Report CM-65, 17 pp.
- Hanssen, J.E., J.P. Rambaek, A. Semb, and E. Steinnes, 1980: Atmospheric deposition of trace elements in Norway. Proceedings of the International Conference on the Ecological Impact of Acid Precipitation, March 11-14, 1980, Sandefjord, Norway, 116-117.
- Hansson, H.-C., B.G. Martinsson, and H.O. Lannefors, 1984: Long range aerosol transport in Southern Sweden: An example of multivariate statistical evaluation methodology. Nucl. Instrum. Methods Phys. Res. B3: 483-488.
- Harris, J.M., 1982: The GMCC Atmospheric Trajectory Program. NOAA Tech. Memo. ERL ARL-116, NOAA Environmental Research Laboratories, Boulder, CO, 30 pp.
- Harris, J.M., 1984: Trajectories during AGASP. Geophys. Res. Lett. 11(5): 453-456.
- Heffter, J.L., 1980: Air Resources Laboratories Atmospheric Transport and Dispersion Model (ARL-ATAD). NOAA Tech. Memo. ERL AL-81, NOAA Environmental Research Laboratories, Boulder, CO, 17 pp.

- Heffter, J.L., A.D. Taylor, and G.J. Ferber, 1975: A regional-continental scale transport, diffusion, and deposition model. NOAA Tech. Memo. ERL ARL-50, NOAA Environmental Research Laboratories, Boulder, CO, 28 pp.
- Heintzenberg, J., and S. Larssen, 1983: SO<sub>2</sub> and SO<sub>4</sub><sup>=</sup> in the Arctic: Interpretation of observations at three Norwegian Arctic-Subarctic stations. Tellus 35B:255-265.
- Heintzenberg, J., H.-C. Hansson, and H. Lannefors, 1981: The chemical composition of Arctic haze at Ny-Alesund, Spitsbergen. Tellus 33:162-171.
- Helms, L., and R. Jaenicke, 1985: Hidden information within series of measurements--Four examples from atmospheric science. J. Atmos. Chem. 3:171-185.
- Henderson, R.G., and Weingartner, 1982: Trajectory analysis of MAP3S precipitation chemistry data at Ithaca, NY. Atmos. Environ. 16(7):1657-1665.
- Henmi, T., and J.F. Bresch, 1985: Meteorological case studies of regional high sulfur episodes in the Western United States. Atmos. Environ. 19(11):1783-1796.
- Higuchi, K., and S.M. Daggupati, 1985: On variability of atmospheric CO<sub>2</sub> at Station Alert. Atmos. Environ. 19(12):2039-2044.
- Hitchcock, D.R., and M.S. Black, 1984: <sup>34</sup>S/<sup>32</sup>S evidence of biogenic sulfur oxides in a salt marsh atmosphere. Atmos. Environ. 18(1):1-17.
- Hobbs, P.V., and J.C. Yates, 1985: Atmospheric aerosol measurements over North America and the North Atlantic Ocean. Atmos. Environ. 19(1):163-179.
- Hoff, R.M., and N.B.A. Trwett, 1984: Ground-based measurements of Arctic haze made at Alert, N.W.T., Canada, during the Arctic Gas and Aerosol Sampling Project (AGASP). Geophys. Res. Lett. 11(5):389-392.
- Holt, B.D., R. Kumar, P.T. Cunningham, M. Bouchard, A. Englekemeir, S.A. Johnson, E.L. Neilsen, and J.D. Shannon, 1978: Regional Oxygen-18 variations in particulate sulfate and water vapor at three sampling sites about 100 km apart. Environ. Sci. Tech. 12(13):1394-1398.
- Hoppel, W.A., 1985: Ion-aerosol attachment coefficients, ion depletion, and the charge distribution on aerosols. J. Geophys. Res. 90(D4):5917-5923.
- Hoppel, W.A., J.W. Fitzgerald, and R.E. Larson, 1985: Aerosol size distributions in air masses advecting off the East Coast of the United States. J. Geophys. Res. 90(D1):2365-2379.
- Hoppel, W.A., G.M. Frick, and R.E. Larson, 1986: Effect of nonprecipitating clouds on the aerosol size distribution in the marine boundary layer. Geophys. Res. Lett. 13(1):125-128.
- Hov, O., S.A. Penkett, I.S.A. Isaksen, and A. Semb, 1984: Organic gases in the Norwegian Arctic. Geophys. Res. Lett. 11(5):425-428.



- Husain, T., and S.M. Khan, 1983: Air quality network design for Saudi Arabia using long-range transport model. Atmos. Environ. 17(5):921-926.
- Husain, L., J.S. Webber, E. Canelli, V.A. Dutkiewicz, and J.A. Halstead, 1984: Mn/V ratio as a tracer of aerosol sulfate transport. Atmos. Environ. 18(6): 1059-1071.
- Huschke, R.E. (ed.), 1959: Glossary of Meteorology. American Meteorological Society, Boston, MA, 638 pp.
- Isaac, G.A., J.W. Strapp, H.A. Wiebe, W.R. Leaitch, J.B. Kerr, K.G. Anlauf, P.W. Summers, and J.I. MacPherson, 1983: The role of cloud dynamics in redistributing pollutants and the implications for scavenging studies. In Precipitation Scavenging, Dry Deposition and Resuspension. H.R. Pruppacher et al. (Eds.), Elsevier, New York, 1-13.
- Isaac, G.A., W.R. Leaitch, J.W. Strapp, and K.G. Anlauf, 1986: Summer aerosol profiles over Algonquin Park, Canada. Atmos. Environ. 20(1):157-172.
- Israel, G.W., H.-W. Bauer, and K. Wengenroth, 1984: The Berlin Smog Project--Description and summary of results. Atmos. Environ. 18(10):2071-2088.
- Iversen, T., 1984: On the atmospheric transport of pollution to the Arctic. Geophys. Res. Lett. 11(5):457-460.
- Iversen, T., and E. Joranger, 1985: Arctic air pollution and large scale atmospheric flows. Atmos. Environ. 19(12):2099-2108.
- Iwasaka, Y., H. Minoura, and K. Nagaya, 1983: The transport and spacial scale of Asian dust-storm clouds: A case study of the dust-storm event of April 1979. Tellus 35B:189-196.
- Joranger, E., J. Schaug, and A. Semb, 1980: Deposition of air pollutants in Norway. Proceedings of the International Conference on the Ecological Impact of Acid Precipitation, March 11-14, 1980, Sandefjord, Norway, 120-121.
- Kadlecek, J., S. McLaren, N. Camarota, V. Mohnen, and J. Wilson, 1983: Cloud water chemistry at Whiteface Mountain. In Precipitation Scavenging, Dry Deposition and Resuspension. H.R. Pruppacher et al. (Eds.), Elsevier, New York, 103-113.
- Kadlecek, I., S. McLaren, V. Mohnen, B. Mossi, A. Kadlecek, and N. Camarota, 1984: Wintertime cloudwater chemistry studies. Atmospheric Sciences Research Center Report, 44 pp. + appendices.
- Kelly, T.J., R.L. Tanner, L. Newman, P.J. Galvin, and J.A. Kadlecek, 1984: Trace gas and aerosol measurements at a remote site in the Northeast U.S. Atmos. Environ. 18(12):2565-2576.
- Kemp, K., 1985: Receptor models for tracing long range transport. Workshop on Advancements in Air Pollution, Freiburg, West Germany, June 3-6, 1985. World Meteorological Organization, Geneva, Switzerland, 1-8.

- Kleinman, L.I., 1986: Photochemical of peroxides in the boundary layer. J. Geophys. Res. 91(D10):10889-10904.
- Kuo, Y.-H., M. Skumanich, P.L. Haagenson, and J.S. Chang, 1985: The accuracy of trajectory models as revealed by the observing system simulation experiments. Mon. Weather Rev. 113(11):1852-1867.
- Kurtz, J., and W.A. Scheider, 1981: An analysis of acidic precipitation in South-Central Ontario using air parcel trajectories. Atmos. Environ. 15(7):1111-1116.
- Kurtz, J., A.J.S. Tang, R.W. Kirk, and W.H. Chan, 1984: Analysis of an acidic deposition episode at Dorest, Ontario. Atmos. Environ. 18(2):387-394.
- Lannefors, H.O., T.B. Johansson, L. Granat, and B. Rudell, 1977: Elemental concentration and particle size distributions in an atmospheric background aerosol. Nucl. Instrum. Methods 142:105-110.
- Lannefors, H., H.-C. Hansson, and L. Granat, 1983a: Background aerosol composition in Southern Sweden--Fourteen micro and macro constituents measured in seven particle size intervals at one site during one year. Atmos. Environ. 17(1):87-101.
- Lannefors, H., J. Heintzenberg, and H.-C. Hansson, 1983b: A comprehensive study of physical and chemical parameters of the Arctic summer aerosol; Results from the Swedish Expedition Ymer-80. Tellus 35B:40-54.
- Lazrus, A.L., P.L. Haagenson, G.L. Kok, B.J. Huebert, C.W. Kreitzberg, G.E. Likens, V.A. Mohnen, W.E. Wilson, and J.W. Winchester, 1983: Acidity in air and water in a case of warm frontal precipitation. Atmos. Environ. 17(3):581-591.
- Lehmhaus, J., J. Saltbones, and A. Eliassen, 1985: Deposition patterns and transport sector analyses for a four-year period. EMEP/MSC-W Report 1/85, The Norwegian Meteorological Institute, Oslo, Norway.
- Lioy, P.J., P.J. Samson, R.L. Tanner, B.P. Leaderer, T. Minnich, and W. Lyons, 1980: The distribution and transport of sulphate "species" in the New York metropolitan area during the 1977 summer aerosol study. Atmos. Environ. 14:1391-1407.
- Lowenthal, D.H., and K.A. Rahn, 1985: Regional sources of pollution aerosol at Barrow, Alaska during winter 1979-80 as deduced from elemental tracers. Atmos. Environ. 19(12):2011-2024.
- Loye-Pilot, M.D., J.M. Martin, and J. Morelli, 1986: Influence of Saharan dust on the rain acidity and atmospheric input to the Mediterranean. Nature 321:427-428.
- Lurmann, F.W., B. Nitta, K. Ganesan, and A.C. Lloyd, 1984: Modeling potential ozone impacts from natural sources---III. Ozone modeling in Tampa/St. Petersburg, Florida. Atmos. Environ. 18(6):1133-1143.

- Mamane, Y., and T.G. Dzubay, 1986: Characteristics of individual particles at a rural site in the Eastern United States. J. Air Pollut. Control Assoc. 36:906-911.
- Martin, D., D. Cheymal, M. Imbard, and B. Skauss, 1984: Classement automatique des trajectoires du panade de l'eina. Third European Symposium, Physico-Chemical Behaviour of Atmospheric Pollution, Varese, Italy, Commission of European Committees, Brussels, Belgium, 509-525.
- Martinsson, B.G., H.-C. Hansson, and H.O. Lannefors, 1984: Southern Scandinavian aerosol composition and elemental size distribution characteristics dependence on air-mass history. Atmos. Environ. 18(10):2167-2182.
- Meinel, A.B., M.P. Meinel, and G.E. Shaw, 1976: Trajectory of the Mt. Augustine 1976 eruption and cloud. Science 193:420-422
- Merrill, J.T., R. Bleck, and L. Avila, 1985: Modeling atmospheric transport to the Marshall Islands. J. Geophys. Res. 90(D7):12927-12936.
- Merrill, J.T., R. Bleck, and D. Boudra, 1986: Techniques of Lagrangian trajectory analysis in isentropic coordinates. Mon. Weather Rev. 114:571-581.
- Miller, J.M., 1981a: A five-year climatology of five day back trajectories from Barrow, Alaska. Atmos. Environ. 15(8):1401-1405.
- Miller, J.M., 1981b: A five-year climatology of back trajectories from Mauna Loa Observatory, Hawaii. Atmos. Environ. 15(9):1553-1558.
- Miller, J.M., and J.M. Harris, 1985: The flow climatology to Bermuda and its implication for long-range transport. Atmos. Environ. 19(3):409-414.
- Miller, J.M., J.N. Galloway, and G.E. Likens, 1978: Origin of air masses producing acid precipitation at Ithaca, New York: A preliminary report. Geophys. Res. Lett. 5(9):757-760.
- Miller, J.M., D. Martin, and B. Strauss, 1987: A comparison of results from two trajectory models used to produce flow climatologies to the western Mediterranean. NOAA Tech. Memo. ERL ARL-151, NOAA Environmental Research Laboratories, Boulder, CO, 11 pp.
- Mueller, S.F., 1984: Analyses of errors in the estimation of wind vectors for medium/long-range air pollutant transport models. In The Meteorology of Acid Deposition. Transactions, APCA Speciality Conference, Hartford, CT, Oct. 16-19, 1983, P.J. Samson (Ed.), Air Pollution Control Association, Philadelphia, 274-289.
- Munn, R.E., G.E. Likens, B. Weisman, J.W. Hornbeck, C.W. Martin, and F.W. Bormann, 1984: A meteorological analysis of the precipitation chemistry event samples at Hubbard Brook (N.H.). Atmos. Environ. 18(12):2775-2779.
- Musold, G., and O. Lindqvist, 1983: Correlations between meteorological data and water-soluble sulphur compounds in five aerosols. Atmos. Environ. 17(7):1253-1260.

- Nieman, B.L., A.A. Hirata, and L.F. Smith, 1979: Application of a regional transport model to the simulation of multi-scale sulfate episodes over the Eastern United States and Canada. WMO Symposium on Long-Range Transport of Pollutants and Its Relation to General Circulation Including Stratospheric/Tropospheric Exchange Processes, Sofia, Bulgaria, Oct. 1-5, 1979. World Meteorological Organization, Geneva, 337-346.
- Norton, R.B., and J.F. Noxon, 1985: Dependence of stratospheric NO<sub>3</sub> upon latitude and season. J. Geophys. Res. 91(D5):5323-5330.
- Nyberg, A., 1977: On transport of sulfur over the North Atlantic. Meeting on Education and Training in Meteorological Aspects of Atmospheric Pollution and Related Environmental Problems, WMO. Research Triangle Park, NC, Jan.-Feb. 1977. World Meteorological Organization, Geneva, 1-5.
- Oblad, M., and E. Selin 1986: Measurements of elemental composition in background aerosols on the west coast of Sweden. Atmos. Environ. 20(7):1419-1432.
- Ockelmann, G., and H.W. Georgii, 1984: The distribution of sulfur dioxide over the Norwegian Arctic Ocean during summer. Tellus 36B:179-185.
- OECD (Organization for Economic Cooperation and Development), 1979: The OECD Programme on Long Range Transport of Air Pollutants: Measurements and findings. Second Edition, OECD, Paris, France.
- Ogen, J., and H. Rodhe, 1986: Measurements of the chemical composition of cloudwater at a clean air site in central Scandinavia. Tellus 38B:190-196.
- Olson, M.P., and K.K. Oikawa, 1980: Trajectory and concentration forecasts for the PEPE Project. Report AQRB-81-017-T, Atmospheric Environment Service, 29 pp.
- Olson, M.P., and K.K. Oikawa, 1983: A preliminary evaluation of the sulfate concentration forecasts for the PEPE Project. In The Meteorology of Acid Deposition. Transactions, APCA Speciality Conference, Hartford, CT, Oct. 16-19, 1983, P.J. Samson (Ed.), Air Pollution Control Association, Philadelphia, 367-386.
- Olson, M.P., K.K. Oikawa, and A.W. Macafee, 1978: A trajectory model applied to the long-range transport of air pollutants: A technical description and some model intercomparisons. Project 2.2.2, Atmospheric Environment Service, Canada, 24 pp.
- Ottar, B., 1978: An assessment of the OECD study on Long Range Transport of Air Pollutants (LRTAP). Atmos. Environ. 12:445-454.
- Ottar, B., and J.M. Pacyna, 1984: Sources of Ni, Pb, and Zn during the Arctic episode in March 1983. Geophys. Res. Lett. 11(5):441-444.
- Ottar, B., and J.M. Pacyna, 1986: Aircraft measurements of air pollution in the Norwegian Arctic. Atmos. Environ. 20(1):87-100.

- Ozkaynak, H., P.B. Ryan, and L.F. Bosart, 1982: Modeling of pollutant transport and removal during a regional sulfate episode. Water, Air, Soil Pollut. 18:157-171.
- Pack, D.H., J.E. Lovelock, G. Cotton, and C. Curthoys, 1977: Halocarbon behavior from a long time series. Atmos. Environ. 11:329-344.
- Pack, D.H., G.J. Ferber, J.L. Heffter, K. Telegadas, J.K. Angell, W.H. Hoecker, and L. Machta, 1978: Meteorology of long-range transport. Atmos. Environ. 12:425-444.
- Pacyna, J.M., and B. Ottar, 1985: Transport and chemical composition of the summer aerosol in the Norwegian Arctic. Atmos. Environ. 19(12):2109-2120.
- Pacyna, J.M., A. Semb, and J.E. Hanssen, 1984a: Emission and long-range transport of trace elements in Europe. Tellus 36B:163-178.
- Pacyna, J.M., V. Vitols, and J.E. Hanssen, 1984b: Size-differentiated composition of the Arctic aerosol at Ny-Alesund, Spitsbergen. Atmos. Environ. 18(11):2447-2459.
- Pacyna, J.M., B. Ottar, U. Tomza, and W. Maenhaut, 1985: Long-range transport of trace elements to Ny-Alesund, Spitsbergen. Atmos. Environ. 19(6): 857-865.
- Parekh, P.P., and L. Husain, 1982: Ambient sulfate concentrations and wind-flow patterns at Whiteface Mountain, New York. Geophys. Res. Lett. 9(1): 79-82.
- Parungo, F.P., C.T. Nagamoto, J. Rosinski, and P.L. Haagenson, 1986: A study of Maine aerosols over the Pacific Ocean. J. Atmos. Chem. 4:199-224.
- Peterson, J., K.J. Hanson, B.A. Bodhaine, and S.J. Oltmans, 1980: Dependence of CO<sub>2</sub>, aerosol, and ozone concentrations on wind direction at Barrow, Alaska during winter. Geophys. Res. Lett. 7(5):349-352.
- Pierson, W.R., W.W. Brachaczek, R.A. Gorse, Jr., S.M. Japar, and J.M. Norbeck, 1986: On the acidity of dew. J. Geophys. Res. 91(D3):4083-4096.
- Pitchford, M., and A. Pitchford, 1985: Analysis of regional visibility in the southwest using principal component and back trajectory techniques. Atmos. Environ. 19(8):1301-1316.
- Poirot, R.L., and P.R. Wishiniski, 1986: Visibility, sulfate and air mass history associated with the summertime aerosol in northern Vermont. Atmos. Environ. 20(7):1457-1469.
- Pratt, G.C., R.C. Hendricksen, B.I. Chevone, D.A. Christophersen, J.V. O'Brien, and S.V. Krupa, 1983: Ozone and oxide of nitrogen in the rural upper-Midwestern USA. Atmos. Environ. 17(10):2013-2023.
- Pratt, G.C., M.R. Coscio, and S.V. Krupa, 1984: Regional rainfall chemistry in Minnesota and West Central Wisconsin. Atmos. Environ. 18(1):173-182.

- Raatz, W.E., 1984: Tropospheric circulation patterns during the Arctic Gas and Aerosal Sampling Program (AGASP), March/April 1983. Geophys. Res. Lett. 11(5):449-452.
- Raatz, W.C., and G.E. Shaw, 1984: Long-range tropospheric transport of pollution aerosols into the Alaskan Arctic. J. Clim. Appl. Meteorol. 23:1052-1062.
- Rahn, K.A., and D.H. Lowenthal, 1984: Elemental tracers of distant regional pollution aerosols. Science 223:132-139.
- Raynor, G.S., and J.V. Hayes, 1982: Effects of varying air trajectories on spatial and temporal precipitation chemistry patterns. Water, Air, Soil Pollut. 18:173-189.
- Raynor, G.S., J.V. Hayes, and D.M. Lewis, 1983: Testing of the Air Resources Laboratories Trajectory Model on cases of pollen wet deposition after long-distance transport from known source regions. Atmos. Environ. 17(2):213-220.
- Reisinger, L.M., and T.L. Crawford, 1982: Interregional transport: Case studies of measurements versus model predictions. J. Air Pollut. Control Assoc. 32(6):629-633.
- Rindsberger, M., M. Magaritz, I. Carmi, and D. Gilad, 1983: The relation between air mass trajectories and the water isotope composition of rain in the Mediterranean Sea area. Geophys. Res. Lett. 10(1):43-46.
- Roberts, J.M., F.C. Fehsenfeld, S.C. Liu, M.J. Bollinger, C. Hahn, D.L. Albritton, and R.E. Sievers, 1984: Measurements of aromatic hydrocarbon ratios and NO<sub>x</sub> concentrations in the rural troposphere: Observation of air mass photochemical aging and NO<sub>x</sub> removal. Atmos. Environ. 18(11):2421-2432.
- Rodhe, H., C. Persson, and O. Akesson, 1972: An investigation into regional transport of soot and sulfate aerosols. Atmos. Environ. 6:675-693.
- Samson, P.J., 1978: Ensemble trajectory analysis of summertime sulfate concentrations in New York State. Atmos. Environ. 12:1889-1893.
- Samson, P.J., 1981: Trajectory analysis of summertime sulfate concentrations in the Northeastern United States. J. Appl. Meteorol. 19:1382-1394.
- Samson, P.J., and M.J. Small, 1984: Atmospheric trajectory models for diagnosing the sources of acid precipitation. In Modeling of Total Acid Precipitation Impacts. J.L. Schnoor (Ed.), Butterworth Publ., Boston, MA, 1-24.
- Samson, P.J., G.D. Thurston, and J.D. Spengler, 1984: Meteorological analysis of trace element data. APCA Annual Meeting, 77th, San Francisco, CA, June 24-29, 1984. Air Pollution Control Association, Philadelphia, 1-14.
- Shaw, G.E., 1980: Transport of Asian Desert aerosol to the Hawaiian Islands. J. Appl. Meteorol. 19:1254-1259.



- Shopauskas, K.K., S.Y. Armalie, A.V. Galvonaite, B.I. Gedraitis, A.Y. Girgzelis, K.K. Kuetkus, A.A. Milukaite, I.A. Shakalis, and D.A. Shopauskene, 1983: On the regularities in the formation of regional atmospheric pollution background in the Baltic Sea area. Proceedings of 2nd International Symposium on Integrated Global Monitoring, Tbilisi, USSR, Leningrad Gigrometeoizdat, Leningrad, USSR, 152-159.
- Slanina, J., and W.A.H. Asman, 1983: Tracking the source regions for wet deposition in the Netherlands by a combination of cluster analysis and meteorological interpretation. In Precipitation Scavenging, Dry Deposition and Resuspension. H.R. Pruppacher et al. (Eds.), Elsevier, New York, 239-247.
- Slanina, J., J.H. Baard, W.L. Zijp, and W.A.H. Asman, 1983a: Tracing the sources of the chemical composition of precipitation by cluster analysis. Water, Air, Soil Pollut. 20:41-45.
- Slanina, J., F.G. Romer, and W.A.H. Asman, 1983b: Investigation of the source regions for acid deposition in the Netherlands. In Acid Deposition. S. Beilke and A.J. Elshout (Eds.), Reidel, Dordrecht, The Netherlands, 129-141.
- Small, M.J., and P.J. Samson, 1983: Stochastic simulation of atmospheric trajectories. J. Clim. Appl. Meteorol. 22:266-277.
- Smith, F.B., and R.D. Hunt, 1978: Meteorological aspects of the transport of pollution over long distances. Atmos. Environ. 12:461-477.
- Sturges, W.T., and R.M. Harrison, 1986: Bromine: Lead ratios in airborne particles from urban and rural sites. Atmos. Environ. 20(3):577-588.
- Sykes, R.I., and L. Hatton, 1976: Computation of horizontal trajectories based on the surface geostrophic wind. Atmos. Environ. 10:925-934.
- Thurston, G.D., J.D. Spengler, and P.J. Samson, 1982: An assessment of relationships between regional pollution transport and trace elements using wind trajectory analysis. Specialty Conference on Receptor Models Applied to Contemporary Pollution Problems, Oct. 17-20, 1982. Air Pollution Control Association, Philadelphia, 1-11.
- Tragardh, C., 1982: Air chemistry measurements in the lower atmosphere over Sweden--Data evaluation. Atmos. Environ. 16(6):1451-1456.
- Vierkorn-Rudolph, B., K. Bachmann, B. Schwarz, and F.X. Meixner, 1984: Vertical profiles of hydrogen chloride in the troposphere. J. Atmos. Chem. 2:47-63.
- Viezee, W., W.B. Johnson, and H.B. Singh, 1983: Stratospheric ozone in the lower troposphere--II. Assessment of downward flux and ground-level impact. Atmos. Environ. 17(10):1979-1993.
- Walmsley, J.L., and J. Mailhot, 1983: On the numerical accuracy of trajectory models for long-range transport of atmospheric pollutants. Atmos.-Ocean 21(1):14-39.

- Warner, T.T., R.G. de Pena, J.F. Takacs, and R.R. Fizz, 1982: Application of a three-dimensional model to the calculation of trajectories: A case study. In Air Pollution Modeling and Its Application, III. Proceedings of the 13th International Technical Meeting, Ile des Embiez, France, Sept. 14-17, C. de Wispeleare (Eds.), Plenum Press, New York, 643-649.
- Warner, T., R. Fizz, and N.L. Seaman, 1983: A comparison of two types of atmospheric transport models--Use of observed winds versus dynamically predicted winds. J. Clim. Appl. Meteorol. 22:394-406.
- Whelpdale, D.M., 1978: Large-scale atmospheric sulfur studies in Canada. Atmos. Environ. 12:661-670.
- Whelpdale, D.M., and L.A. Barrie, 1982: Atmospheric monitoring network operations and results in Canada. Water, Air, Soil Pollut. 18:7-23.
- Wilson, J.W., V.A. Mohnen, and J.A. Kadlecsek, 1982: Wet deposition variability as observed by MAP3S. Atmos. Environ. 16(7):1667-1676.
- Wilson, J., R. Graham, and J. Robertson, 1983: Correlation of intrastorm sequential precipitation chemistry with storm meteorology. In Precipitation Scavenging, Dry Deposition and Resuspension. H.R. Pruppacher et al. (Eds.), Elsevier, New York, 229-238.
- Winkler, P., 1983: Trend development of precipitation-pH in Central Europe. In Acid Deposition. S. Beilke and H.J. Elshout (Eds.), Reidel, Dordrecht, The Netherlands, 114-122.
- Wolff, G.T., N.A. Kelly, and M.A. Ferman, 1982: Source region of summertime ozone and haze episodes in the Eastern United States. Water, Air, Soil Pollut. 18:65-81.
- Wolff, G.T., P.J. Liroy, R.E. Meyers, R.T. Cederwall, G.D. Wight, R.E. Pasceri, and R.S. Taylor, 1977: Anatomy of two ozone transport episodes in the Washington, D.C., to Boston, Mass., corridor. Environ. Sci. Technol. 11(5):506-510.
- Wolff, G.T., M.L. Morrissey, and N.A. Kelly, 1984: An investigation of the sources of summertime haze in the Blue Ridge Mountains using multivariate statistical methods. J. Clim. Appl. Meteorol. 23(9):1333-1341.
- Wolff, G.T., N.A. Kelly, M.A. Ferman, M.S. Ruthkosky, D.P. Stroup, P.E. Korsog, 1985a: Measurements of sulfur oxide, nitrogen oxides, haze and fine particles at a rural site on the Atlantic Coast. Report GMR-5062, Environmental Science Dept., General Motors Research Laboratories, Detroit, MI, 24 pp.
- Wolff, G.T., P.E. Korsog, N.A. Kelly, and M.A. Ferman, 1985b: Relationships between fine particulate species, gaseous pollutants, and meteorological parameters in Detroit. Atmos. Environ. 19(8):1341-1349.
- Wolff, G.T., M.S. Ruthkosky, D.P. Stroup, P.E. Korsog, M.A. Ferman, G.J. Wendel, and D.H. Stedman, 1986: Measurements of SO<sub>x</sub>, NO<sub>x</sub>, and aerosol species on Bermuda. Atmos. Environ. 20(6):1229-1239.