# **FULL SCALE STUDY OF**

# **DISPERSION OF STACK GASES**

**A Summary Report** 

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## FULL-SCALE STUDY OF DISPERSION OF STACK GASES

A Summary Report

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#### INTRODUCTION

### Description of the Study

During fiscal years 1958-1962 the Tennessee Valley Authority conducted an air pollution research project entitled "Full-Scale Study of Dispersion of Stack Gases" under the sponsorship of the Public Health Service. In this project advantage was taken of unique opportunities for full-scale appraisal of dispersion of air pollutants from large coalburning, steam-electric generating plants. Advantages offered for diffusion studies included: (1) large isolated sources where intermixture with extraneous pollutants is not significant; (2) complete plant operational data and emission rates; (3) sufficient fly ash emission to provide a visible plume aloft out to distances of 10-15 miles under meteorological conditions of special interest; (4) a helicopter equipped with special instruments for sampling and recording SO<sub>2</sub> concentrations, as well as extensive auxiliary instruments; (5) tower-mounted meteorological instruments for providing basic information on wind and temperature parameters; and (6) computer facilities for data analysis.

#### Work Plan

The initial work plan envisaged the compilation of sufficient field measurements for reasonably adequate definition of dispersion during inversion conditions, high wind conditions, and low wind conditions. While it is considered that dispersion was defined for inversion and high wind conditions, sampling techniques employed proved unsuitable for effective definition of dispersion during low wind and unstable conditions where excessive variability was presented by looping of the plume.

In addition to the primary studies to determine diffusion parameters, a limited investigation was made of plume rise or effective stack heights. A reasonably accurate estimate of effective stack height is required for useful application of diffusion parameters. Some corollary studies were considered desirable for appraising the validity or reliability of derived diffusion parameters. Principal among these corollary studies was an extensive investigation of the oxidation of SO<sub>2</sub> in the atmosphere after emission from the stack. Oxidation was studied with ground-based facilities and also in the plume at various distances and travel times, and under various weather conditions. In the course of this investigation interrelationships among SO<sub>2</sub>, H<sub>2</sub>SO<sub>1</sub>, and fly ash also were studied.

### Location of Field Studies

The Colbert Steam Plant (figures 1 and 2) located on the south bank of the Tennessee River 8 miles west of Tuscumbia, Alabama, was the site of most of the fieldwork. One flight used in the studies to define dispersion from a single stack was made at the Gallatin Steam Plant near Gallatin, Tennessee. The Colbert plant has four 200,000-kw units with four 300-foot stacks. The Colbert plant was selected for study because the plant is located in an area of reasonably flat topography, and the weather regime of the area includes a wide range of wind speeds and environmental temperature lapse rate conditions. The axis of the line of four stacks is oriented in a northwest-southeast direction (figure 3). This plant is located in a broad, relatively flat valley with the exception of a range of hills beginning about 3 miles southwest of the plant and extending 300-400 feet above the general valley floor.

#### Instrumentation

Helicopter and Auxiliary Equipment -- Equipment used in the Bell Model 47-D-1 helicopter (figure 4) included:

- 1. A portable Model 26-103 Titrilog, with Esterline-Angus recorder, for measuring continuous SO<sub>2</sub> plume concentrations. The Titrilog was positioned on a cushion mount between the pilot and the flight director.
- 2. A sample intake probe for the Titrilog extended about 12 inches forward from left bottom of cockpit canopy. A constant sample rate of about 1,000 cc per minute was maintained by utilizing the manifold vacuum of the helicopter.
- 3. A Model 8425 Cole-Parmer thermistor thermometer with interchangeable probe for taking ambient air vertical temperature profiles, as well as special temperatures in and out of the plume. The probe extended about 12 inches immediately forward from center bottom of the cockpit canopy.
- 4. A precision spring-wound clock for time documenting of flight sampling and observing.
- 5. A standard aircraft-type altimeter for indicating and maintaining desired heights aboveground.
- 6. A standard aircraft-type airspeed indicator for obtaining desired sampling airspeeds.
- 7. A secretarial-type voice recorder for recording temperature, height aboveground, and pertinent plume geometry observations.

Meteorological Facilities -- A fixed meteorological station, common to most TVA steam plants, was located 0.78 mile southeast of the Colbert

Steam Plant (figure 1). Station instrumentation included: (1) a 220-foot steel tower; (2) an anemograph model wind system for continuous recording of wind speed and wind direction at the 220-foot tower level; (3) a Brown temperature instrument, with 3-channel, sequential-type recorder, for continuous recording of ambient air temperature and wet-bulb depression at the 4-foot tower level and continuous temperature difference between the 220- and 4-foot tower levels; and (4) a standard cotton-region temperature shelter with hygrothermograph and maximum-minimum thermometers.

Wind profiles, using single theodolite, were obtained at a launching point near the fixed meteorological station. Hourly (later in the study, half hourly) pibals, using 10-gram ceiling balloon, were released during the sampling period for providing wind speed and wind direction profile data between surface and 2,000 feet.

Ambient air temperature profiles, using helicopter, were made at 100-foot vertical intervals from surface to 500 feet above the plume at a distance of about 1 mile from the plume. Temperature readings within the plume were considered unreliable, as the flight time through the plume generally was insufficient to allow probe readings to become stabilized to conditions encountered in transecting the plume, e.g., traversing through cooler ambient air into warmer plume air.

#### Aerial Sampling Plan

The procedural objective of the sampling plan was to provide adequate definition of SO<sub>2</sub> distribution in plume cross sections while allowing sufficient time for sampling and other essential measuring and observational activities. During inversion conditions, replicate flights were made across the plumes at the observed top and bottom elevations and

at subjectively selected centerline and quarter section elevations (figure 5). Cross-sectional flights were made at selected distances, usually 1/2, 3/4, 1, 3, and 5 to 10 miles from the plant. During high wind and neutral conditions, the flight plan was modified to take care of the greater variations in plume geometry than were found during inversion conditions. Replicate flights were made across the plume at the observed top of the plume and at successively lower 100- to 200-foot elevations to the bottom of the plume. Cross-sectional flights were made at selected distances of 1/2, 1, 2, and 3 miles from the plant. During high wind and neutral conditions, the SO<sub>2</sub> concentrations beyond 3 miles from the plant had diminished to such low levels that plume definition, based on SO<sub>2</sub> recorder registration, was not attainable.

#### Flight Speed and Sampling Rate

The flight speed for sampling was set at the minimum safe forward speed of the helicopter, 30 mph or 44 fps. Because of the excessive friction in the sample line and the fixed sample rate, it was not practical to attain isokinetic sampling at this airspeed. The SO<sub>2</sub> sample was drawn through a 0.075-inch-diameter orifice at the point of takeoff (figure 6), which provided a sample flow of about 20 fps.

Laboratory tests made under simulated field conditions showed that 90 percent of the average SO<sub>2</sub> concentration was being recorded. To compensate for this factor, the instrument factor applied to all Titrilog charts was increased by 10 percent.

## Data Collection and Analysis

The project consisted of four principal activities: (1) field sampling and observations, (2) reduction and consolidation of data, (3) data

analysis and formulation, and (4) summary of results and final reporting.

From table 1, which lists all helicopter flights in chronological order, it will be noted that from September 10, 1957, to October 28, 1960, 59 flights were made, totaling 149 hours of flight time. Classification of flights included: 15 for experimental developing and testing of sampling techniques and special instrumentation, 12 for successful dispersion definition of inversion plumes, 12 for successful dispersion definition of migh wind plumes, 4 for observing and recording plume rise, and 12 for studying SO<sub>2</sub> oxidation. Because of sudden changes in meteorological conditions, 2 flights were discontinued; 3 flights were of limited value because of voice recorder failure; and 1 flight was made for tracking a constant-volume tetroon.

To provide information on plant emission rates for correlation with dispersion data, average SO<sub>2</sub> concentrations were obtained by concurrent sampling of flue gas during all sampling flights. Sampling was obtained for successive 30-minute periods using the iodometric titration procedure. Representative samples of coal were taken concurrently from each plant unit in operation for analysis of average sulfur content of coal. Additional design and operational data included coal consumption per unit day, temperature of flue gas, exit velocity of flue gas, diameter and height of stack, etc. (table 2).

Following each flight all charts, recordings, data sheets, and observations were labeled. Later the data were abstracted, compiled, and tabulated or graphed for convenient use in the investigative studies of plume dispersion. Data made available for these dispersion studies included:

- 1. Average and axial SO2 concentration in cross section
- 2. Plume width and depth
- 3. Plume height aboveground
- 4. Average wind speed in plume
- 5. Vertical temperature gradient, <sup>o</sup>F./1,000 ft., in plume environmental area
- 6. Plume cross-sectional area
- 7. Standard deviation along the y and z axes
- 8. Plume direction in relation to line of stacks
- 9. SO<sub>2</sub> (flux) in cross sections, expressed in arbitrary units in a 1-foot plume segment

Following the establishment of suitable mathematical models, the consolidated data were applied initially in manual calculation of diffusion parameters. A program was developed in which the data were subjected to more extensive analysis through use of TVA's computer facilities.

Analysis was limited to generalized dispersion equations. Data are presented in sufficient detail and completeness for independent study and use by others working in this field.

#### Order of Reporting

To facilitate review and appraisal of extensive data analysis, tabulations, and illustrations, this report is arranged as follows.

Summary - Part I. Diffusion in Inversion Conditions

Summary - Part II. Diffusion in High Wind Neutral Conditions

Summary - Part III. Plume Rise

Summary - Part IV. Corollary Studies of SO2 Oxidation

Because of the volume of tables and figures in the analysis of data, only summaries of the four parts are being published. However, a limited number of copies of the data analysis which has been summarized will be made available to interested persons upon request.

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#### FULL-SCALE STUDY OF DISPERSION OF STACK GASES

#### A SUMMARY REPORT

#### Part I. Diffusion in Inversion Conditions

Diffusion in inversion conditions was defined on 12 sampling days. SO<sub>2</sub> registrations, from Titrilog recorder charts, were obtained during repetitive plume transections from 1/2 mile to 10 miles from the plant source (figure 7). From these sampling activities data shown in table 3 were collected and compiled.

#### Plume Geometry

Perimeter--The plume transection widths were determined from the length of the record trace registrations, e.g., chart speed of 1-1/2 inches per minute and flight speed of 44 feet per second. The depth or vertical distance of the plume was determined from altimeter readings at the top and bottom of the plume. A typical plume cross section developed for day 1 is shown in figure 8.

<u>Cross-Sectional Area--The cross-sectional area was determined</u>
from the formula:

Area = 
$$\frac{\pi \times \text{Width} \times \text{Depth}}{4}$$

l

#### Piume SO2 Data

Maximum Axial and Average SO<sub>2</sub> Concentrations--Maximum axial SO<sub>2</sub> concentration was determined from the flight transection through the center-line of the plume. Average SO<sub>2</sub> concentration was determined from planimetric analysis of the area under the SO<sub>2</sub> distribution curves.

SO<sub>2</sub> Flux-SO<sub>2</sub> flux, expressed in terms of cubic feet of SO<sub>2</sub> per linear foot of plume, was determined from the plume cross-sectional area and average SO<sub>2</sub> concentration in the plume section.

### Meteorological Data

Helicopter soundings provided data on vertical temperature gradient at 100-foot intervals from surface to heights well above the plume top (figure 9). Pilot balloon observations provided vertical wind direction and wind speed profile data from surface to 3,000 to 5,000 feet aboveground (figure 10). Supplementary data, including wind direction, wind speed, vertical temperature gradient, and wet bulb depression, were provided from the fixed meteorological station near the plant area.

### Plant Operational Data

Concurrently with field sampling activities, coal and flue gas samples were taken and coal consumption rates were noted, thus providing a measure of  $SO_2$  and heat emission rates.

#### Data Analysis

The objectives of the analysis of diffusion data were to express the results in terms of the mathematical model which best fits observed dispersion patterns and to develop coefficient values appropriate to the selected mathematical model.

Mathematical Diffusion Model--Examination of records or measured distribution of concentrations about the plume centerline in these studies indicates that Gaussian distribution is closely approximated. In a few instances the distribution is slightly skewed, and two maxima occurred in some instances when the plumes from separate stacks had not become uniformly

blended. Where Gaussian distribution exists, a plot of the distribution on normal probability paper yields a straight line. This test, applied to data taken on day 2 (figure 7), is illustrated in figure 11. While skewness exists in a few instances, most of the points have a very good straight-line fit.

The symmetry and results of the analytical test of distribution about the plume centerline are considered to justify the use of Gaussian distribution in mathematical analysis of diffusion of plumes in an inversion. From the basic equation for Gaussian distribution,

$$\overline{\underline{X}} = \overline{\underline{X}}_{\text{max}} e^{-1/2 \left(\frac{\underline{x}_{g}}{\sigma}\right)^{2}}$$

the following general equation for distribution at any single section of the plume in both horizontal and vertical directions is developed:

$$\overline{X} = \frac{Q}{2\pi\sigma_{y}\sigma_{z}u} \exp \left[-1/2\left(\frac{y^{2}}{\sigma_{y}^{2}} + \frac{z^{2}}{\sigma_{z}^{2}}\right)\right]$$

where the maximum concentration at the plume centerline is given by

$$\overline{\underline{X}}_{\text{max}} = \frac{Q}{2\pi\sigma_{\mathbf{v}}\sigma_{\mathbf{z}}\mathbf{u}}$$
 3a

Where the diffusion parameters  $C_y$ ,  $C_z$ , and m are constants under fixed meteorological conditions, the variability of the standard deviations,  $\sigma_y$  and  $\sigma_z$ , along the y and z axes has the following relationship with x, distance downwind from the source:

$$\sigma_{\mathbf{y}} = C_{\mathbf{y}}^{\mathbf{m}} \mathbf{y}$$

$$\sigma_{\mathbf{z}} = C_{\mathbf{z}}^{\mathbf{m}} \mathbf{z}$$

or when  $m_v = m_z$  or  $m_v + m_z = m$ 

$$\sigma_{\mathbf{y}} = C_{\mathbf{y}}^{\mathbf{x}}$$
 6

$$\sigma_{\mathbf{z}} = C_{\mathbf{z}} \mathbf{x}$$
 7

Then the following equations express the distribution at any point in the plume, along the x, y, and z axes.

$$\overline{X} = \frac{Q}{2\pi C_y C_{zux}^m} \exp \left[ -\frac{1}{2x^m} \left( \frac{y^2}{C_y^2} + \frac{z^2}{C_z^2} \right) \right]$$

or

$$\overline{X} = \frac{Q}{2\pi C_y C_z ux^{(m_y + m_z)}} \exp \left[ -\frac{1}{2} \left( \frac{y^2}{C_y C_z x^{2m_y}} + \frac{z^2}{C_z C_z x^{2m_z}} \right) \right] 8a$$

General dispersion formulas 8 and 8a correspond to the Sutton equation for distribution in an elevated plume when the diffusion coefficients  $C_y$  and  $C_z$  are multiplied by  $\sqrt{2}$  and the value of m or  $(m_y + m_z)$  is set at 2-n.

Standard Deviation--Since  $SO_2$  concentration was measured with a continuous recording instrument at a known uniform rate of speed through the plume, the curves show actual distribution in relation to time and distance. The area under the curves representing  $SO_2$  distribution along the line of flight can be obtained by integrating equation 2 between the limits of  $\overset{*}{\smile} \infty$  which give

$$\sigma = \frac{\text{Area}}{X_{\text{max}} \sqrt{2\pi}}$$

The area under the curve is equal to the base (width or depth in feet) times the average height (average SO<sub>2</sub> concentration in ppm along the axis).

Computer Program - Diffusion Parameters--A FORTRAN program was developed for calculation of  $\sigma_y$ ,  $\sigma_z$ , m, m, m, m, c, c, and  $\overline{X}$  from parameters measured in the plume and from SO<sub>2</sub> emission rates. These values were determined as follows.

$$\sigma_{y} = \frac{\text{(Plume width)(Average SO_2 concentration)}}{\text{(Peak SO_2 concentration)}} \sqrt{2\pi}$$

$$\sigma_{z} = \frac{\text{(Plume depth)(Average SO_2 concentration)}}{\text{(Peak SO_2 concentration)}} \quad 11$$

$$m = \log\left(\frac{\underline{x}_1}{\underline{\overline{x}}_2}\right) \log\left(\frac{x_1}{x_2}\right)$$

$$m_y = \log\left(\frac{\sigma_{y_1}}{\sigma_{y_2}}\right) / \log\left(\frac{x_1}{x_2}\right)$$
 13

$$m_z = \log\left(\frac{\sigma_{z_1}}{\sigma_{z_2}}\right) / \log\left(\frac{x_1}{x_2}\right)$$

 $C_y$  and  $C_z$  from equations 4, 5, 6, and 7, and  $\overline{X}$  at 36 points in the quarter section of the plume illustrated by figure 12 from equation 8a.

The computed values of SO<sub>2</sub> concentrations determined for field conditions from the measured values of standard deviation and axial and average SO<sub>2</sub> concentrations at points along the y and z axes (figure 12) were plotted for each section sampled, and are illustrated for day 2 in figure 13. From these plots, the values of average SO<sub>2</sub> concentration and plume widths and depths were calculated. With the exception of two cases, the calculated values of plume widths and depths agreed closely with

field-measured values (figure 14). The ratio of the measured axial concentration to the average SO<sub>2</sub> concentration (figure 15) was approximately 2.33:1. For the computed values, the ratio was 2.18:1 (figure 16).

Parameters  $m_y$  and  $m_z$ --Average values of the parameters  $m_y$  and  $m_z$  for each day (table 3) were determined for field conditions from the ratio of the standard deviation along each axis to the distance. Average  $m_z$  values were much less than average  $m_y$  values, indicating that the rate of diffusion is much greater along the y axis and that separate m values for the y and z axes would be required.

Since an appreciable range in the values of parameters  $m_y$  and  $m_z$  was evident for the 12 sampling days, values considered to be the most representative of the more accurate dispersion data were grouped into four ranges of decreasing stability (table 4). Variation of these parameters with stability is illustrated in figure 17.

Coefficients  $C_y$  and  $C_z$ --Values of  $m_y$  and  $m_z$  (table 4) were determined for the various ranges of stability and were used to develop estimates of  $C_y$  and  $C_z$  for each cross section for days grouped according to stability, from which the average values of  $C_y$  and  $C_z$  (table 4) were obtained. The variation of these coefficients with stability is shown in figure 17.

While the parameters  $m_y$ ,  $m_z$ ,  $C_y$ , and  $C_z$  developed to this point have a limited value for application to general diffusion problems, they should provide reasonably accurate estimates of diffusion during the full range of inversion conditions at distances of more than 1 mile from the source.

### Modification of Line-Source Parameters to Point-Source Parameters

To provide broader application for use in general diffusion problems, all measured plume dimensional data and axial and average plume concentrations were modified to approximate a single stack or point source. Modification was derived from the relationships between the wind direction and the observed horizontal and vertical spread of the plume width and depth with a line of stacks (figure 18).

Analysis of data measured in a plume from a line of two to four stacks affirms that both the plume width and plume depth vary with respect to the relation between plume direction and direction of the line of stacks. Thus, wind direction, per se, may effectuate variations in the diffusion coefficients for identical meteorological parameters. Minimum widths of plumes from a line of stacks occur when the azimuth direction of the plume is the same as the alignment axis of the stacks; and conversely, maximum widths occur when the plume direction is at right angles to the alignment axis of the stacks. On the other hand, minimum depths of plumes occur when the azimuth direction of the plume is at right angles to the alignment axis of the stacks; and conversely, maximum depths occur when the plume direction is the same as the stack alignment axis. With the addition of one stack, at a stack separation distance  $\triangle$  (figure 19), the plume width is increased by an increment equivalent to distance  $\triangle$  when the plume direction is at right angles to the alignment axis of the stacks. As the plume direction varies from 90° to 0° from the stack alignment, the increase in plume width decreases from  $\triangle$  to zero. The data plotted in figure 18 indicate that for the Colbert Steam Plant the magnitude of increase in plume width is approximately four times the decrease in plume depth. Thus the process of

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converting line-source plume dimensions, plume width and depth, to dimensions representative of one stack is approximated by:

- 1. Reducing the width by  $(n-1) \triangle \sin \theta$
- 2. Reducing the depth by  $(n-1) \frac{\triangle}{4} \cos \theta$  where n = Number of stacks
  - $\theta$  = Degrees plume direction varies from stack alignment
  - $\Delta$  = Linear separation between stacks

When a normal distribution exists for both point- and line-source emissions, axial and average concentrations for the single stack point source should have the same ratio as axial and average SO<sub>2</sub> concentrations for a line source. This relation is confirmed from field data with 2-, 3-, and 4-stack sources (figure 15). On the basis of this relation, point-source concentrations were estimated from line-source data by using the following formula.

SO<sub>2</sub> Concentration (point source) =

Plume cross-sectional area (line) x Line-source SO2 concentration | Number of units on line

Thus the line-source values were adjusted according to this single stack point-source formula. These data were then processed through the computer in the same manner as the original field data for compiling estimates of  $m_v$ ,  $m_z$ ,  $C_v$ , and  $C_z$  for a point source (table 5).

Parameters  $m_y$  and  $m_z$ —In the same manner as for line sources, the values of  $m_y$  and  $m_z$  were determined from the combined values considered to be most representative of the more accurate dispersion data and were grouped into four ranges of decreasing stability from which the best estimates of the average values (table 6) were obtained. The variation of the  $m_y$  and  $m_z$  parameters with stability is shown in figure 20.

Coefficients  $C_y$  and  $C_z$ --Values of  $m_y$  and  $m_z$  (table 6) derived for the four ranges of stability were used to develop  $C_y$  and  $C_z$  values for each cross section according to classification of stability. Average  $C_y$  and  $C_z$  coefficients (table 6) were then determined for each of the groups of stability. Variation of these coefficients with stability is given in figure 20.

#### Modification of Point-Source Parameters for General Diffusion Problems

As the number of stacks is increased beyond one, the plume widths and depths are increased (figure 19) in accordance with the following equations.

$$\Delta_{y_n} = (n-1) \triangle \sin \theta$$
 16

$$\Delta_{\mathbf{z_n}} = (\mathbf{n-1}) \frac{\Delta}{4} \cos \theta$$
 17

Where:

n = Number of stacks

 $\triangle_{y_n}$  = Increase along y axis

 $\Delta_{\mathbf{z}_n}$  = Increase along  $\mathbf{z}$  axis

 $\triangle$  = Distance between stacks

 $\theta$  = Degrees plume off line of stacks

These relationships provided the means for estimating the rate of change in parameters  $m_y$  and  $m_z$  (figure 21) and  $C_y$  and  $C_z$  (figure 22) in relation to  $\Delta_{y_n}$  and  $\Delta_{z_n}$ , as a point or single stack source is enlarged to a line of two or more stacks.

To utilize the point-source parameter for line-source determinations, the following steps are necessary.

- 1. Determine the increase in plume widths,  $\triangle_{y_n}$ , and plume depths,  $\triangle_{z_n}$ , due to additional stacks and plume direction from equations 16 and 17.
- 2. Determine the line-source parameters  $m_y$ ,  $m_z$ ,  $C_y$ , and  $C_z$  by multiplying the corresponding point-source parameters by their respective ratios shown by the curves in figures 21 and 22 for the increase in plume widths,  $\Delta_{y_n}$ , and plume depths,  $\Delta_{z_n}$ .

Table 7 shows the line-source parameters developed for each day from the best estimate of the point-source parameters, and gives the field values of the axial concentration shown in line B, table 8.

# <u>Comparison - Field Concentrations and Concentrations Derived from Calculated Diffusion Parameters</u>

Table 8 summarizes the results from field sampling and mathematical analysis.  $\underline{A}$  values in this table are axial concentrations measured in the plume, and  $\underline{B}$  values are concentrations calculated from point-source parameters modified to simulate line-source field conditions.

Reasonably good agreement exists between measured and calculated values for days 1, 2, 3, and 9. The relation of measured A values and calculated B values is evident in the following table. Data from these four

Paraant

	Mea	asured A Dista		centratio	n
Day	1/5	3/4	1	2	x
1	117	106	90	83	77
2	101	95	95	92	86
3	82	79	104	110	-
9		-	90	_	85

days suggest that use of the calculated diffusion coefficients in the general dispersion equation should yield concentrations within ±25 percent of the actual field values and that SO<sub>2</sub> distribution should closely approximate the actual plume geometry or dimensions.

Greater differences in measured and calculated concentrations were evident for the other sampling days. These differences primarily are attributed to use of diffusion coefficients based on a single fixed temperature gradient or stable condition to calculate diffusion over a 2- to 2-1/2-hour period. Actually, stability over such an interval may undergo significant changes which affect both measured concentrations and diffusion rates. Simultaneous sampling at two or more sections would be required to overcome this problem.

## Part II. Diffusion in High Wind and Neutral Conditions

Field instrumentation and sampling procedures used during high winds and neutral conditions were similar to those used in the study of dispersion during inversion conditions. Because of the greater variance in the plume configurations, in comparison to the relatively stable and unvarying conditions of the inversion plume, the aerial sampling plan was modified slightly. Additional horizontal flights across the plume from top to bottom at successively lower elevations were made to determine the cross-sectional areas. Some flights were made in the plume between 1/2 mile and 3 miles from the source and along paths parallel to the x axis of the plume. The purpose of these flights was to compare the SO<sub>2</sub> distribution along the vertical axes with the concentrations determined from the cross-sectional flights. In most cases the plume was widely dispersed in both horizontal

and vertical directions within a relatively short distance or travel time. Because of the larger cross sections and the increase in time required for sampling each cross section, the maximum sampling distance from the plant was restricted to 3 miles. Generally, SO<sub>2</sub> concentrations at this distance had diminished to such a low level that plume definition from recorded charts no longer was discernible.

#### Plume Geometry

Constant shifting of the plume along the vertical axis, primarily attributed to cyclical variations in wind speed, interfered with precise definition of plume cross sections. As a result of continuous vertical shifting, the measured depth of the plume (table 9), based on the difference in elevation between the bottom of plume and the first higher elevation when no SO<sub>2</sub> was recorded, may have varied slightly with the true depth of the plume. Such discrepancies were dependent upon vertical shifting and the progression status of the sampling flight. The transit widths, determined from the recorder chart speed and flight speed, indicated that during the sampling of a single cross section it could be possible to traverse the plume centerline (figure 23) more than once. The plume width used in this analysis (figure 23) was the maximum width determined by several transection flights through a representative plume segment which was selected to best define the plume depth.

#### Plume SO2 Data

Maximum and average SO<sub>2</sub> concentrations (table 9) associated with each cross section were determined from SO<sub>2</sub> charts (figure 24), as they were in the analysis of data taken in inversion conditions. The SO<sub>2</sub> distribution

determined from these Titrilog charts was similar to that observed in inversion conditions and approximated a Gaussian distribution. The average ratio of maximum to average concentrations (figure 25) was about 2.00:1.

To estimate SO<sub>2</sub> flux, a true cross section was required. This condition was unobtainable because of plume fluctuation during the sampling period (figure 23).

#### Meteorological Data

The same type of meteorological information obtained under inversion conditions was obtained for high wind and neutral conditions.

In the 11 sampling days average temperature change with elevation (figure 26) closely approximated the adiabatic lapse rate from the surface to elevations well above the plume. The average vertical temperature gradient in the plume during all sampling periods varied from -4° F. to -6.5° F. per 1,000 feet.

Average wind speed in the plume section (table 9) based on pibal observations (figure 23) varied from 8 to 23 miles per hour.

### Data Analysis

The same approach and procedures as those used in the analysis of data for inversion conditions were followed in the analysis of field data for high wind and neutral conditions (table 9), with the exception of the determination of standard deviation about the z axis. Here the value of  $\sigma_z$  was determined indirectly from  $\sigma_y$  through the general dispersion equation 3a. This indirect determination is considered preferable to the determination of  $\sigma_z$  from the variable SO<sub>2</sub> distribution along the z axis (figure 27). While there is only 7-percent variability in the average of values by the two methods, significant variability exists in individual sections.

To facilitate the application to general problems, values of the parameters for line-source field conditions (table 9) were modified as outlined in Part I to approximate values representative of a single stack point source (table 10). Thus field values of standard deviation were modified to approximate the standard deviation along the y and z axes and were used in calculation of the diffusion parameters m, m<sub>y</sub>, m<sub>z</sub>, C<sub>y</sub>, and C<sub>z</sub> for a single stack point source (table 10). Finally, the values of parameters developed for a single stack point source were modified to approximate line-source field sampling conditions. The modified parameters were used to calculate SO<sub>2</sub> concentrations for comparison with field-measured values.

Parameters m, m<sub>y</sub>, and m<sub>z</sub>--The values of m, m<sub>y</sub>, and m<sub>z</sub> determined for each sampling day show a range of 0.500 to 0.847 with an average of 0.686 for m<sub>y</sub> and a range of 0.800 to 0.968 with an average of 0.759 for m<sub>z</sub> (table 10). The relationship of  $\sigma_y$  values to  $\sigma_z$  values (figure 28) indicates that the average values of m<sub>y</sub> and m<sub>z</sub> probably were equal. The maximum average values for m<sub>y</sub> and m<sub>z</sub> (line 4, figure 29) for all days were about 0.75.

Diffusion Coefficients  $C_y$  and  $C_z$ —The values of  $C_y$  and  $C_z$  are determined from formulas 4 and 5 for each plume cross section for each day, using values of  $m_y = m_z = 0.80$ , 0.75, and 0.70 (table 11). The values 0.80 and 0.70 selected to bracket the average value 0.75 show a significant variation among the days. Analyses of  $C_y$  and  $C_z$  values plotted against average wind speed (figures 30, 31, and 32) reveal a slight scatter; also the relationship reveals a trend for decreasing  $C_y$  and  $C_z$  values with increasing wind speed. The best estimate of  $C_y$  and  $C_z$  values at wind speed intervals of 2 miles per hour is given in table 12. The variation of the coefficients  $C_y$ ,  $C_z$ , and  $C_z$  for  $m_y = m_z = 0.75$  to wind speed is given in figure 33.

### Modification of Point-Source Parameters for General Problems

As the number of stacks is increased, plume widths and plume depths are increased (figure 19) in the amount of  $\triangle_{y_n}$  and  $\triangle_{z_n}$  in accordance with equations 16 and 17. These increases in plume widths and depths were used in estimating the change in  $m_y$  and  $m_z$  (figure 34) and  $C_y$  and  $C_z$  (figure 35) in relation to  $\triangle_{y_n}$  and  $\triangle_{z_n}$  as plant units were added.

To utilize the point-source parameters for line-source determinations, the following steps are necessary.

- 1. Determine the increase in plume widths,  $\triangle_{y_n}$ , and plume depths,  $\triangle_{z_n}$ , due to additional stacks and plume direction from equations 16 and 17.
- 2. Determine the line-source parameters  $m_y$ ,  $m_z$ ,  $C_y$ , and  $C_z$  by multiplying the corresponding point-source parameters by their respective ratios shown by the curves in figures 34 and 35 for the increase in plume widths,  $\Delta_{y_n}$ , and plume depths,  $\Delta_{z_n}$ .

The point-source values for  $m_y = m_z = 0.75$  (table 12) modified to represent line-source field conditions outlined above are shown in table 13 for each cross section sampled.

#### Summary of Results

Final  $m_y$ ,  $m_z$ ,  $C_y$ , and  $C_z$  values estimated for field line-source conditions (table 13) were employed in equation 8a to calculate axial concentrations. Field-measured and calculated concentrations are summarized in table 14 and are plotted in figure 36. Good agreement is indicated in most of the sections sampled. The variation for individual points is attributed to application of a steady-state mathematical model to the relatively variable plume pattern.

#### Part III. Plume Rise

This over-all project was concerned primarily with the investigation of diffusion rates in steam plant smoke plumes. While detailed data on plume rise were obtained (on a limited basis) during each plume diffusion sampling period, the scope of data collections, restricted by project objectives, was not considered extensive enough to support a comprehensive study designed specifically to improve present analytical procedures for determining plume rise.

Plume rise observed during each sampling day for temperature inversion and lapse conditions is shown in figures 37 and 38. During inversion conditions maximum plume rise usually was attained within the first 1/2 mile from the emission source. During lapse conditions most of the plume rise was attained in the first 1/2 mile, but a slight continuous increase in plume height occurred beyond this point.

The average observed relationships of plume rise to wind speed and stability for cases occurring within the first 2-mile section of the plume during inversion conditions and within the 1/2- and 1-mile sections during lapse conditions are given in figures 39 through 43. The wide variation of plume rise with wind speed during inversion conditions (figure 39) probably is due to the variance in stability for comparable wind speeds.

#### Part IV. Corollary Studies of SO2 Oxidation

In the analysis of data taken in inversion conditions, some variability of SO<sub>2</sub> flux was noted in progressive plume cross sections. The apparent consistency in a trend of decreasing SO<sub>2</sub> flux with distance, along with published information on SO<sub>2</sub> oxidation, indicated a need for study

of oxidation of SO<sub>2</sub> in a steam plant plume. The following principal phases of the SO<sub>2</sub> oxidation studies were defined.

- Develop equipment and techniques for the collection
   of representative samples of flue gas and fly ash from
   steam plant ducts or stacks.
- 2. Collect and analyze sufficient samples of flue gas and fly ash to establish the relative proportions and concentrations of SO<sub>2</sub> and SO<sub>3</sub>, as well as pertinent physical and chemical characteristics of fly ash.
- 3. Develop facilities for controlled dilution and cooling of flue gas simulating atmospheric dispersion and cooling.
- 4. Develop instrumentation for evaluating changes in sulfur oxides and fly ash subjected to controlled dilution and cooling.
- 5. Modify instrumentation and techniques developed in the preceding step for study of sulfur oxides and fly ash in the dispersed plume.
- 6. Collect and analyze sufficient plume samples to establish the relative proportions of SO<sub>2</sub> and SO<sub>3</sub>.
- 7. Interpret and analyze data and observations.

In steps 1 through 4, flue gas and fly ash samples were taken at ground level from the duct section connecting the mechanical fly ash collectors and the induced draft fan, or from a dilution chamber adjoining this duct. A report on these studies follows.

#### Oxidation Studies in Duct and Dilution Chamber

Gas samples were collected from the duct for  $SO_2$  and  $SO_3$  analyses. The tests considered to yield the most reliable values for 30-minute average concentrations of  $SO_2$  and  $SO_3$  are tabulated below.

SO<sub>2</sub>-SO<sub>3</sub> Analysis

Colbert Steam Plant Flue Gas

Date	SO3 ppm as SO2	SO <sub>2</sub>
7/10/59 7/15/59 7/21/59 7/21/59 7/21/59 7/29/59 7/29/59	42 66 18 12 17 17	2317 1886 2388 2241 2192 2258 2312

These data suggest that only 1 to 2 percent of sulfur in coal exists as SO<sub>3</sub> in flue gas at the Colbert plant.

Fly ash samples were collected directly from the duct for size consist, identification of principal physical characteristics, and chemical analysis.

Concurrent with the collection of samples directly from the duct, flue gas was aspirated into a large dilution chamber (figure 44) where it was held for 2 to 5 hours. Gas samples were taken from the chamber for SO<sub>2</sub> and SO<sub>3</sub> analyses and fly ash samples were taken for other chemical analysis. While exceptions and unexplained events were noted, the data suggest an increase in SO<sub>2</sub> oxidation with holding time. In these tests, data from the Titrilog and autometer indicate that 10 to 30 percent of SO<sub>2</sub> in the trailer was oxidized in 2 to 3 hours (table 15).

Using the difference in recorded values for the same sample on an autometer and a Titrilog as a measure of oxidation is not a recommended

technique. However, data taken with and without a filter for removal of acid aerosol indicate that the technique has some validity. In figure 45 where acid aerosol is removed, Titrilog and autometer values are approximately equal. In figure 46 where the acid is not removed and is presumably registered on the autometer, a significant difference exists.

In a second series of tests, oxidation in the dilution chamber was evaluated by direct sampling of SO<sub>2</sub> and SO<sub>3</sub>. While results are random (table 16), the extent of oxidation was similar to that derived indirectly from autometer-Titrilog data and was as high as 30 percent on two tests.

#### Fly Ash Studies

Fly ash may influence the oxidation of SO<sub>2</sub> in coal flue gases because of its catalytic and nucleating properties. Therefore, concurrent with the estimate of SO<sub>2</sub> oxidation in the dilution chamber, studies were made of the physical and chemical characteristics of fly ash. Principal data are presented in table 17.

Because of deposition in the transfer line and aspirator system, fly ash transfer from the duct to the dilution chamber was only 25-75 percent efficient. However, a number of factors were indicative of oxidation of SO<sub>2</sub>.

- l. Large crystals of aluminum sulfate were identified on the aluminum foil electrostatic precipitator liners. This is attributed to the reaction of precipitated  $\rm H_2SO_4$  aerosol with the aluminum foil lining.
- 2. Sulfate content of fly ash increased from about 2.5 percent in the duct to 5 to 10 percent in the chamber. This increase is attributed to deposition of sulfuric acid mist or aerosols on fly ash particles.

3. The pH of fly ash generally decreased with holding time in the chamber--from about neutral in the duct to a low of 4.5.

Peripheral fly ash studies included size analysis, study of acid-alkaline characteristics, and microchemical-petrographic studies. While the information disclosed by these studies is of general interest from an operational and industrial hygiene viewpoint, its relation to SO<sub>2</sub> oxidation is indirect. However, the acid-alkaline characteristics of fly ash are of particular interest.

Fly ash taken in the duct remained essentially neutral for all dilutions. However, the pH of fly ash from the chamber increased with increasing dilution and time. Investigation suggested that the low pH with minimum dilution was due to rapid solution of acid aerosol on the fly ash surface. As dilution was increased, the solubility of calcium oxides increased, and a part of the initial acidity was neutralized with resultant higher pH values. Data on these tests are provided in table 18. Figure 47 illustrates a change in the pH of a fly ash sample taken from the mechanical collector from 4.4 to 12.0 in about 2 hours. The heterogenous acid and alkaline fly ash characteristics were clearly demonstrated when colorimetric acid indicator on fly ash samples revealed random distribution of acid subsamples. Microchemical-petrographic tests indicated that the minor component responsible for the alkalization of fly ash is a dehydrated form of calcium sulfate, probably calcium oxide-calcium hydroxide formed during brief retention in the fire chamber.

### Studies in Power Plant Plumes

While the SO<sub>2</sub> oxidation studies were beset with numerous problems and limitations, some useful information was disclosed; and, through elimination, the most satisfactory sampling and analytical procedures were identified. Thus, for sampling in the plume, filter paper was used to collect the acid aerosol, and SO<sub>2</sub> was collected in a subsequent series H<sub>2</sub>O<sub>2</sub> scrubber (figure 48). SO<sub>3</sub> concentrations were based on the sulfate content of the filter.

Samples were taken in an inversion plume on 8 days from 1/2 mile to 10 miles from the steam plant. Plume travel time from the source ranged from 5 to 108 minutes. Data from the 8 sampling days are compiled in tables 19 and 20. Data for 5 sampling days (August 2, September 2, and October 14, 26, and 28, 1960) do not indicate significant oxidation of SO<sub>2</sub>. Oxidation of this magnitude, 1 to 3 percent, approximates that determined for undiluted hot flue gas. Relatively high oxidation, 8 to 55 percent, was observed on 3 days (May 3, August 19, and October 11, 1960).

More tests of this type are needed for confirmation of these limited data. However, the data derived with these sampling and analytical techniques suggest that in periods of 1 to 2 hours, oxidation of SO<sub>2</sub> in the plume may range from almost none to 50 percent. Moisture within the plume or ambient strata apparently is the factor which exerts predominant control over the rate of oxidation. When relative humidity is below 70 percent, oxidation is very slow. Atmospheric moisture above this level but at less than saturation conditions produces a maximum initial rate of oxidation.

A primary reason for initiating a study of atmospheric oxidation of SO<sub>2</sub> was to appraise its effect on diffusion parameters based on data

from the Titrilog which does not record acidified  $SO_2$ . Analysis suggests that  $SO_2$  oxidation may not have significantly affected the values of diffusion parameters during most days when field dispersion measurements were taken. On most days the observed variation may be ascribed to limitations of the field sampling procedure rather than to the sampling instrument. Also, meteorological criteria established for dispersion measurements excluded high humidity conditions favorable to a high rate of  $SO_2$  oxidation.

APPENDIXES

# TENNESSEE VALLEY AUTHORITY Division of Health and Safety

and

PUBLIC HEALTH SERVICE Division of Air Pollution

### APPENDIX A

TABLES

Chattanooga, Tennessee August 1964

Table 1
HELICOPTER FLIGHTS

Date	Dura Hr.	ation Min.	Type of Flight	Date	Dura Hr.	Min.	Type of Flight
9/10/57 9/11/57 9/12/57 9/17/57 9/23/57 9/23/57 9/25/57 9/26/57 9/26/57 10/4/57 10/10/57 10/10/57 10/10/57 10/11/57 10/15/57 3/28/58 3/21/58 4/2/58 4/10/58 4/11/58 4/11/58 4/11/58 4/16/58 3/27/59	Hr. 03310434314443200032324312263	Min. 45 20 0 15 350 55 0 0 0 5 20 5 5 20 5 30 15 30 15 5 20 15 30 15 30 15 5 20 15 30 15	Experimental Experimental Experimental Experimental Experimental Experimental Inversion Experimental Experimental Experimental Experimental Experimental High wind High wind High wind High wind Experimental High wind Experimental High wind High wind Experimental High wind Experimental High wind Experimental High wind Experimental High wind	Date  4/3/59 4/7/59 4/17/59 4/20/59 4/23/59 4/24/59 1/26/60 2/24/60 2/24/60 8/12/60 8/12/60 8/19/60 10/11/60 10/13/60 10/25/60 10/25/60 10/28/60			of Flight  High wind Inversion  Voice recorder out  SO2 - SO3 Plume observation SO2 - SO3 Tetroon SO2 - SO3 Tetroon SO2 - SO3 and tetroon
4/1/59 4/2/59	5 0	0 25	High wind High wind				release

Table 2
PLANT DESIGN AND OPERATIONAL DATA

	Stea	am Plant
	Colbert	Gallatin
Total rated capacity, kw	720,000	450,000
Number of units	4	2
Unit rated capacity, kw	180,000	225,000
Unit capability, kw	200,000	250,000
Total capability, kw	800,000	500,000
Number of stacks	4	1
Spacing of stacks, feet, approximately	100	-
Height of stack, feet	300	500
Diameter of stack, feet	16.5	25
Exit velocity of flue gas, fps	47	44
Temperature of flue gas, OF.	290	290
Coal consumption per unit day, tons	1,800	1,928
Sulfur content of coal, percent	1-3.5	3.2

Table 3 SUMMARY - PRINCIPAL DATA, BY SECTIONS (FIELD-MEASURED VALUES)

<u>Day</u>	Dist.	Time	Units Operating	Plume Dir.a	Elev. (Ft.)b	Pl Width (Ft.)	ume Depth (Ft.)	SO <sub>2</sub> (ppr Max.		Wind Speed (mph)	Temp. Gradient OF./1,000' Plumed	Standa Deviat (Ft.	ion	(Dimens	Diffusion sionless) mz	Parameter  (Ft. 2)  Cy	(Ft. 2)	Plume Cross- Sectional Area (Sq. Ft.)	SO <sub>2</sub> Emission Rate (cfs)	Measured SO <sub>2</sub> Flux (Cu. Ft. per Lin. Ft.)
1	1/2 3/4 1 2 6 9-1/2	0636 0649 0700 0716 0741 0801	2, 3, & 4 2, 3, & 4	84 84 84 84 84	680 680 680 660 660 680	1660 1856 2188 2837 3441 3848	460 460 420 400 440 480	18.1 16.0 16.3 12.3 7.9 6.1	8.5 7.5 7.1 5.5 3.4 2.3	9.8 9.9 10.0 10.0 10.1 10.0	8.3 7.4 7.4 6.8 5.0 4.0	299 334 394 511 619 693	83 83 76 72 79 86	0.346	0.053	19.629 19.059 20.355 20.774 17.213	54.655 53.492 48.239 44.051 45.598	599,426 670,202 721,384 890,818 1,188,521 1,449,926	41.8	2.6 2.5 2.6 2.5 2.0 1.7
2	1/2 3/4 1 2 8	0655 0704 0645 0740 0752	2, 3, & 4 2, 3, & 4 2, 3, & 4 2, 3, & 4 2, 3, & 4	3 3 3 3	720 820 7 <sup>1</sup> 40 800 1080	1026 1041 1192 1373 2173	530 540 575 575 625	28.7 24.1 20.1 12.9 5.6	12.8 9.8 7.4 5.9 3.0	7.8 7.6 7.3 6.8 6.6	4.3 4.6 2.6 3.0 4.0	181 183 210 242 382	93 95 101 101 110	0.262	0.081	23.041 20.951 22.299 21.436 23.543	49.021 48.452 50.322 47.565 46.284	426,867 441,280 538,039 619,738 1,066,128	47.4	2.8 2.2 2.2 1.9 1.6
3	1/2 3/4 1 2	0623 0631 0638 0657	2, 3, & 4 2, 3, & 4 2, 3, & 4 2, 3, & 4	15 15 15 15	670 680 680 600	875 1328 2037 2248	550 525 515 560	25.4 21.1 13.9 9.0	11.6 10.1 6.4 4.7	12.0 12.0 11.9 11.6	9.3 7.8 7.4 8.4	168 255 391 432	106 101 99 108	0.849	0.069	0.208 0.224 0.269 0.165	61.494 56.974 54.747 56.930	377,781 547,302 823,508 988,221	48.3	2.2 2.8 2.7 2.3
14	1/2 3/4 1 2 8	0637 0647 0657 0707 0737	2, 3, & 4 2, 3, & 4 2, 3, & 4 2, 3, & 4 2, 3, & 4	39 39 39 39 39	720 750 720 750 <b>7</b> 40	1117 1418 1494 2233 4904	550 650 720 750 820	17.2 16.9 15.2 9.6 4.4	6.9 6.5 5.9 3.8 1.6	13.2 13.4 13.7 14.0 12.8	4.2 3.5 2.6 2.0 0.7	173 220 232 346 760	85 101 112 116 127	0.493	0.197	3.545 3.690 3.376 3.577 3.964	18.058 19.813 20.763 18.765 15.643	482,265 723,535 844,409 1,314,679 3,156,705	46.3	1.7 2.4 2.5 2.5 2.6
5	1/2 3/4 1 2 1C-1/2	0638 0646 0655 0703 0809	1, 2, 3, & 4 1, 2, 3, & 4 1, 2, 3, & 4 1, 2, 3, & 4 1, 2, 3, & 4	89 89 89 89	750 750 700 720 850	1388 1554 1766 2113 5116	850 750 870 825 1050	24.4 18.2 19.1 16.8 4.5	10.7 8.6 8.0 6.0 1.8	9.4 9.2 9.0 8.9 9.5	5.7 2.9 2.3 1.8 0.8	232 260 295 353 854	142 125 145 138 175	0.380	0.152	11.589 11.131 11.321 10.407 13.400	42.764 35.390 39.292 33.648 33.145	926,143 914,918 1,206,090 1,368,432 4,216,863	69.2	5.0 4.0 4.8 4.1 3.8
6	1/2 3/4 1 2 9	0805 0757 0748 0734 0638	1, 2, 3, & 4 1, 2, 3, & 4 1, 2, 3, & 4 1, 2, 3, & 4 1, 2, 3, & 4	67 67 67 67 67	1300 1350 1050 1000 560	1147 1403 1464 1750 5749	780 900 950 950 540	20.1 15.2 14.5 16.8 13.2	9.2 7.0 6.4 7.3 5.0	7.6 7.5 7.4 7.2 6.6	-2.6 -2.3 -2.7 -2.1 +8.7	200 244 255 305 1000	136 157 165 165 94	0.432	0.199	6.650 6.809 6.284 5.571 9.538	28.317 30.154 29.925 26.067 11.006	702,308 991,220 1,091,778 1,305,063 2,437,001	70.1	3.3 3.5 4.5 4.8 6.1
7	1/2 3/4 1 2 9	0712 0720 0731 0744 0618	1, 2, 3, & 4 1, 2, 3, & 4 1, 2, 3, & 4 1, 2, 3, & 4 1, 2, 3, & 4	74 74 74 74 74	950 1050 850 900 550	1660 1509 2082 2384 4904	700 900 900 900 290	20.1 19.1 22.4 20.5 11.2	11.2 10.1 10.9 6.6 4.5	6.8 6.4 5.9 6.1 9.5	2.7 4.9 5.3 4.4 20.7	300 287 377 432 888	127 163 163 163 52	0.428	o <b>.3</b> 85	10.286 8.272 9.607 8.182 8.833	6.107 6.705 6.002 4.595 0.821	912,170 1,119,096 1,470,933 1,684,296 1,116,396	55.4	5.1 5.3 8.0 5.6 2.5
8	1/2	0740 0727 0708	1, 2, 3, & 4 1, 2, 3, & 4 1, 2, 3, & 4	74 74 74	850 850 1000	1841 2792 4104	680 700 820	25.1 6.6 4.6	7.7 3.1 1.8	7.3 8.0 8.3	3.1 3.3 0.6	287 436 640	106 109 128	0.579	0.136	3.009 3.061 3.009	36.294 33.963 36.294	982,726 1,534,204 2,641,745	74.7	3.8 2.4 2.4
9	1 4 <b>-1/</b> 2	0648 0810	1, 2, & 3 1, 2, & 3	69 69	650 <b>7</b> 00	2501 5009	510 4 <b>7</b> 0	8.0 3.3	2.8 1.4	13.8 11.5	5.3 2.3	388 776	79 73	0.461	0.0	7.469 7. <b>4</b> 69	79.000 73.000	1,001,510	33.6	1.4 1.3
10	1 2	0815 0845	1 & 2 1 & 2	0	, 1100 1800	2022 2983	900 1200	2.2 0.9	1.2 0.4	14.0 11.0	-2.1 -2.4	400 591	1 <b>7</b> 8 238	0.563	0.419	3.204 3.204	4.901 4.901	1,428,543 2,809,986	39.4	0.9 0.6
11	1/2 5-3/4	0659 0 <b>7</b> 38	1 & 3 1 & 3	71 71	550 550	1724 3903	500 480	12.1 7.5	5.4 3.0	9.6 8.2	*	288 652	84 80	0.335	0.0	20.640 20.640	84.000 80.000	676,670 1,470,650	37.1	1.9 2.2
12	1/2 5	0706 0746	1 & 2 1 & 2	33 33	600 500	1188 3 <b>3</b> 92	440 335	17.5 9.1	8.7 3.8	13.7 11.5	15.7 11.0	217 621	81 61	0.457	0.0	5.944 5.944	81.000 61.000	410,335 892,011	29.4	1.8 1.7

aDegrees off line of stacks.

bElevation of flight where maximum concentration was recorded.

CAverage wind speed along elevation of maximum concentration.

dFrom approximately bottom to top of plume.

\*No data.

Table 4 BEST ESTIMATE OF AVERAGE  $m_y$ ,  $m_z$ ,  $c_y$ , AND  $c_z$  BY RANGES OF STABILITY (FIELD-MEASURED VALUES)

	Temperature Gra		,		C C-			
Group	Range	Av.	$m_{\underline{\mathbf{y}}}$	$m_Z$	$\overline{c}^{\lambda}$	$\frac{C_{\mathbf{Z}}}{C_{\mathbf{Z}}}$		
, 1	13.4	13.4	.396	0	11.067	76.500		
. 5	(6.5 to 8.2)	7.4	.430	.090	8.667	48.323		
- 3	(2.3 to 3.8)	3.0	.458	.174	5.710	23.976		
24	(-0.2 to -2.3)	-1.3	.531	.273	3.255	14.606		

Table 5

SUMMARY - PRINCIPAL DATA, BY SECTIONS

FOR SINGLE STACK POINT SOURCE

			* .						Diffusion	n Parameters		
Day	Dist. (M:.)	Pij Width (Ft.)	Depth (Ft.)	SO <sub>2</sub> (pr Max.		σ <sub>y</sub> (Ft.)	σ <sub><b>z</b></sub> (Ft.)	(Dimensi	ionless) mz	(Ft. 2)	(Ft. $\frac{m_Z}{c_Z}$ )	SO <sub>2</sub> Emission (cfs)
1 .	1/2 3/4 1 2 6	1461 1657 1989 2638 3242	455 455 415 395 435	6.9 6.0 6.0 4.5 2.8	3.3 2.8 2.6 2.0 1.2	263 298 358 475 584	32 82 75 71 78	0.381	0.054	13.111 12.731 13.708 13.971 11.307	53.696 52.538 47.316 43.155 44.692	13.9
2	1/2 3/4 1 2 8	1016 1031 1182 1363 2163	460 490 525 525 575	11.1 8.9 7.4 4.7 2.0	4.8 3.6 2.7 2.2 1.1	179 181 208 240 381	84 86 92 92 101	0.264	0.089	22.280 20.246 21.561 20.712 22.789	41.529 `41.004 42.751 40.182 38.970	15.8
3	1/2 3/4 1 2	823 1276 1985 2196	502 477 467 512	9.8 8.1 5.2 3.4	4.5 3.9 2.4 1.8	158 245 381 422	96 92 90 98	0.883	0.067	0.151 0.163 0.197 0.118	56.457 52.647 50.513 52.494	16.1
Į,	1/2 3/4 1 2 8	991 1292 1368 2107 4778	511 611 681 711 781	6.9 6.6 5.8 3.6 1.6	2.8 2.5 2.3 1.4 0.6	154 200 212 327 741	79 95 106 110 121	0.529	0.209	2.385 2.500 2.276 2.433 2.648	15.225 16.820 17.673 15.867 13.063	15.4
5	1/2 3/4 1 2 10-1/2	1088 1254 1466 1813 4816	849 749 869 824 1049	7.8 5.7 5.8 4.9	3.4 2.7 2.4 1.8 0.5	182 209 245 303 804	142 125 145 138 175	0.447,	0.152	5.388 5.162 5.321 4.828 6.108	42.764 35.390 39.292 33.648 33.145	17.3
6	1/2 3/4 1 2 9	871 1127 1188 1474 5473	751 871 921 921 511	6.9 4.9 4.6 5.1 3.7	3.2 2.3 2.0 2.2 1.4	152 196 207 256 952	131 152 160 160 89	0.506	0.206	2.831 2.974 2.716 2.366 4.113	25.846 27.586 27.367 23.725 9.681	17.5
7	1/2 3/4 1 2 9	1372 1221 1794 2096 4616	679 879 879 879 269	6.3 6.1 6.7 6.0 3.2	3.5 3.0 3.3 1.9	248 221 325 379 835	123 159 159 159 159	0.524	0.396	3.989 2.874 3.635 2.947 2.951	5.422 5.969 5.326 4.047 0.687	13.9
8	1/2 1 2	1553 2504 3816	659 679 799	7.7 1.9 1.3	2.4 0.9 0.5	253 408 622	107 111 130	0.649	0.140	1.524 1.567 1.524	35.385 33.303 35.385	18.7
9	1 4-1/2	2314 4822	492 452	3.0 1.2	1.0 0.5	359 747	76 70	0.487	0.0	5.515 5.515	76.000 70.000	11.2
10	1 2	2022 2983	900 1200	2.2 0.9	1.2	400 591	178 238	0.563	0.419	3.204 3.204	4.901 4.901	39.4
11	1/2 5 <b>-3/</b> 4	1629 3808	492 472	6.6 3.9	2.9 1.5	272 636	82 <b>7</b> 9	0.348	0.0	17.563 1 <b>7.</b> 563	82.000 79.000	18.6
12	1/2 5	1134 3338	419 31 <sup>4</sup>	9.7 4.9	4.8 2.1	208 611	77 57	0.468	0.0	5.210 5.210	77.000 57.000	14.7

Table 6 BEST ESTIMATE OF  $m_{\rm y}$ ,  $m_{\rm z}$ ,  $C_{\rm y}$ , AND  $C_{\rm z}$  FOR POINT SOURCE IN FOUR RANGES OF TEMPERATURE GRADIENT

Group	Average Temperature Gradient	$\overline{m}\lambda$	m <sub>z</sub>	$\overline{c}^{\lambda}$	Cz
1	13.4° F./1,000'	.408	0	9.578	73.750
2	7.4° F./1,000'	.466	.096	5.708	45.945
3	3.0° F./1,000'	.505	.182	3.459	22.110
4	-1.3° F./1,000'	.606	.279	1.490	15.090

Table 7

CALCULATED POINT- AND LINE-SOURCE DIFFUSION PARAMETERS

	Temperature	dient Point Sou				Added D			Line Source				
Day	Gradient F./1,000'	my	m <sub>z</sub>	Cy	$C_{\mathbf{Z}}$	(Ft Width	Depth	my	m <sub>Z</sub>	Cy	$C_{\mathbf{Z}}$		
1	6.5	.466	.096	5.9	41.0	199	5	.419	.095	9.4	41.8		
2	3.7	.505	.182	3.9	25.5	10	50	.500	.169	4.0	29.3		
· 3	8.2	.466	.096	5.9	41.0	52	48	.452	.090	6.6	47.2		
4	2.6	.505	.182	<b>3.</b> 9	25.5	126	<b>39</b> ,	.475	.171	5.3	28.6		
. 5	2.7	.505	.182	3.9	25.5	300	ı	.434	.182	7.7	25.5		
. 6	-0.2	.606	.279	1.7	15.0	276	29	.527	.268	3.2	16.2		
7	7.6	.466	.096	5.9	41.0	288	21	.410	.093	11.3	43.5		
8	2.3	.505	.182	3.9	25.5	288	21	.439	.177	7.5	27.0		
9	3.8	.505	.182	3.9	25.5	187	18	.460	.178	6.0	26.8		
10	-2.3	.606	.279	1.7	15.0	0	0	.606	.279	1.7	15.0		
11	. <b>-</b>	.408	0	9.5	75.0	95	8	.371	0	14.7	75.0		
12	13.4	.408	0	9.5	75.0	54	21	.396	0	10.7	75.0		

Table 8 MEASURED AND CALCULATED AXIAL SO2 CONCENTRATIONS IN PLUME (PPM)

			Dist	tance (Miles	s)	
Day		1/2	3/4	Ţ	2	x
1	(A) (B)	18.1 21.1	16.0 17.0	16.3 14.7	10.2	7.9 6.1
2	(A) (B)	29.8 30.0	24.2 22.9	20.1 19.0	12.9 11.9	5.6 4.8
3	(A) (B)	25.4 20.9	21.1 16.7	13.9 14.4	9.0 9.9	
14	(A) (B)	17.2 15.3	16.9 11.7	15.2 9.7	9.6 6.2	4.4 2.6
5	(A) (B)	24.4 34.7	18.2 26.6	19.1 22.7	16.8 14.8	4.5 5.3
6	(A) (B) (B)*	20.1 37.6 34.1	15.2 27.3 23.8	14.5 21.6 18.5	16.8 12.4 10.0	13.2 3.9 12.5
7	(A) (B)	20.1 31.4	19.1 25.7	22.4 22.4	20.5 16.0	11.2 7.7
8	(A) (B)	25.1 39 <b>.</b> 7		6.6 26.0	4.6 16.9	
9	(A) (B)			8.0 7.2		3.3 2.8
10	(A) (B)			2.2 6.8	0.9 3.7	
11	(A) (B)	12.1 23.1				<b>7.</b> 5 9.5
12	(A) (B)	17.5 13.7				9.1 5.4

<sup>(</sup>A) Field-measured values.(B) Values calculated from point-source coefficients modified

to simulate line-source field conditions.

\* Temperature gradient at 1/2, 3/4, 1, and 2 miles = -2.4° F./1,000 feet.

Temperature gradient at x miles = 8.7° F./1,000 feet.

Table 9

SUMMARY - PRINCIPAL DATA, BY SECTIONS - HIGH WIND AND NEUTRAL CONDITIONS

(FIELD-MEASURED VALUES)

				Do our	Elev. (		73		90. 9		Average	Temp. Gradient Plume			D: 00			\$0 <sub>≥</sub>
Day	Dist.	Time	Units Operating	Degrees Off Line of Stacks	Maximum Conc.	Maximum Plume Width	w. th	Depth (Ft.)	SO <sub>2</sub> C (pp:		Wind Speed in Plume (mph)	Bottom to Top oF./1,000	σ <sub>y</sub> (Ft.)	°z (Ft.)		on Paramensionle		Emission Rate (cfs)
. 1	1/2	1229-1238 9 min.	1, 2, 4	4-1/2	800	800	1465	800	10.0	4.8	8.2	-6.4	279	212	0.969	0.384	0.586	44.86
	1	1242-1331 49 min.	1, 2, 4	4-1/2	800	800	2020	1200	4.0	2.2	8.0	-5.8	385	384				
	2	1340-1449 69 min.	1, 2, 4	4-1/2	1800	1200	2495	1800	2.0	0.9	10.7	-5.9	475	478				
2	1/2	1315-1327 12 min.	1, 2, 4	4	600	600	910	600	15.3	7.8	13.1	-5.8	173	127	1.387	0.704	0.682	40.57
	1	1329-1342 13 min.	1, 2, 4	14	800	800	1310	850	7.9	4.1	11.5	-5.6	250	193				
	2	1346-1417 31 min.	1, 2, 4	4	1000	1000	2410	1450	3.2	1.3	9.1	-5.9	459	327				
3	1/2	1145-1155 10 min.	1, 3, 4	85-1/2	800	800	900	650	12.9	5.8	14.2	-6.5	171	145	1.254	0.563	0.691	41.83
	1	1159-1219 20 min.	1, 3, 4	85-1/2	800	800	1575	1250	4.3	2.4	12.8	-6.3	300	275				
	2	1224-1246 22 min.	1, 3, 4	85-1/2	1000	1000	1957	1300	2.7	1.3	11.9	-6.7	373	378				
4	1/2	1223-1234 11 min.	1, 3, 4	86	1000	1000	826	800	13.9	7.1	21.6	-5.4	157	106	1.665	0.675	0.993	46.24
	1	1236-1249 13 min.	1, 3, 4	86	600	600	1200	1050	3.5 4.0*	2.1	20.4	-4.8	229	202				
	2	1255-1326 31 min.	1, 3, 4	86	800-1600	1000	2100	1650	1.3	0.9	14.9	-5.5	400	420				
5	1/2	0929-0942 13 min.	1, 3, 4	45	1000	600-1000	1186	800	8.9	4.6	13.9	-3.5	226	167	1.035	0.543	0.493	43.09
	ı	0946-1011 25 min.	1, 3, 4	45	1000	800-1000	1705	1000	3.5	1.9	14.5	-4.7	325	284				
	2	1017-1043 26 min.	1, 3, 4	45	800	800-1000	2520	1100	2.1	1.1	14.0	-3.9	480	331				
6	1/2	1326-1337 11 min.	1, 3, 4	2	600-1000	600-1000	957	900	8.4	4.6	9.9	-6.0	182	171	1.309	0.672	0.640	41.83
	1	1340-1403 23 min.	1, 3, 4	2	800-1600	800-1600	1517	1300	6.6	2.8	6.6	-5.8	289	360				
	2	1407-1427 20 min.	1, 3, 4	2	800-1400	800-1400	2427	1500	0.7	0.4	8.5	-6.0	462	415				
7	1/2	1021-1036 15 min.	1, 2, 3, 4	54	800	800	1042	1150	6.0 10.5*	3.2	11.5	-6.5	198	223	0.800	0.384	0.418	49.27
	1	1042-1115 27 min.	1, 2, 3, 4	54	600	600-1000	1476	1450	5.1	2.6	12.9	-6.1	281	289				
	2		1, 2, 3, 4	54	1200	600-1400	1770	1500	1.4 2.8*	8.0	17.3	-6.6	337	398				
8	2	1401-1500 1505-1638	1, 2, 3 1, 2, 3	11 11	1200 1200	400-1200 1200	880 1452	1200 1800	4.7 2.4	2.3	16.1 14.6	-4.6 -6.5	168 277	327 427	1.109	0.721	0.385	38.03
9	1	0929-1000	1, 2, 3	55	500	500	1520	800	3.0 5.2*	1.5	11.1	-5.5	290	271	0.800	0.516	0.291	41.83
	2 3	1016-1037 1039 <b>-</b> 1105	1, 2, 3 1, 2, 3	55 55	1000 800	600 800	2162 2680	1050 1300	1.6	0.8	13.3 15.6	-5.3 -5.4	412 510	352 380				
10	1 2	1115-1445 1129-1506	1, 2, 3 1, 2, 3	65 65	1000 1400	800 800	1012 1408	1100 1400	6.2 2.7	2.7 1.1	20.6 23.1	-5.9 -6.1	193 268	219 <b>323</b>	1.034	0.474	0.561	49.77
11	1/2 1 2-1/2	1238-1300 1302-1344 1418-1519	1, 2 1, 2 1, 2	n n	800 800 1200	700 1000 1200	1408 2640 3430	1000 1500 2500	4.9 2.2 0.7	1.7 1.3 0.4	12.2 8.5 10.4	-5.2 -5.0 -5.2	268 503 653	195 332 656	1.338	0.582	0.755	28.85

<sup>\*</sup>Concentration measured along horizontal flight parallel to x axis.

Table 10

SUMMARY - PRINCIPAL DATA, BY SECTIONS

(FOR SINGLE STACK POINT SOURCE)

		Plu		SO <sub>2</sub> Co				Diffusion Parameters  m m <sub>V</sub> m <sub>Z</sub>						SO <sub>2</sub>
Day	Dist. (Mi.)	Width (Ft.)	Depth (Ft.)	(ppr Max.	Av.	σy (Ft.)	$\frac{\sigma_{\mathbf{z}}}{(\text{Ft.})}$	All	*	All	<u>*</u>	<u>All</u>	<u>*</u>	Emission (cfs)
1	1/2 1 2	1442 1997 24 <b>7</b> 2	725 1125 1725	3.7 1.4 0.7	1.8 0.8 0.3	2 <b>7</b> 5 381 471	198 370 464	0.996	1.411	0.388	0.500	0.614	0.758	14.95
2	1/2 1 2	889 1289 2389	525 775 1375	6.0 2.9 1.1	3.0 1.5 0.5	169 246 455	113 179 313	1.444	1.444	0.714	0.714	0.735	0.735	13.52
3	1/2 1 2	601 1276 1658	644 1244 1294	6.5 1.8 1.1	2.9 1.0 0.5	114 243 316	144 274 377	1.476	1.476	0.735	0.735	0.694	0.811	13.95
4	1/2 1 2	527 901 1801	795 1045 1645	7.3 1.6 0.5	3.7 1.0 0.3	106 172 343	105 201 419	1.896	1.806	0.847	0.847	0.998	0.968	15.41
5	1/2	974 1493 2308	747 947 1047	3.9 1.4 0.8	2.0 0.8 0.4	186 285 440	157 274 321	1.123	1.348	0.621	0.621	0.516	0.660	14.36
6	1/2 1 2	946 1506 2416	825 1225 1425	3.1 2.4 0.2	1.7 1.0 0.1	180 287 460	157 346 401	1.354	1.383	0.677	0.677	0.676	0.676	13.94
7	1/2 1 2	799 1233 1527	1106 1406 1456	2.0 1.6 0.4	1.1 0.8 0.2	152 235 291	215 281 390	0.903	1.007	0.468	0.629	0.430	0.500	12.32
8	1 2	842 1414	1151 1751	1.7 0.8	0.8	161 270	318 418	1.140	1.140	0.746	0.746	0.394	0.500	12.68
9	1 2 3	1356 1998 2516	771 1221 1571	1.2 0.6 0.5	0.6 0.3 0.3	259 381 4 <b>7</b> 9	265 346 374	1.186	1.425	0.560	0.560	0.297	0.500	13.94
10	1 2	831 1227	1079 1379	2.6 1.0	0.4	159 2 <b>3</b> 4	215 319	1.147	1.147	0.557	0.557	0.569	0.569	16.59
11	1/2 1 2-1/2	1389 2621 3411	979 1479 2479	2.5 1.1 0.4	0.9 0.7 0.2	264 499 649	191 328 652	1.365	1.365	0.588	0.739	0.764	0.764	14.43
							Average	1.294	1.418	0.624	0.686	0.625	0.759	

 $m_y = 1.000$  and <2.000  $m_y = 1.000$  and <1.000

Table 11

VALUES OF  $C_y$  AND  $C_z$  (FT.  $\frac{m}{z}$ ) CALCULATED FOR EACH SECTION (POINT SOURCE)

FOR VALUES OF  $m_y = m_z = 0.80, 0.75$ , AND 0.70 OR m = 1.6, 1.5, AND 1.4

Day	Dist. (Mi.)	$\frac{m_{\mathbf{y}} = m_{\mathbf{z}} = m_{\mathbf{z}}}{\underline{C}\mathbf{y}}$	0.80 Cz	$\frac{m_{\mathbf{y}} = m_{\mathbf{z}}}{\underline{c_{\mathbf{y}}}}$	$= 0.75$ $\underline{C_z}$	$\frac{m_{y} = m_{z}}{c_{y}}$	= 0.70 <u>Cz</u>	Wind Speed (fps)
1	1/2 1 2 Average	.401 .285	.363 .398 .280 .347)	.747 .616 .452 (.605)	.538 .598 .445 (.523)	1.107 .944 .719 (.923)	.797 .917 .708 (.807)	12.1 11.8 15.7 (13.2)
2	1/2 1 2 Average	.259 .275	.207 .188 .186 .194)	.459 .397 .437 (.431)	.307 .289 .300 (.299)	.680 .610 .694 (.661)	.455 .444 .478 (.459)	19.2 16.9 13.4 (16.5)
3	1/2 1 2 Average	.256 .191	.264 .288 .228 .260)	.310 .393 .303 (.335)	.391 .443 .361 (.398)	.459 .602 .482 (.514)	.580 .679 .575 (.611)	20.8 18.9 17.5 (19.1)
Ц	1/2 1 2 Average	.181 .207	.192 .211 .253 .219)	.288 .278 .329 (.298)	.285 .325 .402 (.337)	.427 .426 .523 (.459)	.423 .498 .639 (.520)	31.8 30.0 21.9 (27.9)
5	1/2 1 2 Average	.300 .266	.287 .288 .194 .256)	.505 .460 .422 (.462)	.427 .443 .308 (.393)	.749 .706 .671 (.709)	.632 .679 .490 (.600)	20.4 21.3 20.6 (20.7)
6	1/2 1 2 Average	.302 .278	.287 .364 .242 * .298)	.489 .464 .441 (.465)	.427 .559 .385 (.457)	.725 .711 .702 (.713)	.632 .857 .612 (.700)	14.5 9.7 12.4 (12.2)
7	1/2 1 2 Average	.274 .176	.394 .296 .236 .308)	.413 .380 .279 (.357)	.584 .454 .374 (.471)	.612 .582 .444 (.546)	.866 .696 .595 (.719)	16.9 19.0 25.4 (20.4)
8	1 2 Average		.334 .252 .293)	.260 .259 (.260)	.514 .401 (.458)	.399 .412 (.405)	.788 .638 (.713)	23.6 21.5 (22.5)
9	1 2 3 Average	.230 .209	.279 .209 .163 .217)	.418 .366 .339 (.374)	.428 .332 .265 (.342)	.642 .581 .550 (.591)	.657 .528 .430 (.538)	16.3 19.6 22.9 (19.6)
10	1 2 Average	.141	.226 .193 .209)	.257 .225 (.241)	.347 .306 (.327)	.394 .357 (.376)	.533 .487 (.510)	30.3 34.0 (32.2)
11	1/2 1 2-1/2 Average	.525 .328	.350 .345 .329 .341)	.717 .806 .527 (.683)	.519 .530 .529 (.526)	1.063 1.237 .847 (1.049)	.769 .813 .852 (.811)	17.9 12.5 15.3 (15.2)
	Average All days	(.267) (.	267)	(.410)	(.412)	(.631)	(.635)	

Table 12  $\label{eq:wind_speed} \mbox{Wind SPEED, $C_{\mathbf{y}}$ AND $C_{\mathbf{z}}$, $AND $m_{\mathbf{y}}$ AND $m_{\mathbf{z}}$ VALUES}$ 

					Calculated	Values (	Cy and Cz, Ft.	m Z		
Wind	Speed	m	$v = m_z =$	0.80*	m	$y = m_z =$	0.75**	$m_{ m V}$	$- = m_{\mathbf{Z}} = 0$	).70 <del>***</del>
(mph)	(fps)	$c_{\mathbf{y}}$	$\frac{C_{\mathbf{Z}}}{C_{\mathbf{Z}}}$	$C_{y} = C_{z}$	CY	$c_{\mathbf{z}}$	$C_y = C_z$	Cy	$C_{\mathbf{Z}}$	$C_y = C_z$
8	11.76	<b>.33</b> 9	.343	.341	.522	.524	.523	.807	.803	.805
10	14.70	.305	.316	.310	.470	.484	.477	.727	.742	•735
12	17.64	. 275	.291	.283	.423	.447	.435	.654	.687	.670
14	20.58	.248	.268	.257	.381	.412	.396	.589	.635	.612
16	23.52	.223	.247	.234	.343	.381	.361	.530	.588	.558
18	26.46	.201	.227	.214	.309	.351	.330	.477	. 544	.509
20	29.40	.181	.209	.194	.279	.324	300	.430	.503	.465
22	32.30	.163	.193	.177 。	.251	.300	.274	.387	.466	.424
					Ве	est Estim	ate			
		٠.		·	<b>-</b>		<b></b>			•

u - Wind speed in fps

Table 13

DIFFUSION COEFFICIENTS my, mz, Cy, AND Cz - POINT AND LINE SOURCES

		77.		Point-S	Source Value	<u>s</u>			L	ine-Sour	ce Field Va	
Day	Dist. (Mi.)	Wind Speed (fps)	m <sub>y</sub>	m <sub>Z</sub>	Cy m (Ft. 2)	C <sub>z</sub> m (Ft.)	$^{\triangle}$ y $_{n}$ (Ft.)	$rac{ riangle_{\mathbf{z_n}}}{ ext{(Ft.)}}$	my	$m_{\mathbf{Z}}$	Cym (Ft. <sup>2</sup> )	Cz (Ft.
1	1/2 1 2	12.0 11.7 15.7	.75	.75	.510 .515 .450	.510 .515 .465	23	75	.74	.71	.561 .567 .495	.740 .745 .674
2	1/2 1 2	19.2 16.9 13.4	.75	•75	.400 .430 .490	.425 .450 .490	21	75	·7 <sup>1</sup> 4	.71	.440 .473 .539	.616 .653 .711
3	1/2 1 2	20.8 18.7 17.5	<b>.7</b> 5	<b>.7</b> 5	.378 .405 .420	.408 .430 .440	299	6	.60	.74	1.550 1.661 1.722	.409 .430 .440
<b>4</b>	1/2 1 2	31.7 29.9 21.9	.75	.75	.260 .270 .360	.305 .320 .395	299	5	.60	.74	1.066 1.107 1.476	.305 .320 .395
5	1/2 1, 2	20.3 21.2 20.1	.75	<b>.7</b> 5	.380 .370 .385	.410 .400 .420	212	53	.64	.72	1.102 1.073 1.117	•533 •520 •546
6	1/2 1 2	14.5 9.7 12.4	•75	<b>.7</b> 5	.470 .550 .505	.480 .540 .505	11	75	-74	.71	.494 .578 .530	.696 .783 .732
7	1/2 1 2	16.9 18.9 25.4	•75	.75	.430 .400 .320	.450 .425 .360	243	<del>ի</del> կ	.62	.73	1.419 1.320 1.056	.563 .531 .450
8	1 2	23.6 21.4	.75	<b>.7</b> 5	.340 .370	.378 .400	38	49	.73	.72	.408 .444	.476 .504
9	1 2 3	16.3 19.5 22.9	.75	.75	.440 .390 .350	.460 .420 .385	164	29	.66	.74	1.021 .905 .812	.520 .470 .319
10	1 2	29.6 3 <b>3.</b> 9	.75	•75	.275 .240	.320 .285	181	21	.66	.74	.688 .600	.352 .314
11	1/2 1 2 <b>-</b> 1/2	17.9 12.5 15.2	.75	.75	.420 .505 .460	.440 .505 .470	19	21	.74	.74	.462 .556 .506	.484 .556 .517

Table 14 MEASURED AND CALCULATED

## AXIAL SO2 CONCENTRATIONS IN PLUME

				ntrations		
Day		1/2 Mi.	1 Mi.	2 Mi.	2-1/2 Mi.	3 Mi.
1	(A) (B)	10.0 13.4	4.0 5.8	2.0		
2	(A)	15.3 11.6	7.9 4.9	3.2 1.8		
3	(A) (B)	12.9 11.4	4.3 5.1	2.7 2.0		
<b>4</b>	(A) (B)	13.9 16.1	5.3 7.1	2.0 2.3		
5	(A) (B)	8.9 11.0	3.5 5.0	2.1 1.9		
6	(A) (B)	8.4 12.5	6.6 6.2	0.7 2.0		
7	(A) (B)	10.0 12.1	5.1 5.6	2.3 2.4		
8	(A) (B)		4.7 5.3	2.4 1.9		
9	(A) (B)		5.2 4.7	1.6 1.9		1.5 1.5
<b>1</b> 0	(A) (B)		6.2 6.8	2.7 2.9		
11	(A) (B)	4.9 8.4	2.2 3.7		0.7 0.9	

<sup>(</sup>A) = Field-measured values.(B) = Values calculated from point-source coefficients modified to line-source field conditions.

Taole 15

CHAMBER SO<sub>2</sub> OXIDATION STUDIES

Date					Percer	0 مnt SO	xidatio								
<u>(1959)</u>	Type of Run	Uniform	n SO <sub>2</sub> S	upply				No SC	Supp.	Ly - Nat	ural De	ecay	·		
		<u>2a</u>	1	2	2	4	2	<u>6</u>	7	<u>8</u>	2	<u>10</u>	11	12	<u>13</u>
10/13	Fly ash and moisture	16.5	8.6	9.7 /	24.0	25.8	27.0 /	3.7	-1.3						
10/16	Fly ash and dry		-2.1	<b>-2.8</b> /	3.9	15.6	23.5	33.3	22.8	28.4 /	-2.1				
10/19	Fly ash and moisture		<b>-</b> 6.9	-1.8 /	12.2	11.5	9.5 /	19.2	14.0	11.9	14.3	15.3	8.5	-2.3	
10/20	Moisture (Fly ash removal)		-6.5	-6.9 /	11.3	13.2	17.3	12.3 /	11.8	7.6	9.9	2.4	4.6	0	-4.0
10/22	Fly ash and moisture		-1.2	4.3 /	13.5	15.6 /	6.9	7.7	9.7	8.8	8.4	10.4	10.5		
					<u>Ca.</u>	libratio	n Runs								
10/9	SO <sub>2</sub> cylinder (Not through chambe		.0 2.7	6.3 /											
10/23	SO <sub>2</sub> cylinder (Through chamber)			/	0.8	-1.7	-4.9	-4.8	3.9	0					

Negative (-) Values (Titrilog concentration more than autometer concentration)
/ Instruments switched from controlled diluted sample to straight sample

Table 16

CHAMBER SO<sub>2</sub> OXIDATION STUDIES

### COLBERT STEAM PLANT

Date (1959)	Time Sampling Period	Dust Count	ts (mppcf) Millipore Filter	Geometric Mean Particle Size (Microns)	Relative Humidity(%)	Dry Bulb Temp. (°F.)	SO <sub>2</sub> Titrilog (ppm by Vol.)	Total Sulfates as SO <sub>3</sub> (ppm by Vol.)	SO <sub>2</sub> Oxidation (%)
10/29 10/29	1335-1405 1406-1431	1.83 .98	-	-	96 96	65 65	2.0 2.0	.640 1.004	24.2 33.4
11/2 11/2 11/2	1310-1330 1337-1353 1438-1458 1528-1548	2.94 - - -	1.28 1.35 6.7 6.3	.56 1.50 .63 1.29	95 98 99 95	71 72 72 72	2.7 2.6 2.4 2.3	1.03 1.22 .75 1.13	27.6 31.9 23.8 32.9
	Average	2.94	3.90	•99	97	72	2.5	1.06	29.8
11/3 11/3 11/3 11/3	0947-1007 1014-1034 1051-1111 1131-1151	6.37 6.28 3.18 5.4	4.9 4.9 7.2 6.7	.74 .74 .40 .40	98 96 95 97	64 66 69 71	5.9 5.5 5.3 5.3	.88 1.00 .81 .69	13.0 15.4 13.3 11.5
	Average	5.31	5.93	.57	97	68	5.5	.84	13.2
11/3 11/3 11/3 11/3	1351-1400 1416-1436 1500-1520 1546-1606	1.13 1.30 1.47 1.30	3.6 2.7 2.5 2.6	.54 .54 .91 .86	57 58 56 56	79 80 81 81	5.6 5.5 5.5 5.2	2.81 1.07 .84	33.4 16.3 13.9
<del></del> / )	Average	1.30	2.85	.71	57	80	5.5	.96	14.9
11/4 11/4 11/4 11/4	0859-0920 0926-0947 1022-1042 1103-1123	1.87 1.87 1.12 .75	6.3 6.3 5.2 3.5	.48 .48 .32 .32	100 100 99 98	73 73 73 73	9.4 9.6 9.1 9.2	.59 .59 1.05 1.12	5.9 5.8 10.3 10.9
	Average	1.40	5.33	.40	99	73	9.3	.85	8.4

<sup>\*</sup>Average ratio impinger counts to millipore filter counts = 1:2.97.

Table 16 (Continued)

Date (1959)	Time Sampling Period	Dust Coun	ts (mppcf) Millipore Filter	Geometric Mean Particle Size (Microns)	Relative Humidity (%)	Dry Bulb Temp. (°F.)	SO <sub>2</sub> Titrilog (ppm by Vol.)	Total Sulfates as SO <sub>3</sub> (ppm by Vol.)	SO≥ Oxidation (%)
11/4 11/4 11/4 11/4	1340-1410 1412-1440 1442-1504 1505-1530	8.72 3.10 1.42 1.58	- - -	- - - -	96 96 96 96	71 71 71 71	64.7 39.7 19.9 11.8	.94 .31 .29 .60	1.4 0.8 1.4 4.8
	Average			·	96	71		•54	1.6
11/5 11/5	0958-1017 1027-1047	.83 1.15	5.3 5.3	•32 •32	53 49	85 84	5.3 5.6	1.08 1.28	16.9 18.6
	Average	•99	5.3	.32	51	85	5.5	1.19	17.8
11/5 11/5 11/5 11/5	1300-1320 1327-1345 1420-1440 1457-1517	.64 .54 1.07 .94	.91 1.58 1.89 1.37	1.06 .86 .49 .49	45 43 42 40	81 81 81 81	10.4 10.6 10.5 10.4	.45 .20 .36 .53	4.1 1.9 3.3 4.8
	Average	•94	1.44	.73	43	81	10.4	•39	3.6
11/6 11/6 11/6 11/6	0904-0924 0924-0944 1009-1120	.13 .13 .22 .22	.54 .54 1.22 1.22	1.07 1.07 1.06 1.06	90 96 97 97	46 46 46 47	10.3 10.6 10.1 9.2	.03 .03 .05 .05	0.3 0.3 0.5 0.5
	Average	.18	.88	1.07	95	46	10.1	.04	0.4
11/6	1156-1216	1.46	4.8	.80	86	47	22.9	1.15	4.8

<sup>\*</sup>Average ratio impinger counts to millipore filter counts = 1:2.97.

Table 17 CHAMBER SO2 OXIDATION STUDIES

## COLBERT STEAM PLANT

			D)4-	4 . 4	: - D		3		ibber Sa								9	ပ <sub>ဋ</sub> Concent		
Date	Test	Wt.			Resist.	Est.	Minutes		Bb SO <sub>2</sub>	Filter B	Wt.			ly Ash Sa Resist.	Est.	% SO <sub>4</sub>			Autometer Minus Titrilog	*
(1959)	No.	(mg.)	Dil.		(Ohms)	SO <sub>4</sub> (%)	Operated	(ppm)	(ppm)	(mg.)	(mg.) <sup>c</sup>	Dil.		(Ohms)	SO4 (%)	Grav.	(ppm)	(ppm.)	(ppm)	<u>Oxidation</u>
9/30	1 2 3 4 5 6	6.0	10:1 10:1 10:1 10:1	6.4 4.9	16,500 14,300 32,600	6.2 6.4 8.3	30 30 30 30 30	58.3 50.1 29.6 16.9 8.9 4.5	62.9 56.3 32.2	0.0	1287.1	10:1	6.5	52,000	5.6					
10/1	1 2 3 4 5 6	2.5 1.3 0.4 0.5	10:1	4.8			30 30 30 30	20.7 13.8 7.9 6.3 5.2 3.9	20.5 13.2 8.9 6.2 4.9 3.6	0.0 0.0 0.0 0.0 0.0	2584.3	10:1	7.0	77,000	3.9					
10/2	123156	18.3 8.5 4.2	10:1 10:1 10:1 10:1	5.6 6.0 4.9	67,000 44,500 53,000 27,200	4.4 6.4 5.4 9.6	30 30 30 30 60	50.0 44.1 29.1 21.5 15.8 12.5	50.1 45.0 30.4 22.5 17.1 12.3	0.1 0.2 0.0 0.0 0.0	1644.3	10:1	7.2	67,000	4.4					
10/12	1 2 3	2.2	10:1 10:1 10:1	5.0	22,600	11.2	30 30 30	24.9 3.2 0.8	24.2 3.3 0.5	0.6 0.4 0.2	2350.0 1117.4	10:1	7.1	74,000	4.0					
10/13	1 2 3 4 5 6	17.7 12.2 10.2 2.4	10:1 10:1 10:1	4.6	49,000	5.8	30 30 30 30 30 30	41.4 39.7 41.1 20.1 10.5 5.1	40.2 40.6 37.4 20.7 10.0 6.3	1.4 1.0 0.8 1.2 0.7	2750.2 2336.1	10:1	6.7	53,000	5.4		45.19 45.29 43.76 31.52 15.40 9.08	41.31 40.90 36.52 23.96 11.42 6.63	3.88 4.39 7.24 7.56 3.98 2.45	8.59 9.69 16.54 23.98 25.84 26.98
10/ <b>1</b> 6	1 2 3 4 5 6	26.7 26.7 14.0 6.6 3.0	10:1	6.6 <sup>d</sup> 6.6 5.5	42,500 37,000	5.6 <sup>e</sup> 6.6 7.6	30 30 29 29 30	42.3 45.4 38.8 27.9 16.1 11.6	41.0 46.3 38.4 22.8 17.2 10.8	0.8 0.4 0.2 0.0 0.0	924.9 50 <b>7.3</b>	10:1	8.0 6.6d	41,500	6.8		48.81 52.24 49.61 33.03 21.37 14.74	49.83 53.72 47.66 27.89 16.34 9.83	-1.02 -1.48 1.95 5.14 5.03 4.91	-2.09 -2.83 3.93 15.56 23.54 33.31
10/19	1	43.2	10:1	6.1 4.6			33	36.2	31.3	0.6	2219.0	10:1	7.2	68,000	4.4		36.90	39.46	-2.56	-6.94
	2	37.7	10:1	6.3			29	38.5	35.9	0.2	2031.1		3.7d			2.6	38.75	39.46	-0.71	-1.83
	3	11.7	10:1	6.0	•		29	19.9	16.9	0.0							25.22	22.14	3.08	12.21
	4 5 6		10:1				28 30 30	9.9 6.7 5.2	10.4 6.4 4.9	0.0							12.51 7.59 4.59	11.07 6.87 3.71	1.44 0.72 0.88	11.51 9.49 19.17
10/20	1 2 3 4 5 6	35.4 11.5 2.9		6.7 <sup>d</sup>		10.4 <sup>e</sup>	57 58 58	42.0 42.8 24.4 12.1 6.9 4.7	39.6 38.8 21.6 11.2 7.1 4.9	0.5 0.7 0.0 0.7 0.4 0.7	2309.3 1530.6		3.5 <sup>d</sup>			2.3	39.87 38.81 28.22 13.87 7.80 4.72	42.47 41.51 25.04 12.04 6.45 4.14	-2.60 -2.70 3.18 1.83 1.35 0.58	-6.52 -6.96 11.27 13.19 17.31 12.29
10/22	1 2 3 4 5 6	7.7 6.6					30 150	32.8 25.4 10.9 4.7 2.7 2.0			2271.0		3.7 <sup>d</sup>				32.70 25.60 12.60 4.50 2.45 1.95	33.10 24.50 10.90 3.80 2.28 1.80	-0.40 1.10 1.70 0.70 0.17 0.15	-1.22 4.30 13.49 15.56 6.94 7.69

<sup>a. Without filter
b. With filter
c. Sample period, 40-70 minutes
d. Soxhlet extraction entire sample, diluted to 250 ml. and refluxed for 2 hours
e. Gravimetric analysis</sup> 

Table 18

EFFECT OF DILUTION ON FLY ASH pH

Sample	el: Electrostatic precipitator No. 1 collected			Dat	te		
•	from Colbert trailer on 10/20/59, weight	10/2	1/59	10/2	1/59	10/2	2/59
	35.4 mg., and initially diluted to 45 cc.	Time	рН	Time	рН	Time	рН
Inst	rument check against standard solution pH - 7.00		7.00		7.00		7.00
la.	Aliquot of initial dilution, unfiltered	0900	4.50	1334	4.55	0842	4.50
lb.	Aliquot of initial dilution, filtered	0903	4.49	1332	4.50	0843	4.50
lc.	Aliquot, dilution increased to 10 cc. per mg., filtered	0910 0909	6.20 6.22	1346 1335	6.00 6.00	0845	6.16
*1d.	Aliquot, dilution increased to 10 cc. per mg., unfiltered	0913 0912	5.82 5.80	1337	5 <b>.7</b> 0	0900 0843	5.84 5.90
Sample	2: Thimble fly ash sample collected inside unit 2 duct on 10/20/59 - 100 mg. diluted to 100 cc. of distilled water	0050	<del>.</del> 0				
2a.	Aliquot of initial dilution, unfiltered	0950 0947	7.18 7.20	1339	7,30	0849	7,22
2b.	Aliquot of initial dilution, filtered	0958	6.85	1341	6.92	0852	6.88
2c.	Aliquot dilution increased to 10 cc. per mg., filtered	1000	7.00	1343	7.00	0854	6.99
2d.	Aliquot, dilution increased to 10 cc. per mg., unfiltered	0954	6.90	1345	6.99	0856	6.90
Inst	. check against standard solution		7.00		7.01		7.00
pH d	istilled water, unfiltered	0924	6.90				
pH d	istilled water, filtered	0925	6.70				

<sup>\*</sup>This was bottom sample, more visible fly ash

Table 17 CHAMBER SO2 OXIDATION STUDIES

# COLBERT STEAM PLANT

			m						bber Sa			_	. 5				s	U <sub>2</sub> Concent		
Date	Test	Wt.	Electi	rostat	Resist.	itator Sa Est.	Minutes	A <sup>a</sup> S	о <sup>5</sup> Вр	Filter B	Wt.	Duc	t - F.	ly Ash Sa Resist.	Est.	% SOL	Autometer	Titrilog	Autometer Minus Titrilog	4
(1959)	No.	(mg.)	Dil.	рН	(Ohms)	SO <sub>14</sub> (%)	Operated	(ppm)	(ppm)	(mg.)	(mg.) <sup>c</sup>	Dil.	Нq	(Ohms)	SO4 (%)	Grav.	(ppm)	(ppm)	(mgq)	Oxidation
9/30	193456	30.6 6.0	10:1 10:1 10:1 10:1	6.4 4.9	16,500 14,300 32,600	6.2 6.4 8.3	30 30 30 30 30	58.3 50.1 29.6 16.9 8.9	62.9 56.3 32.2	0.0	1287.1	10:1	6.5	52,000	5.6					
10/1	1 2 3 4 5 6	2.5 1.3 0.4 0.5	10:1	4.8			30 30 30 30 30	20.7 13.8 7.9 6.3 5.2 3.9	20.5 13.2 8.9 6.2 4.9 3.6	0.0 0.0 0.0 0.0 0.0	2584.3	10:1	7.0	77,000	3.9					
10/2	1234 56	18.3 8.5 4.2	10:1 10:1 10:1 10:1	5.6 6.0 4.9	67,000 44,500 53,000 27,200	4.4 6.4 5.4 9.6	30 30 30 30 60	50.0 44.1 29.1 21.5 15.8 12.5	50.1 45.0 30.4 22.5 17.1 12.3	0.1 0.2 0.0 0.0 0.0	1644.3	10:1	7.2	67,,000	4.4					
10/12	5 3	2.2	10:1 10:1 10:1	5.0	22,600	11.2	30 30 30	24.9 3.2 0.8	24.2 3.3 0.5	0.6 0.4 0.2	2350.0 1117.4	10:1	7.1	7 <sup>1</sup> 4,000	4.0					
10/13	1 2 3 4 5 6	10.2	10:1 10:1 10:1	4.6	49,000	5.8	30 30 30 30 30 30	41.4 39.7 41.1 20.1 10.5 5.1	40.2 40.6 37.4 20.7 10.0 6.3	1.4 1.0 0.8 1.2 0.7 1.1	2750.2 2336.1	10:1	6.7	53,000	5.4		45.19 45.29 43.76 31.52 15.40 9.08	41.31 40.90 36.52 23.96 11.42 6.63	3.88 4.39 7.24 7.56 3.98 2.45	8.59 9.69 16.54 23.98 25.84 26.98
10/16	1 2 3 4 5 6	26.7 14.0 6.6	10:1 10:1 10:1 10:1	6.6 <sup>d</sup> 6.6 5.5	42,500 37,000	5.6 <sup>e</sup> 6.6 7.6	30 30 29 29 30	42.3 45.4 38.8 27.9 16.1 11.6	41.0 46.3 38.4 22.8 17.2 10.8	0.8 0.4 0.2 0.0 0.0	924.9 507.3	10:1	8.0 6.6d	41,500	6.8		48.81 52.24 49.61 33.03 21.37 14.74	49.83 53.72 47.66 27.89 16.34 9.83	-1.02 -1.48 1.95 5.14 5.03 4.91	-2.09 -2.83 3.93 15.56 23.54 33.31
10/19	1	43.2	10:1	6.1 4.6			33	36.2	31.3	0.6	2219.0	10:1	7.2	68,000	4.4		36.90	39.46	-2.56	-6.94
	2	37.7	10:1				29	38.5	35.9	0.2	2031.1		3.7 <sup>d</sup>			2.6	38.75	39.46	-0.71	-1.83
	3	11.7	10:1				29	19.9	16.9	0.0							25.22	22.14	3.08	12.21
	4 5 6		10:1	5.4			28 30 30	9.9 6.7 5.2	10.4 6.4 4.9	0.0							12.51 7.59 4.59	11.07 6.87 3.71	0.72 0.88	11.51 9.49 19.17
10/20	1 2 3 4 5 6	35.4 11.5 2.9	:	6.7 <sup>d</sup>		10.4e	57 58 58	42.0 42.8 24.4 12.1 6.9 4.7	39.6 38.8 21.6 11.2 7.1 4.9	0.5 0.7 0.0 0.7 0.4 0.7	2309.3 1530.6		3.5 <sup>d</sup>			2.3	39.87 38.81 28.22 13.87 7.80 4.72	42.47 41.51 25.04 12.04 6.45 4.14	-2.60 -2.70 3.18 1.83 1.35 0.58	-6.52 -6.96 11.27 13.19 17.31 12.29
10/22	1 2 3 4 5	7.7 6.6					30 150	32.8 25.4 10.9 4.7 2.7 2.0			2271.0		3.7ª				32.70 25.60 12.60 4.50 2.45 1.95	33.10 24.50 10.90 3.80 2.28 1.80	-0.40 1.10 1.70 0.70 0.17 0.15	-1.22 4.30 13.49 15.56 6.94 7.69

a. Without filter
b. With filter
c. Sample period, 40-70 minutes
d. Soxhlet extraction entire sample, diluted to 250 ml. and refluxed for 2 hours
e. Gravimetric analysis

Table 18

EFFECT OF DILUTION ON FLY ASH pH

Sample	1: Electrostatic precipitator No. 1 collected			Dat			
•	from Colbert trailer on 10/20/59, weight	10/2		10/2		10/2	
	35.4 mg., and initially diluted to 45 cc.	Time	рН	Time	<u>pH</u>	Time	pН
Instr	ument check against standard solution pH - 7.00		7.00		7.00		7.00
la.	Aliquot of initial dilution, unfiltered	0900	4.50	1334	4.55	0842	4.50
lb.	Aliquot of initial dilution, filtered	0903	4.49	1332	4.50	0843	4.50
lc,	Aliquot, dilution increased to 10 cc. per mg., filtered	0910 0909	6.20 6.22	1346 1335	6.00 6.00	0845	6.16
*ld.	Aliquot, dilution increased to 10 cc. per mg., unfiltered	0913 0912	5,82 5.80	1337	5 <b>.7</b> 0	0900 0843	5.84 5.90
Sample	2: Thimble fly ash sample collected inside unit 2 duct on 10/20/59 - 100 mg. diluted to 100 cc. of distilled water						
2a.	Aliquot of initial dilution, unfiltered	0950 094 <b>7</b>	7.18 7.20	1339	7.30	0849	7,22
2b.	Aliquot of initial dilution, filtered	0958	6.85	1341	6.92	0852	6,88
2c.	Aliquot dilution increased to 10 cc. per mg., filtered	1000	7.00	1343	7.00	0854	6.99
2d.	Aliquot, dilution increased to 10 cc. per mg., unfiltered	0954	6.90	1345	6.99	0856	6.90
Inst.	check against standard solution		7.00		7.01		7.00
pH di	stilled water, unfiltered	0924	.6.90				
pH di	stilled water, filtered	0925	6.70				

<sup>\*</sup>This was bottom sample, more visible fly ash

Table 19
SO2 OXIDATION STUDIES - COLBERT STEAM PLANT PLUME

Date (1960)	Sample No.	Travel from Po	pint of Emission Distance (Mi.)	Relative Humidity in Plume (%)	SO <sub>2</sub> Oxidation (%)
					Low Rates:
8/2	1 2 3 4	5 5 5 15	.25-1 .25-1 .25-1 1-1.5		0 0 1.20 0
·9/2	1 2	30 78	2 <b>-</b> 3 8		3.70 2.20
10/14	1 2	12 60	.5-1.5 5-6	62 54	2.15 3.23
10/26	<u>1</u> 2	6 84	.25-1.25 8-9	45 48	1.50 2.70
10/28	1 2	12 84	.5-1.5 8 <b>-</b> 9	68 70	1.10
					High Rates:
5/3	1 2 3	13 13 13	1.1 1.1 1.1		13.80 10.00 19.20
8/19	1 2	108 23	8-10 .75-2		55.50 8.00
10/11	1 2	12 96	.5 <b>-</b> 1.5 8	74 73	21.60 32.00

Table 20 SO2 OXIDATION STUDIES - COLBERT STEAM PLANT FLUME

Date (1960)	Sample No.	Distance from Plant (Miles)	Plume Travel Time (Min.)	T Start	ime Stop	*Sample Elev. (Ft.)	Approx. Wind Speed (mph)	Plume Temp.	Ground Temp.	Rel. Hu Ground	Plume	SO3** (ppm)	SO <sub>2</sub> ** (ppm)	Total** (mqq)	Oxidation (る)	Weather Observations
5/3	1 2 3	1.1 1.1 1.1	13 13 13	0526 0627 0719	0553 .0658 0 <b>7</b> 50	812 <b>713</b> 800	5 5 5	56 56 56	43 45 35			2.4 2.0 3.8	15.0 18.0 16.0	17.4 20.0 19.2	13.6 10.0 19.2	Fair; slight fog at ground. Fair; slight haze to the E and N of plant. Fair.
8/2	1 2 3 4	.25-1 .25-1 .25-1 1-1.5	5 5 5 <b>1</b> 5	0507 0555 0641 0728	0538 0627 0711 0811	660 700 730 920	7 7 7 5	77 77 76 79	74 76 80 83	92 86 80 <b>73</b>		0.0 0.0 0.06 0.0	6.0 6.0 5.0 3.0	6.0 6.0 5.06 3.0	0.0 0.0 1.2 0.0	No clouds; very hazy and smoky; no sunshine. Very hazy and smoky; no sunshine due to haze and smoke. Hazy and smoky; bright red sun visible through smoke and haze.
8/19	1 2	8-10 .75-2	108 23	0514 0630	0607 0655	950 950	5 5	69 67	71 71	+95 +95		1.0	0.8 2.3	1.8 2.5	55.5 8.0	Mist and fog over general area; complete cloud cover. Discontinued because of rain.
9/2	1 2	2-3 8	30 78	0521 '0656	0602 0743	850 1600	5 5	74 74	65 74	96 96		0.07 0.06	1.9 2.4	1.97 2.46	3.7 2.2	Haze; no mist. Fair; sun shining.
10/11	1 2	.5-1.5 8	12 96	0633 0736	0 <b>70</b> 8 0824	800 800	5 5	65 65	57 60	99 98	74 73	1.0	3.6 0.17	4.6 0.25	21.6 32.0	.8 cloud cover; fog and haze in low areas.
10/14	1 2	.5-1.5 5-6	12 60	0630 0724	0652 0814	700 700	5 5	68 70	58 61	96 89	62 54	0.11	5.0 2.1	5.11 2.1 <b>7</b>	2.15 3.23	.6 cloud cover; fog and haze in low areas8 cloud cover; fog and haze in low areas.
10/26	<u>:</u> 2	.25 <b>-1.</b> 25 8-9	6 84	0628 0724	0704 0820	600 <b>7</b> 50	6 6	62 60	58 62	89 75	45 48	0.07 0.03	4.6 1.1	4.67 1.13	1.5 2.7	High overcast of clouds; no sunlight. High overcast of clouds; no sunlight.
10/28	1 2	.5-1.5 8-9	12 84	0647 0800	0721 0855	800 900	5 5	50 48	47 56	99 8 <b>3</b>	68 70	0.04 0.03	3.6 0.71	3.64 0.74	1.1	Clear and sun shining. Clear; slightly hazy near ground.

<sup>\*</sup>Elevation above ground level at point of emission.

\*\*Based on soluble sulfate; first decimal determinations were gravimetric; second decimal determinations were colorimetric.

# TENNESSEE VALLEY AUTHORITY Division of Health and Safety

and

PUBLIC HEALTH SERVICE Division of Air Pollution

### APPENDIX B

**FIGURES** 

Chattanooga, Tennessee August 1964

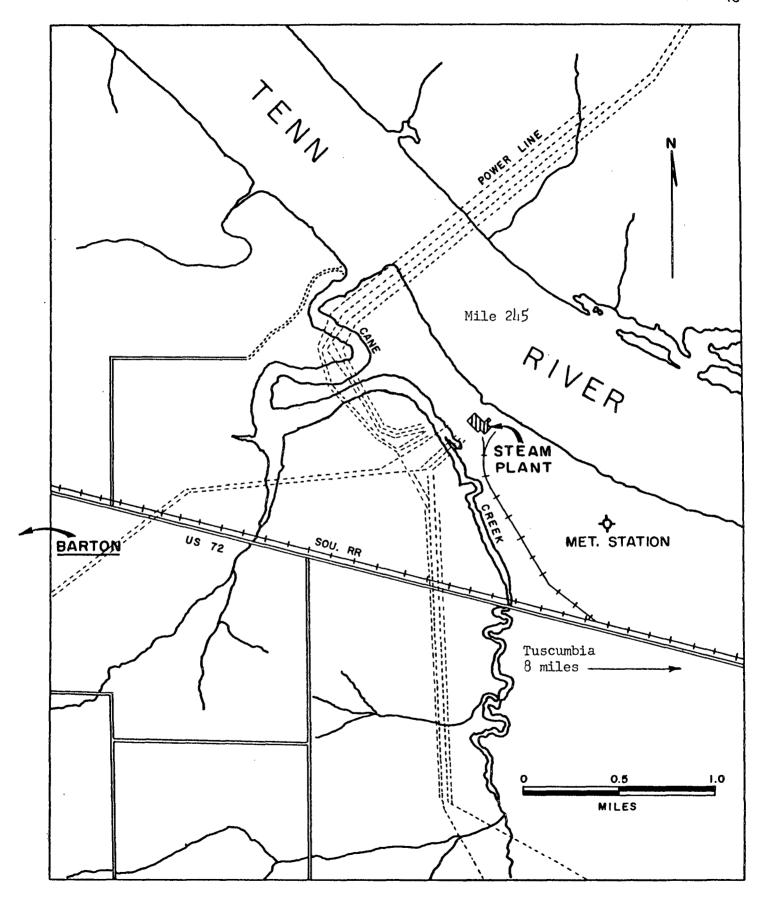


Figure 1. MAP OF COLSERT STEAM PLANT SITE - NEAR TUSCUMBIA, ALABAMA

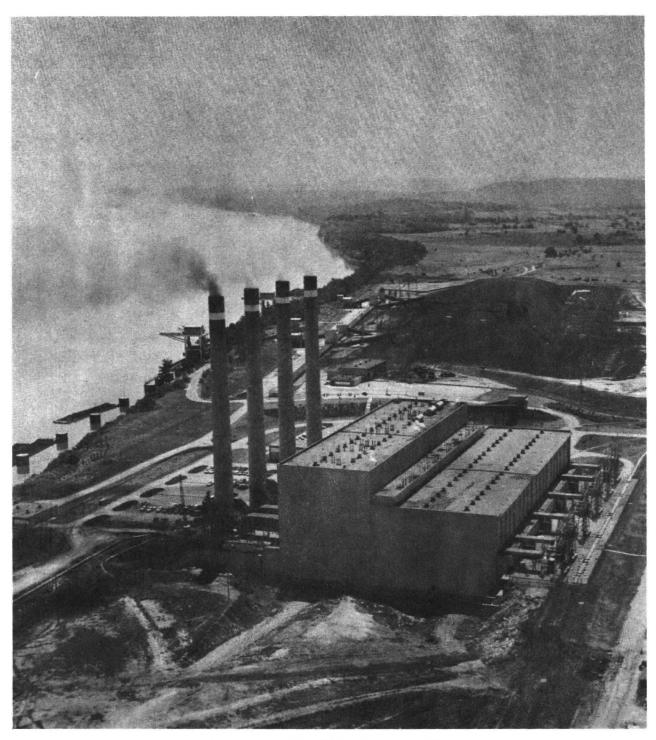


Figure 2. COLBERT STEAM PLANT

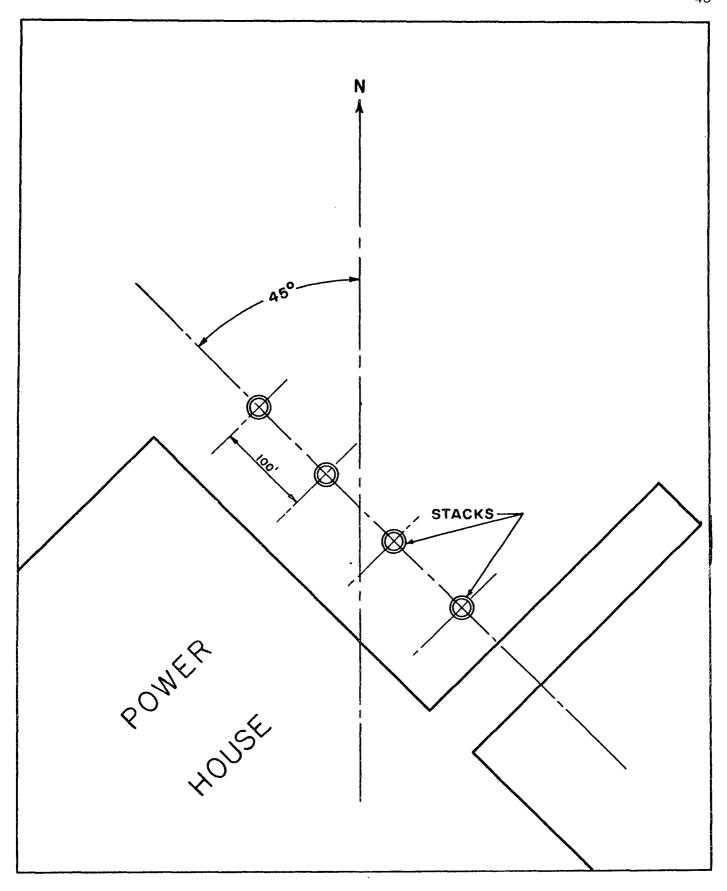


Figure 5. SECTRATION AND CRIMINATION OF STACKS - COLESPI STRAM PLANE

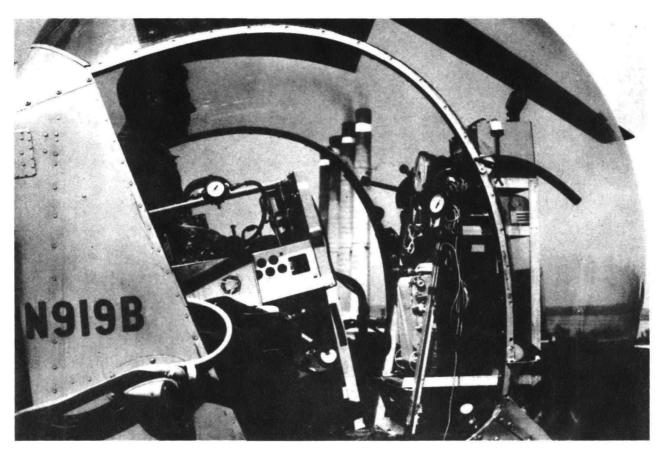


Figure 4. AIR SAMPLING AND AUXILIARY INSTRUMENTS IN HELICOPTER

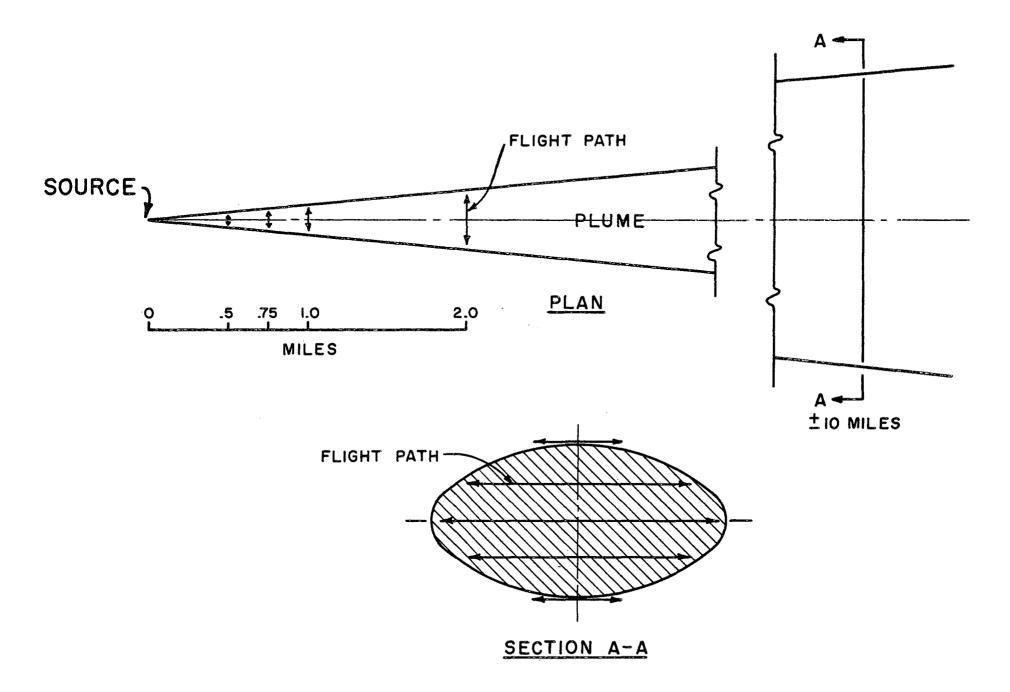


Figure 5. SAMPLE PLAN - INVERSION CONDITIONS

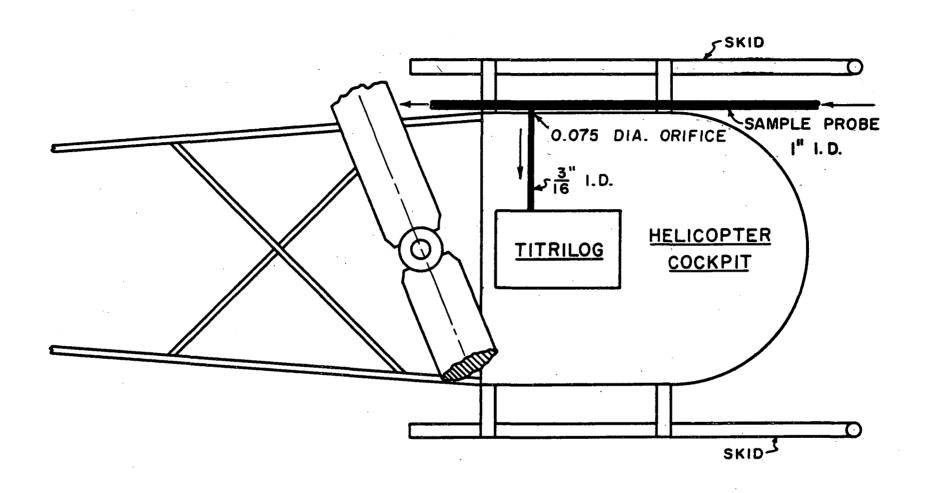
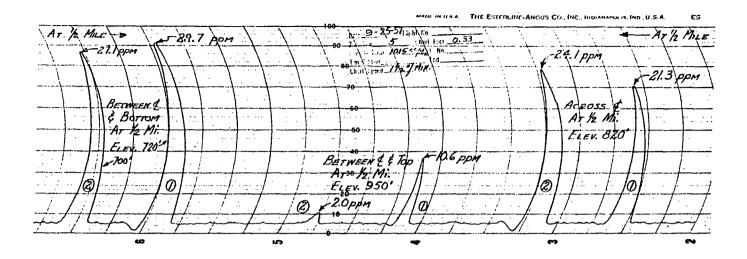
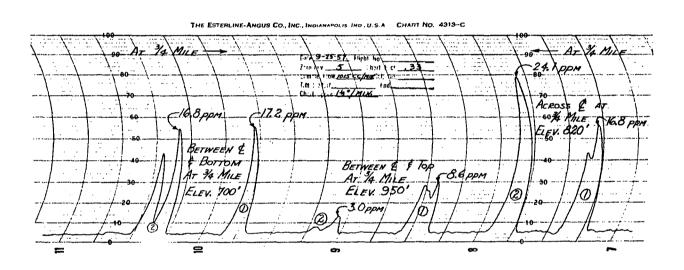


Figure 6. SCHEMATIC PLAN - HELICOPTER AIR SAMPLING EQUIPMENT





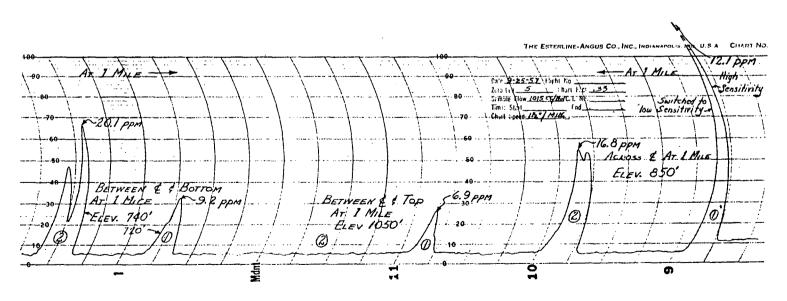


Figure 7. FIRELOG CHART ILLUSTRATING SO2 DISTRIBUTION, DAY 2

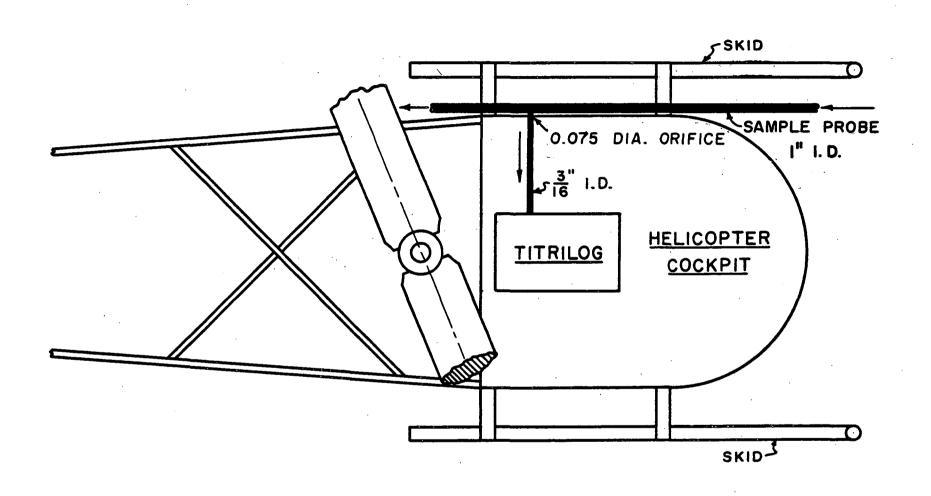
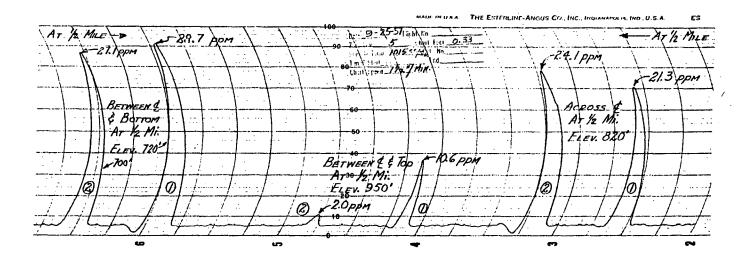
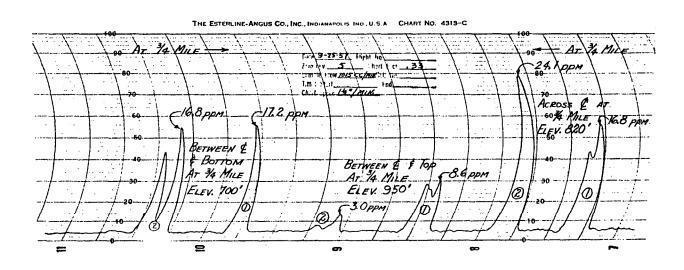


Figure 6. SCHEMATIC PLAN - HELICOPTER AIR SAMPLING EQUIPMENT





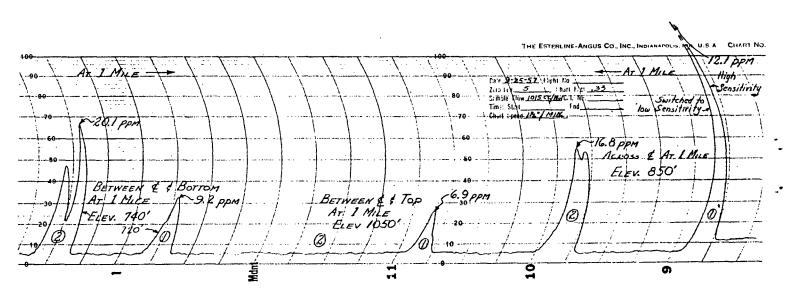
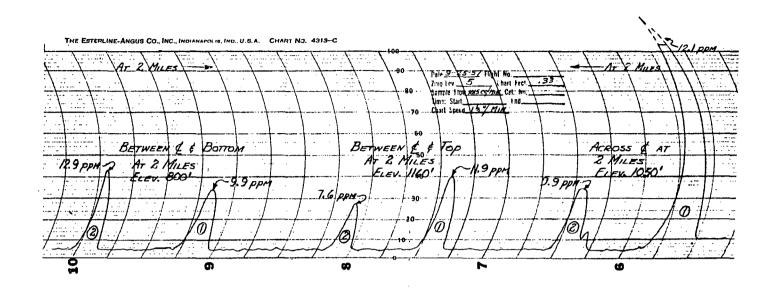
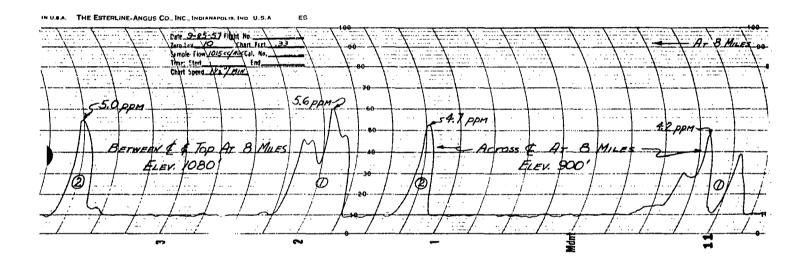


Figure 7. FIRRILG CHART ILLUSTRATING SO2 DISTRIBUTION, DAY 2





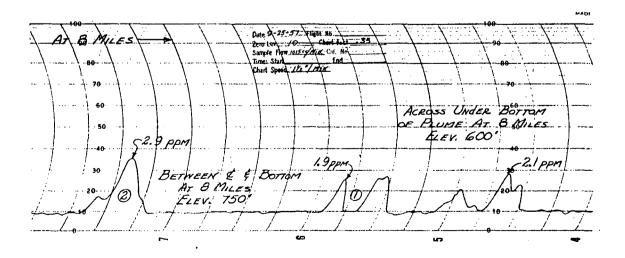


Figure 7. TITRILOG CHART ILLUSTRATING SO2 DISTRIBUTION, DAY 2

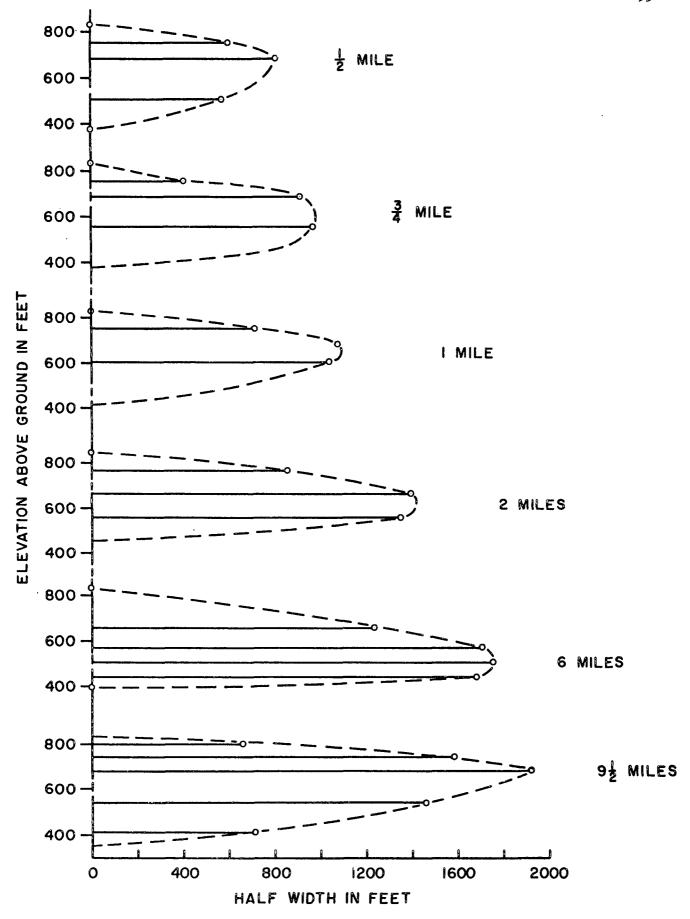


Figure 8. TYPICAL FLOME CROSS SECTION, 9/24/57, DAY 1

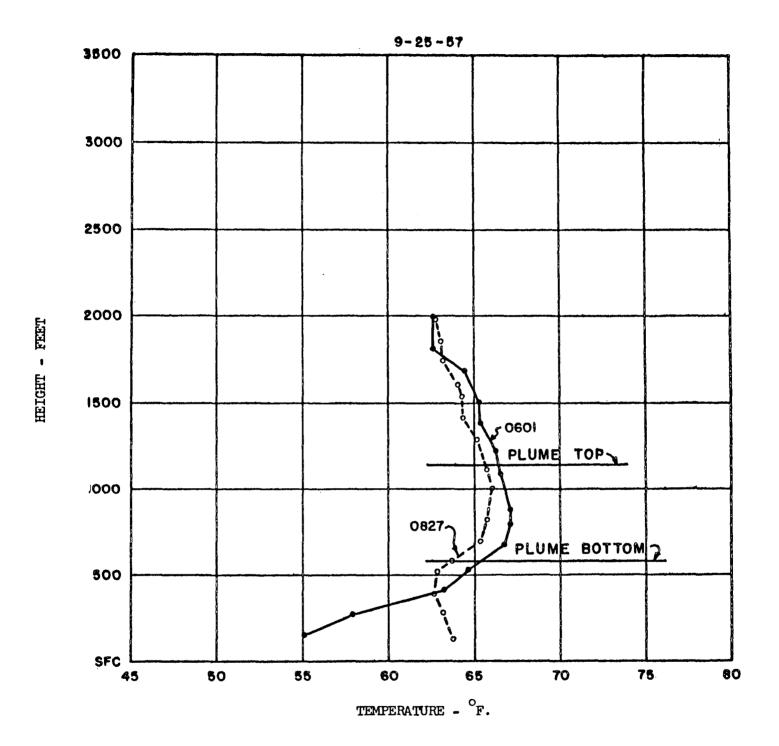


Figure 9. TEMPERATURE PROFILES, DAY 2

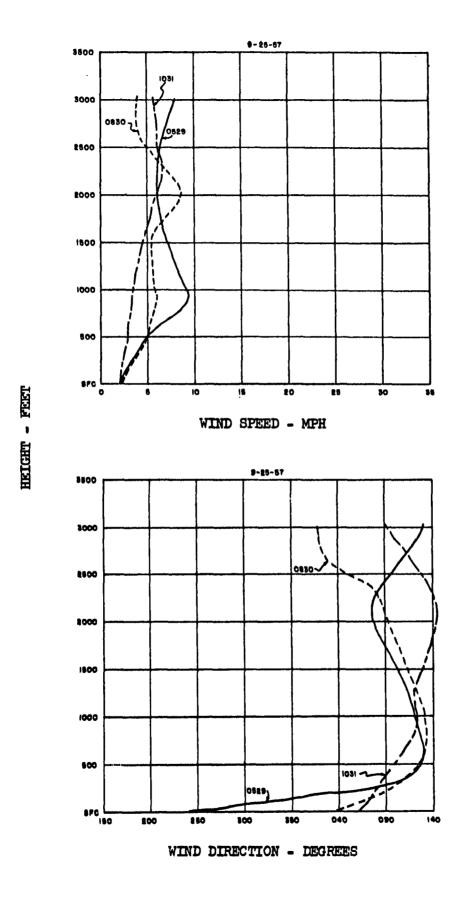


Figure 10. RELATION - WIND SPEED AND WIND DIRECTION TO ELEVATION, DAY 2

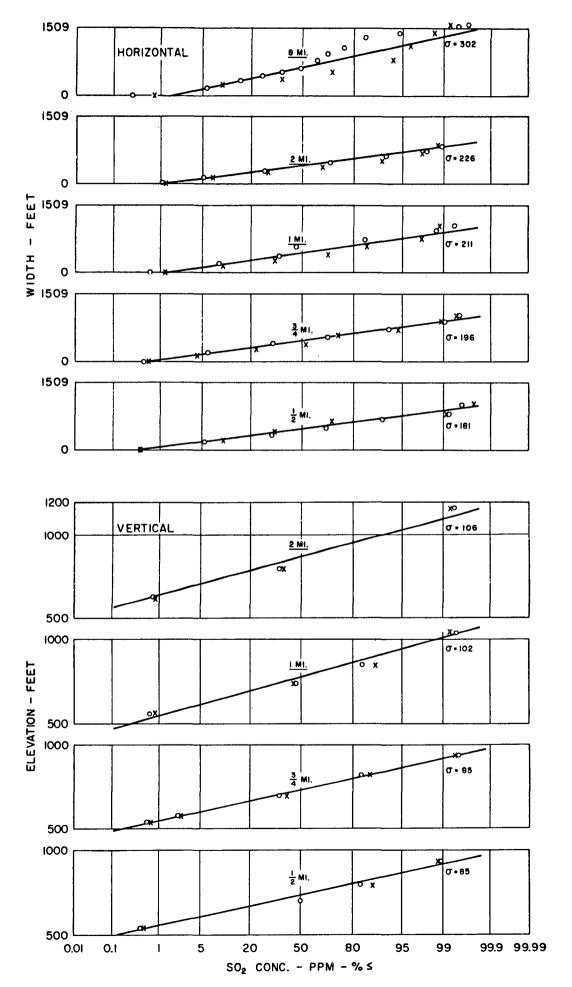


Figure 11. DISTRIBUTION OF SO2 IN PLUME, DAY 2

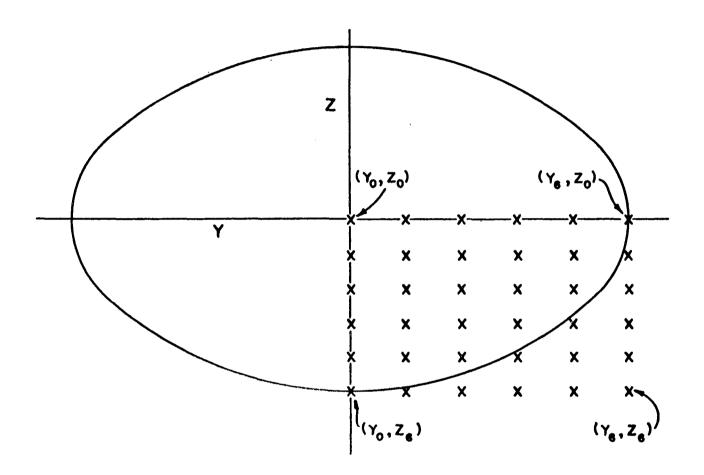


Figure 12. DISTRIBUTION OF POINTS FOR CALCULATED  $SO_2$  CONCENTRATION

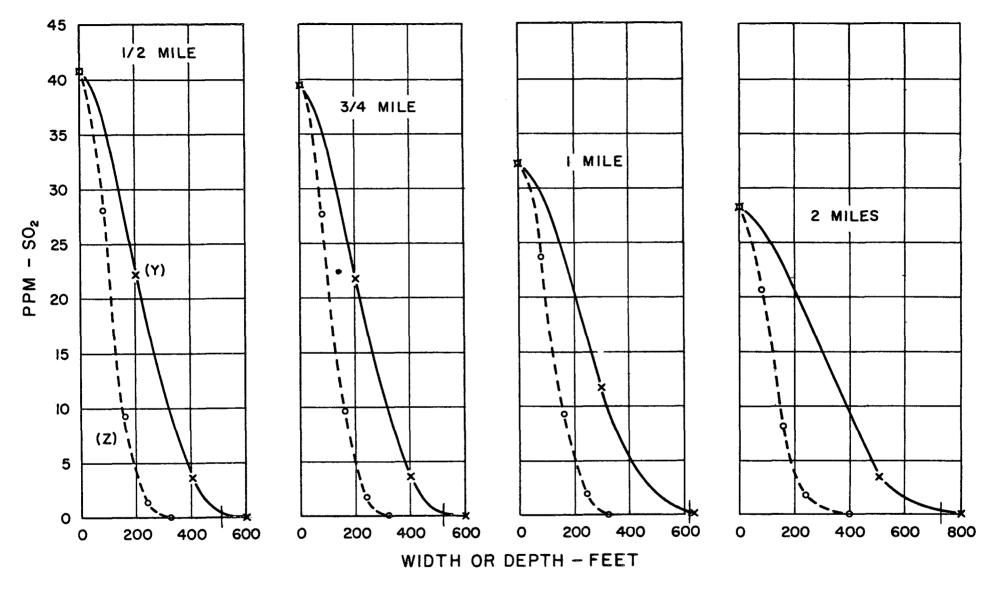


Figure 13. CALCULATED SO2 DISTRIBUTION ALONG Y AND Z AXES, DAY 2

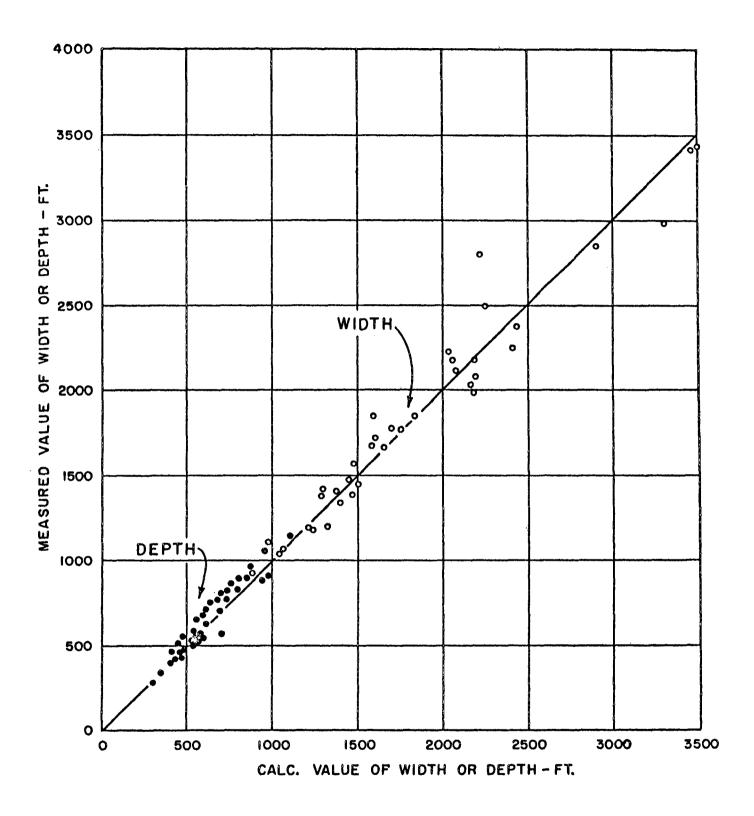


Figure 14. RELATION - MEASURED AND CALCULATED PLUME WIDTH AND DEPTH

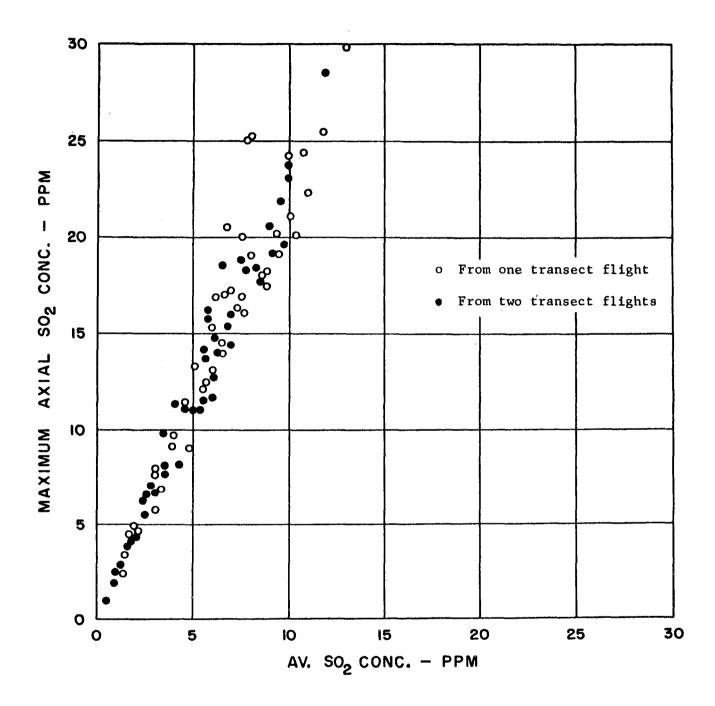


Figure 15. RELATION - MAXIMUM AXIAL CONCENTRATION TO AVERAGE CONCENTRATION ALONG PLUME AXES (AS MEASURED)

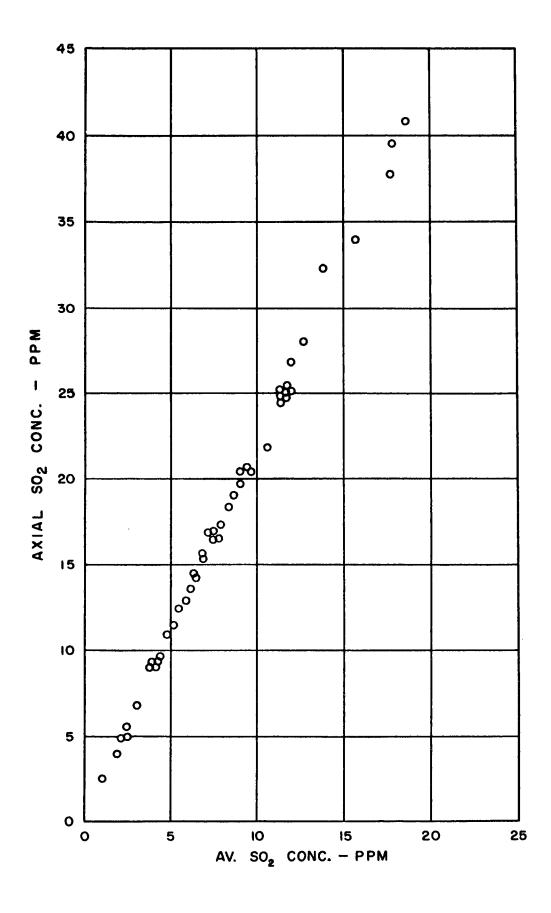


Figure 16. RELATION - CALCULATED AXIAL AND AVERAGE CONCENTRATIONS

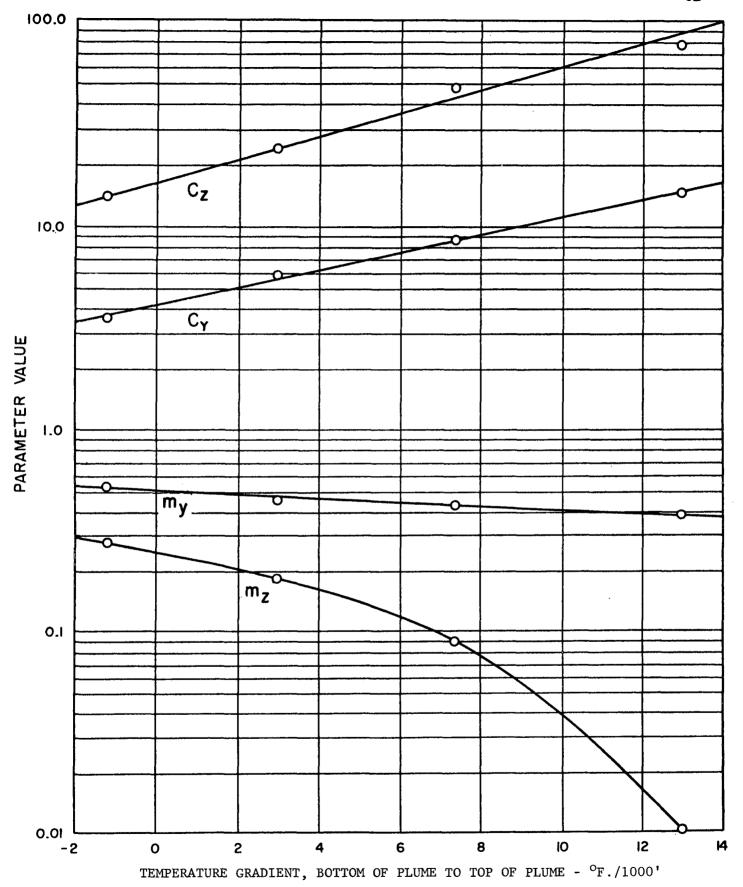


Figure 17. RELATION -  $m_{\mathbf{y}}$ ,  $m_{\mathbf{z}}$ ,  $C_{\mathbf{y}}$ , AND  $C_{\mathbf{z}}$  TO TEMPERATURE GRADIENT (LINE SOURCE)

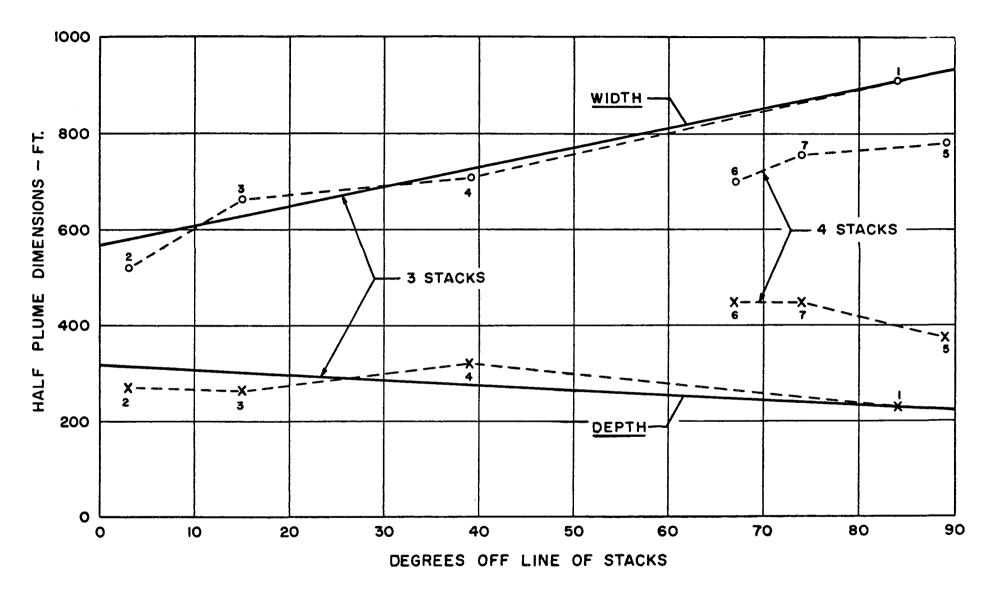
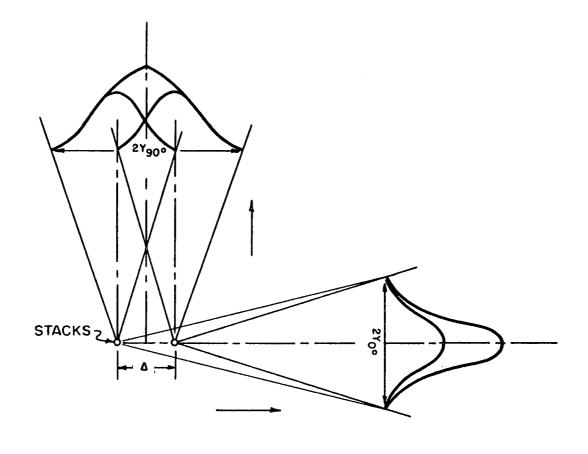


Figure 18. RELATION - PIUME WIDTH AND DEPTH TO WIND DIRECTION AND STACK ALIGNMENT



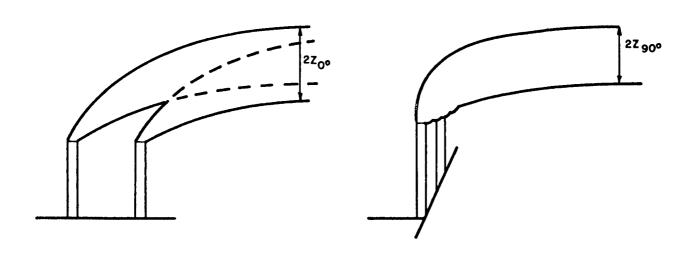


Figure 19. RELATION - PLUME WIDTH AND DEPTH TO WIND DIRECTION (FROM TWO OR MORE STACKS)

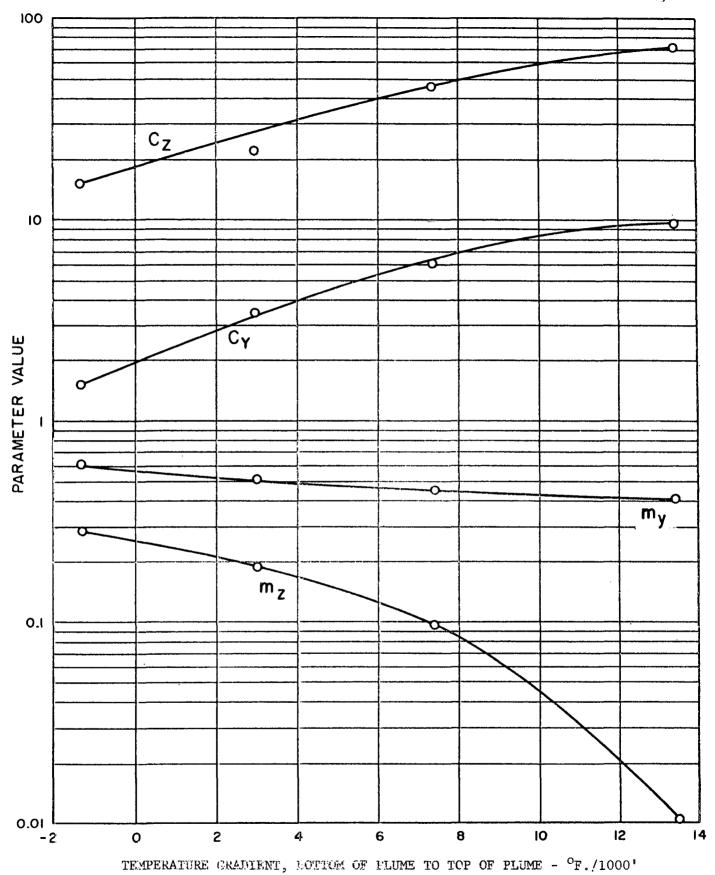
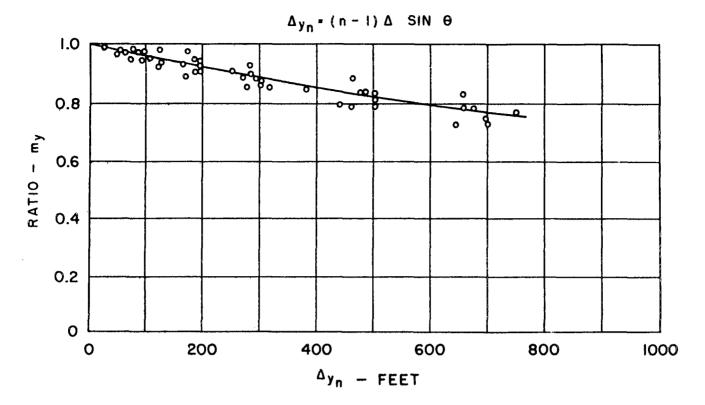
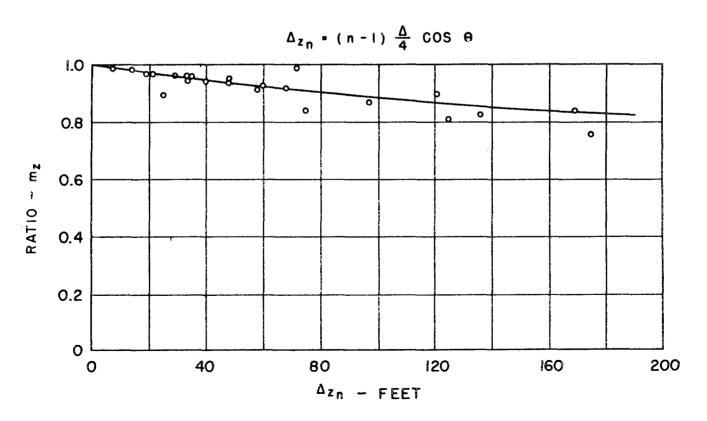


Figure 20. BEST ESTIMATE  $m_y$ ,  $m_z$ ,  $c_y$ ,  $c_z$  - POINT SOURCE



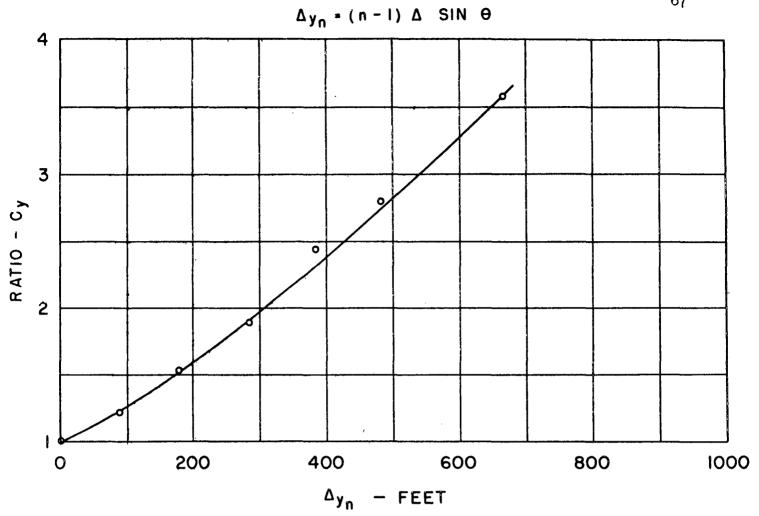
RELATION - RATIO OF  $\mathbf{m}_y$  TO  $\triangle_{y_n}$ 



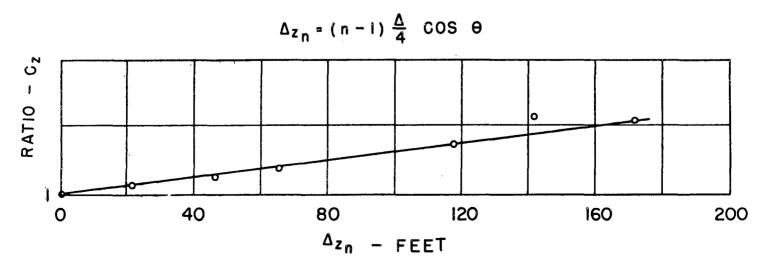
RELATION - RATIO OF  $m_z$  TO  $\triangle_{z_n}$ 

Figure 21





RELATION - RATIO OF  $c_y$  to  $\triangle_{y_n}$ 



RELATION - RATIO OF  $c_{\mathbf{z}}$  TO  $\Delta_{\mathbf{z}_n}$ Figure 22

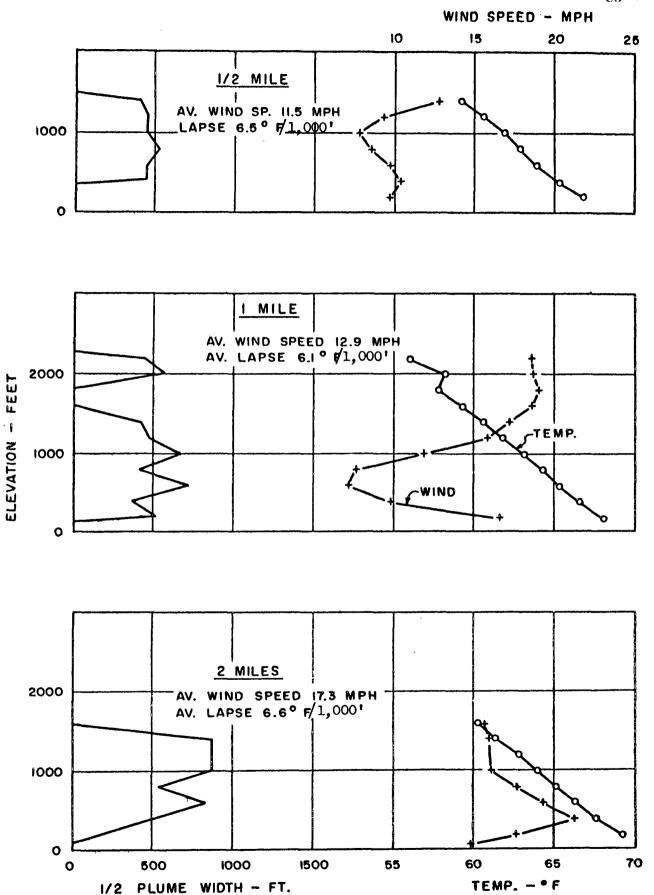


Figure 23. PLUME CROSS SECTION, TEMPERATURE PROFILE, AND WIND SPEED PROFILE, DAY 7

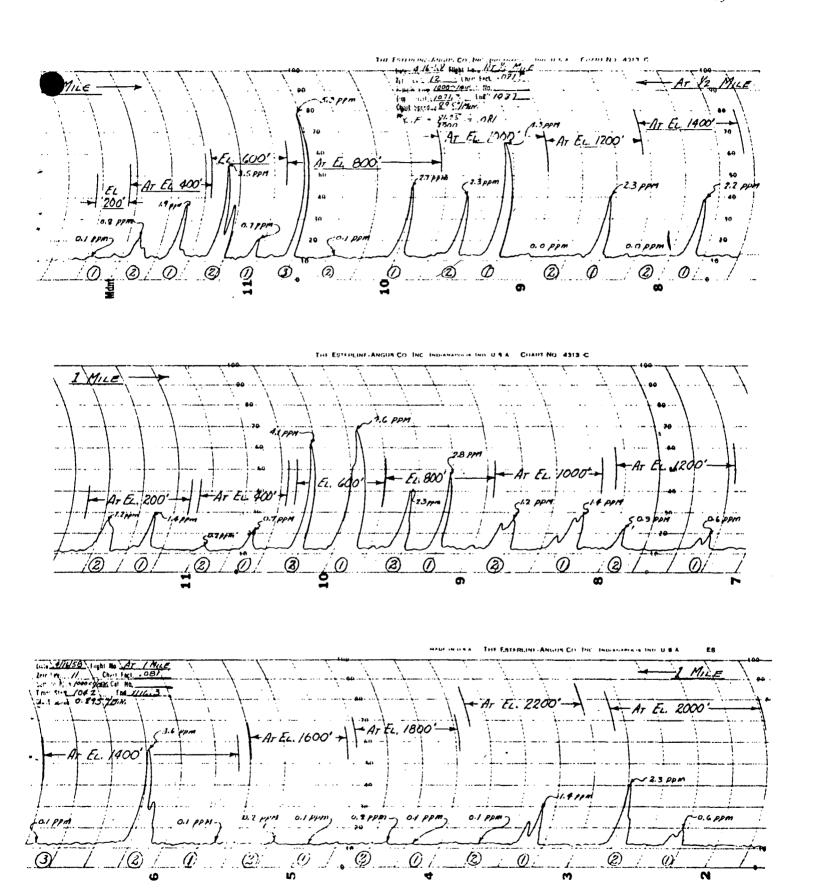
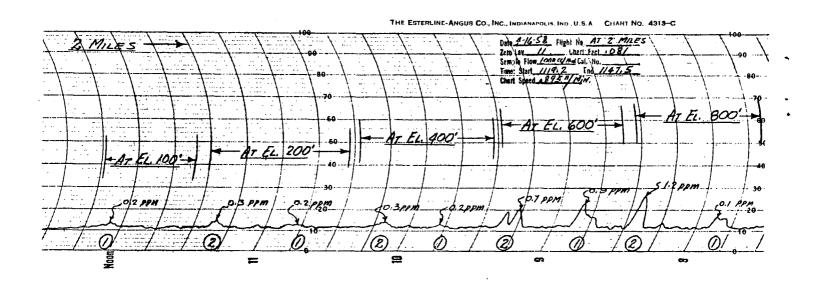


Figure 24. TITRILOG CHARTS ILLUSTRATING SO2 DISTRIBUTION, DAY 7



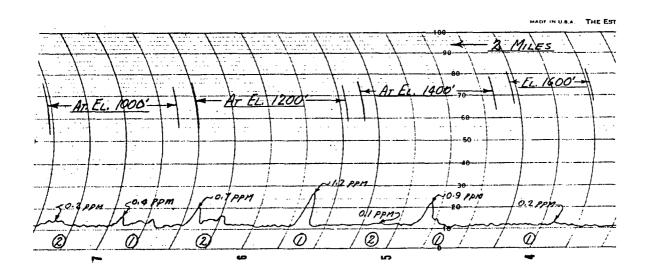


Figure 24. TITRILOG CHARTS ILLUSTRATING SO2 DISTRIBUTION, DAY 7

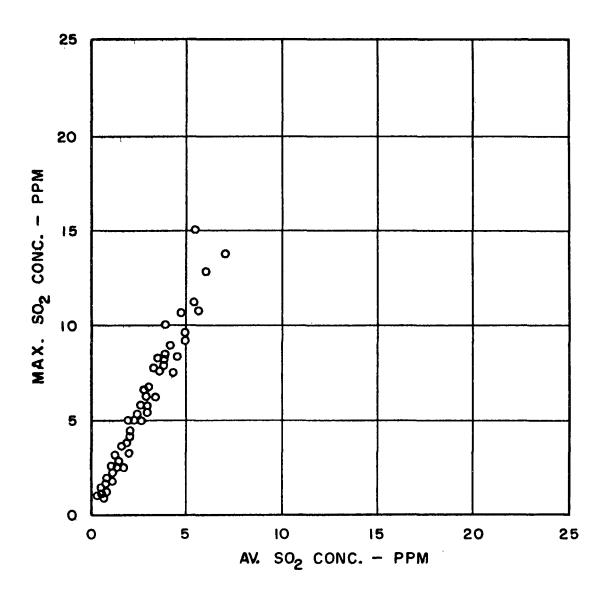


Figure 25. RELATION - MAXIMUM TO AVERAGE SO<sub>2</sub> CONCENTRATION

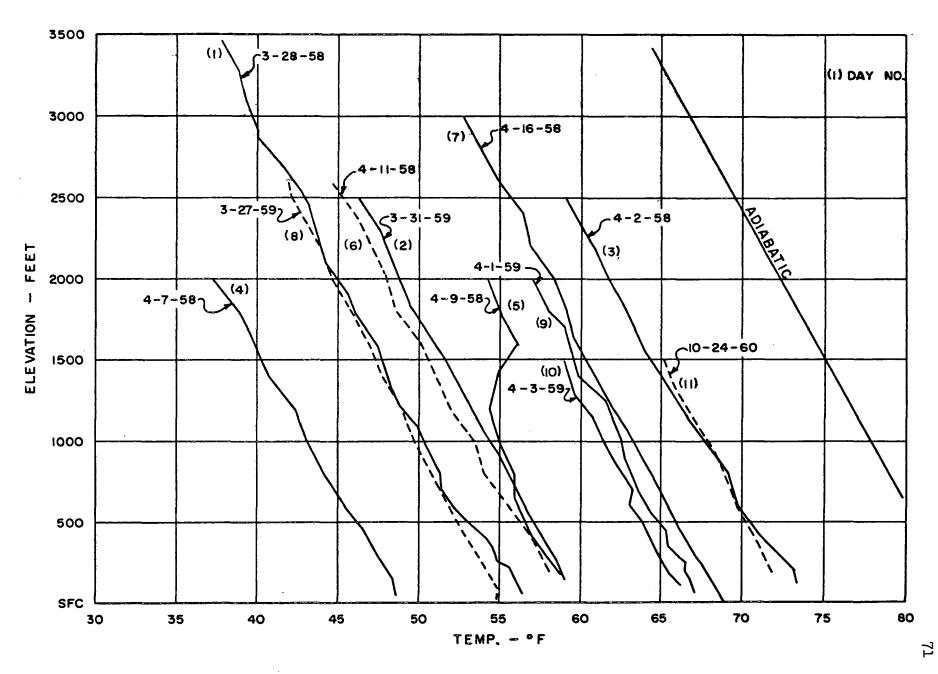


Figure 26. AVERAGE CHANGE OF TEMPERATURE WITH ELEVATION - EACH SAMPLING FERIOD

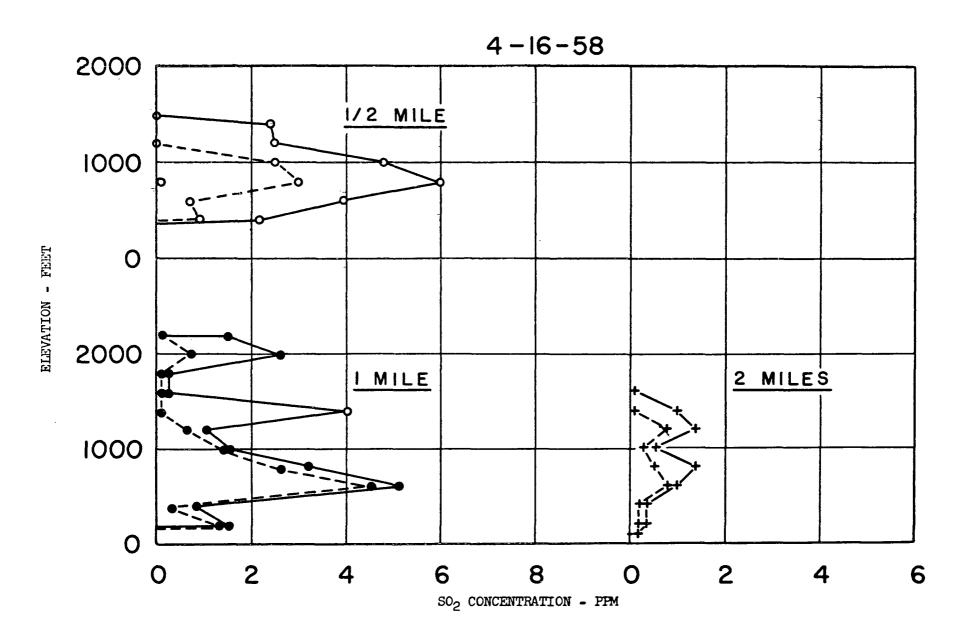


Figure 27. PROFILE OF SO<sub>2</sub> ALONG Z AXIS, DAY 7

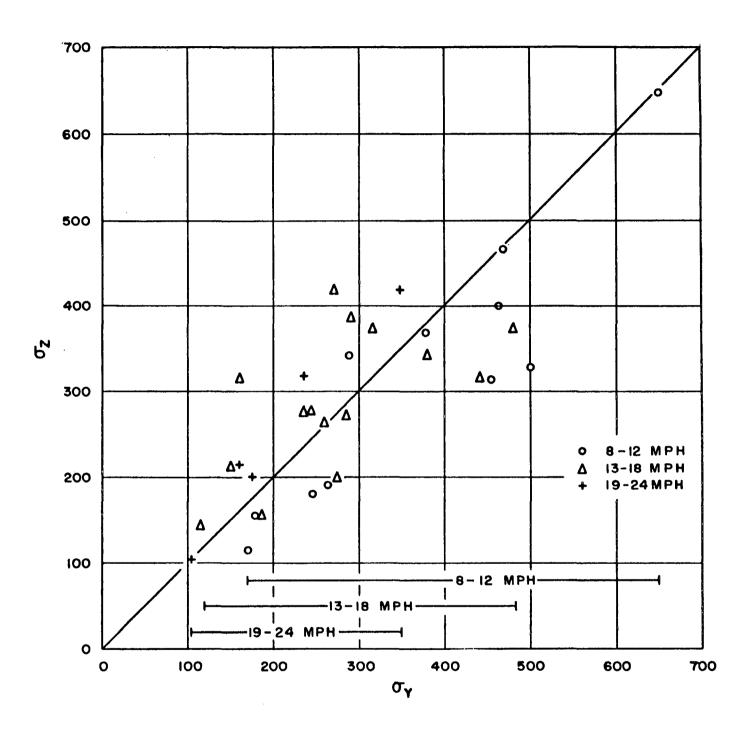


Figure 28. RELATION -  $\sigma_{\mathbf{y}}$  AND  $\sigma_{\mathbf{z}}$ 

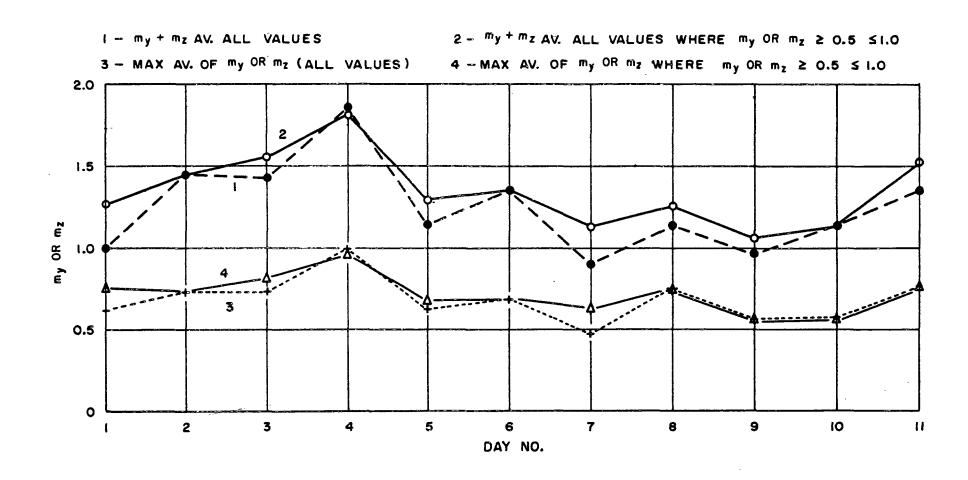


Figure 29. m. Alle m $_{\mathbf{z}}$  - BACH SAMPLING DAY

LOG. Cy = -0.101167 - 0.015439 X WIND SPEED STANDARD ERROR OF ESTIMATE = 0.054586 CORRELATION COEFFICIENT = 0.87

 $m_y = m_z = 0.75$ 

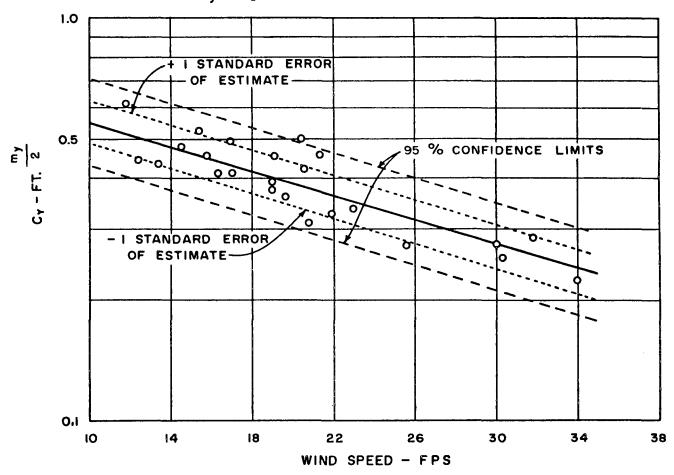


Figure 50. STATISTICAL ANALYSIS - RELATION  $\mathbf{c}_{\mathbf{y}}$  TO WIND SPEED

LOG  $C_z$  = -0.141965 -0.011797 X WIND SPEED STANDARD ERROR OF ESTIMATE = 0.0561885 CORRELATION COEFFICIENT = 0.80  $m_y$  =  $m_z$  = 0.75

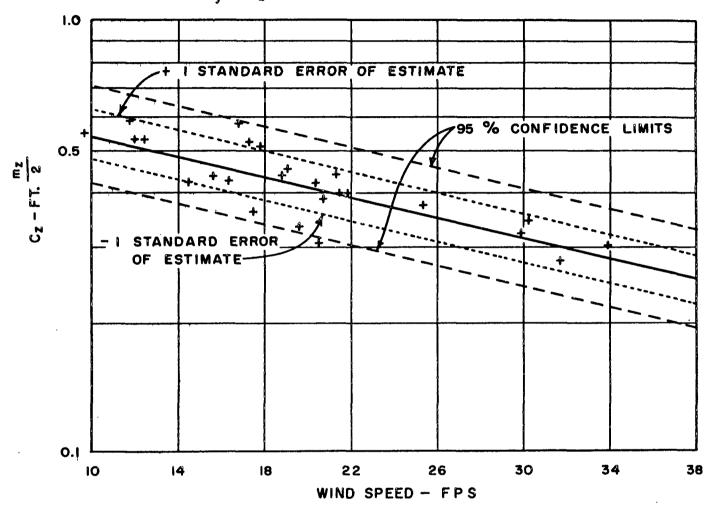


Figure 31. STATISTICAL ANALYSIS - RELATION  $\mathbf{C_z}$  TO WIND SPEED

LOG C<sub>Y</sub> OR C<sub>Z</sub> = -0.121340 - 0.013635 X WIND SPEED STANDARD ERROR OF ESTIMATE = 0.0578577 CORRELATION COEFFICIENT = 0.82

 $m_y = m_z = 0.75$ 

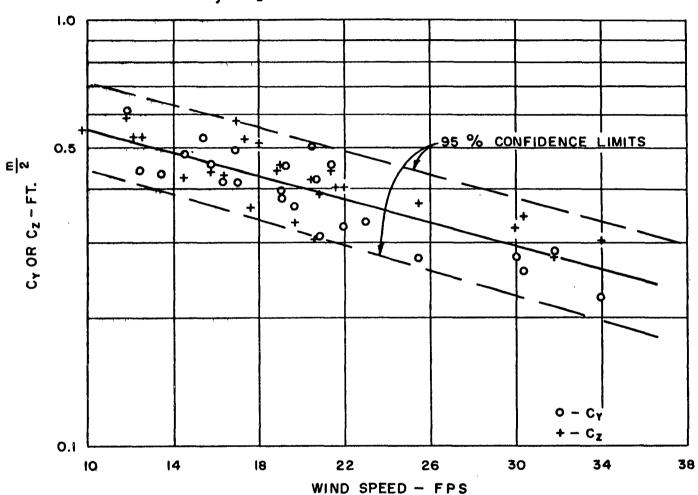


Figure 32. STATISTICAL ANALYSIS - RELATION  $\mathbf{C}_{\mathbf{y}}$  AND  $\mathbf{C}_{\mathbf{z}}$  TO WIND SPEED

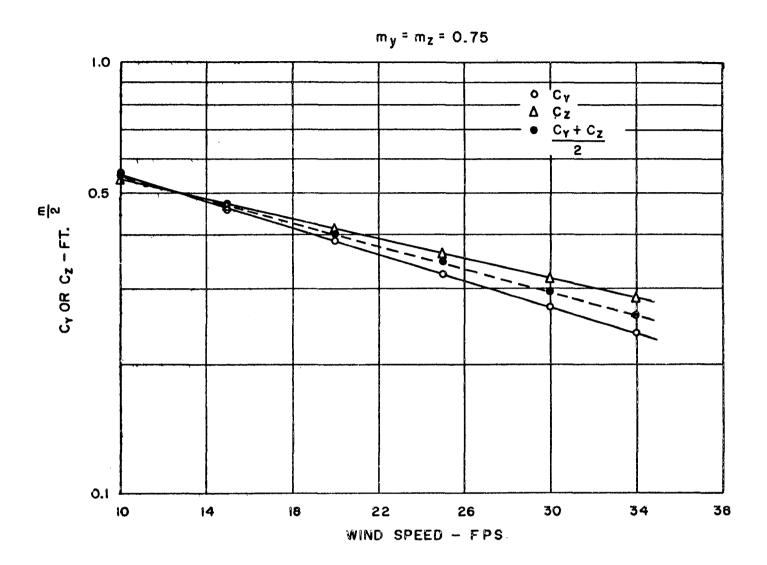


Figure 33. RELATION -  $C_y$ ,  $C_z$  AND AVERAGE C, FT.  $\frac{m}{2}$ , TO WIND SPEED

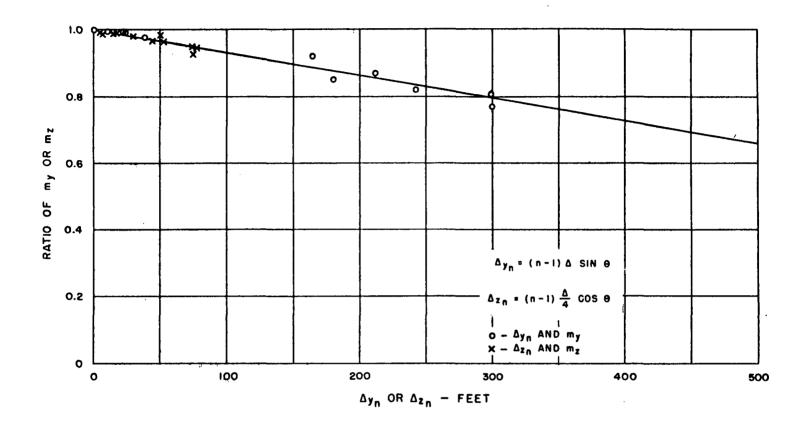


Figure 34. RELATION - RATIO OF  ${\rm m_y}$  AND  ${\rm m_z}$  FOR LINE AND POINT SOURCES TO  $\triangle_{y_n}$  AND  $\triangle_{z_n}$ 

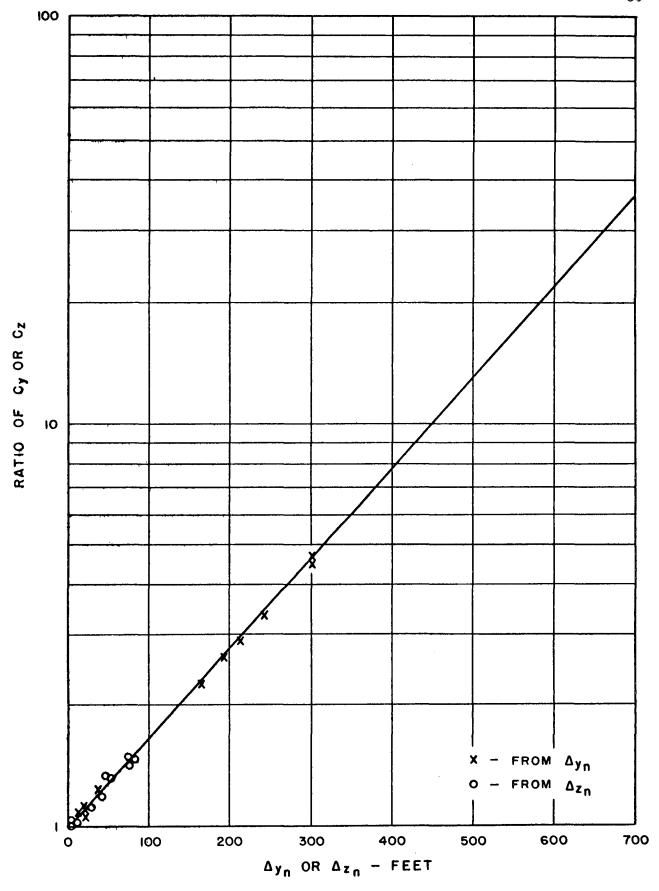


Figure 35. RELATION - RATIO OF  ${\tt C_y}$  AND  ${\tt C_z}$  FOR LINE AND POINT SOURCES TO  $\triangle_{y_n}$  AND  $\triangle_{z_n}$ 

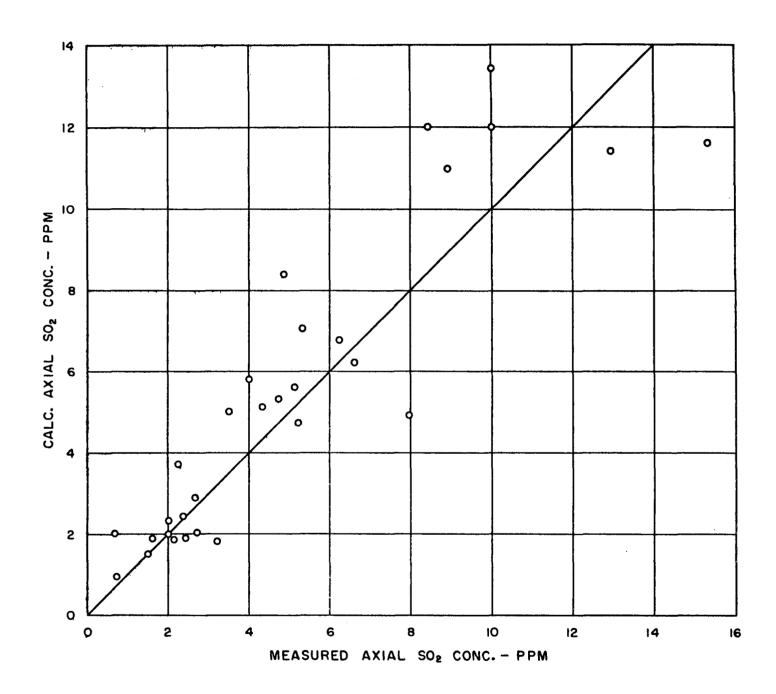


Figure 36. RELATION OF CALCULATED AXIAL SO<sub>2</sub> CONCENTRATION TO MEASURED AXIAL SO<sub>2</sub> CONCENTRATION

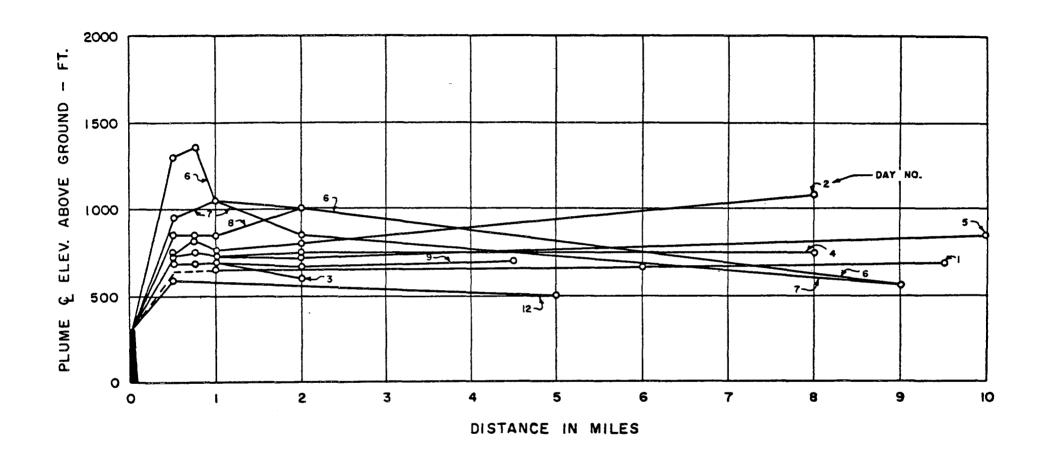


Figure 37. OBSERVED PLUME CENTERLINE ELEVATION WITH DISTANCE - INVERSION CONDITIONS

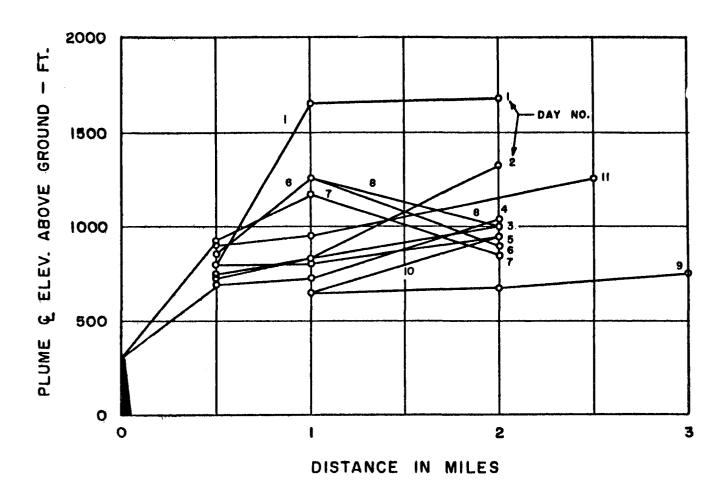


Figure 38. AVERAGE OBSERVED PLUME CENTERLINE ELEVATION WITH DISTANCE - LAPSE CONDITIONS

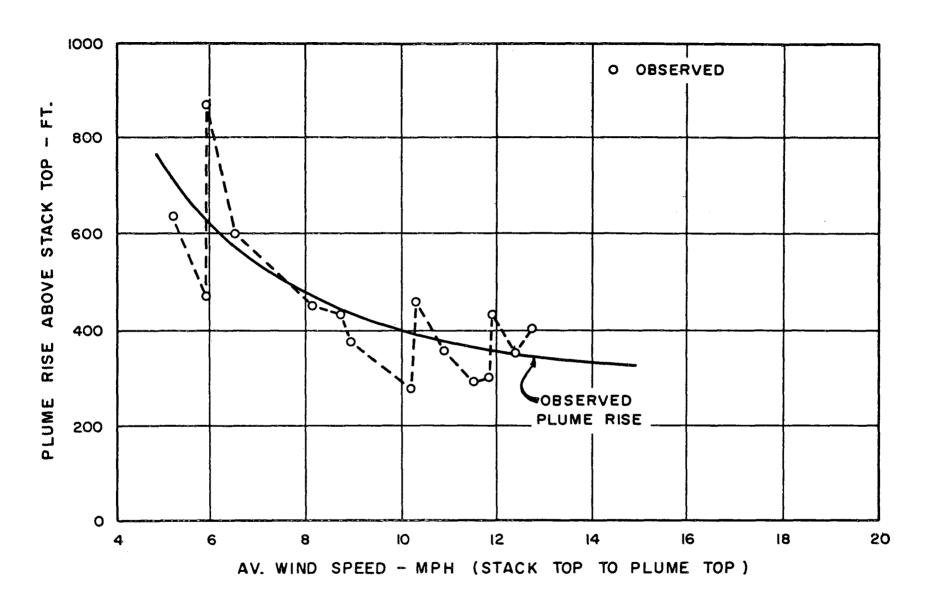


Figure 39. RELATION OF PLUME RISE TO WIND SPEED - INVERSION CONDITIONS

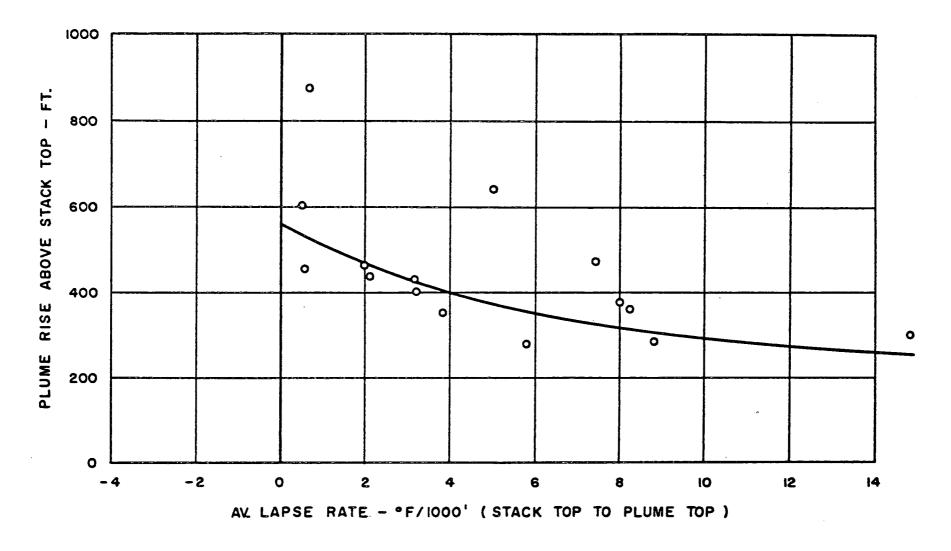


Figure 40. RELATION OF PLUME RISE TO TEMPERATURE GRADIENT, STACK TOP TO PLUME TOP - INVERSION CONDITIONS

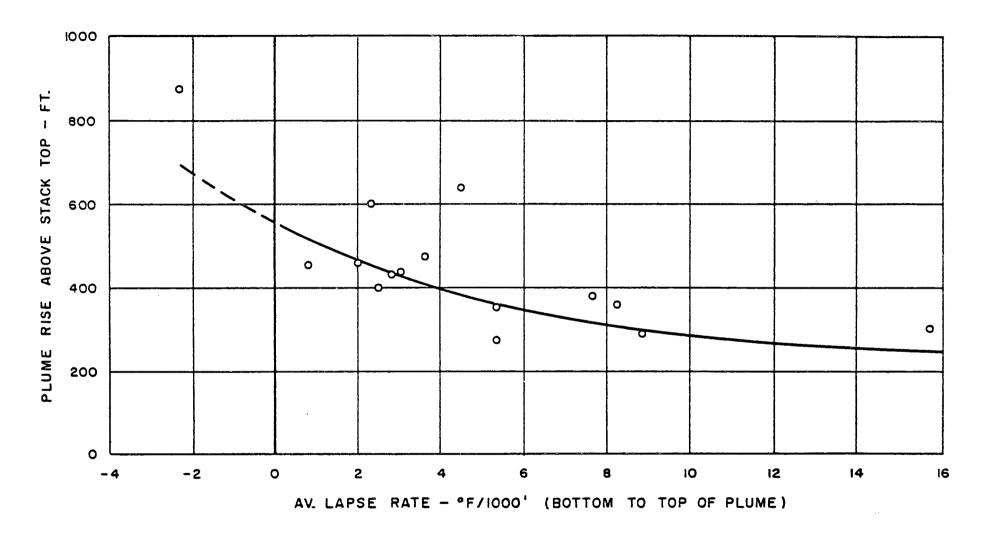


Figure 41. RETATION OF FIUME RISE TO TEMPERATURE GRADIENT, IN PLUME - INVERSION CONDITIONS

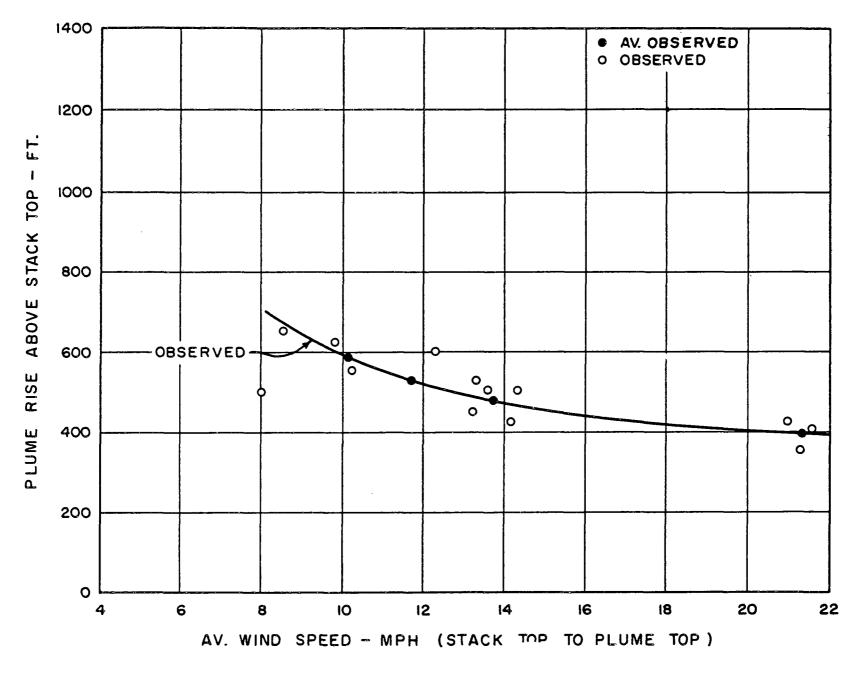


Figure 42. RELATION OF AVERAGE PROME RISE TO WIND SPEED AT 1/2 AND 1 MILE FROM SOURCE - LAPSE CONDITIONS

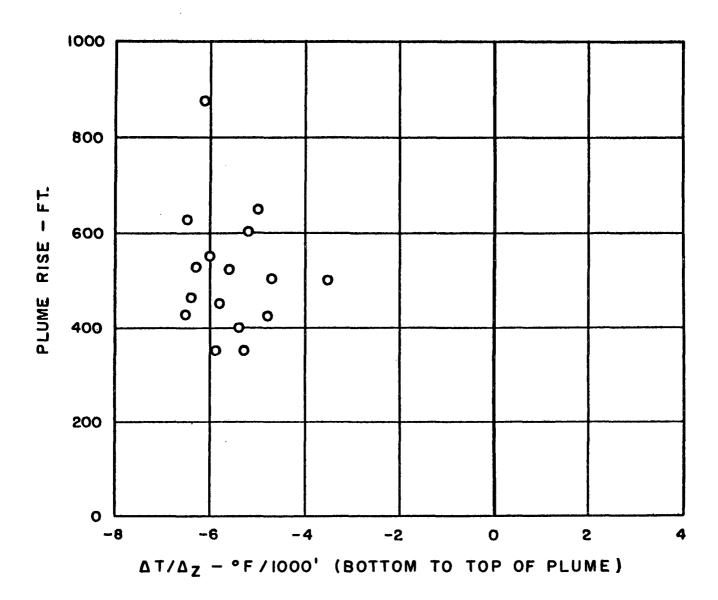


Figure 43. RELATION OF AVERAGE PLUME RISE AT 1/2 AND 1 MILE TO TEMPERATURE GRADIENT - LAPSE CONDITIONS

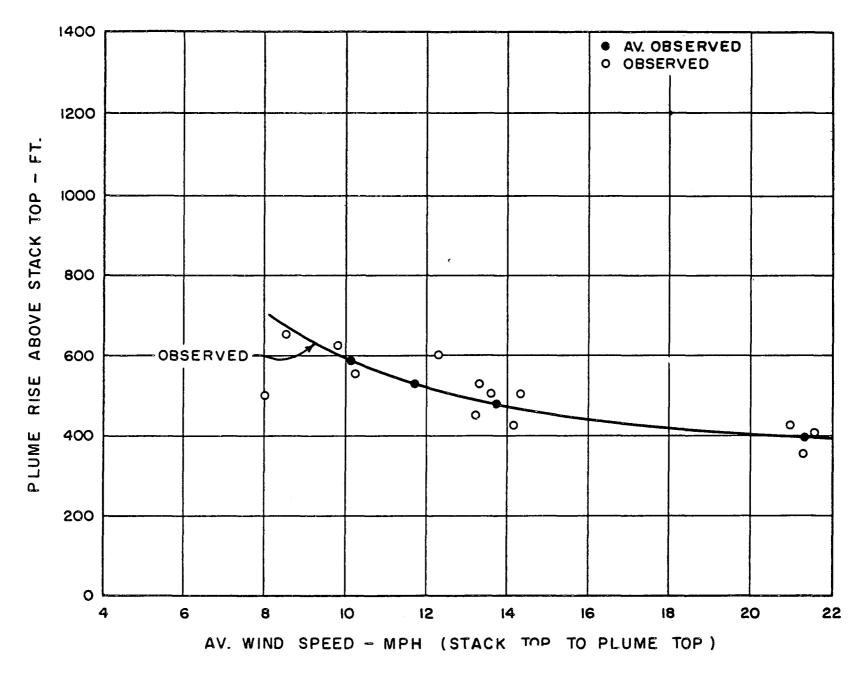


Figure 42. RELATION OF AVERAGE PLUME RISE TO WIND SPEED AT 1/2 AND 1 MILE FROM SOURCE - LAPSE CONDITIONS

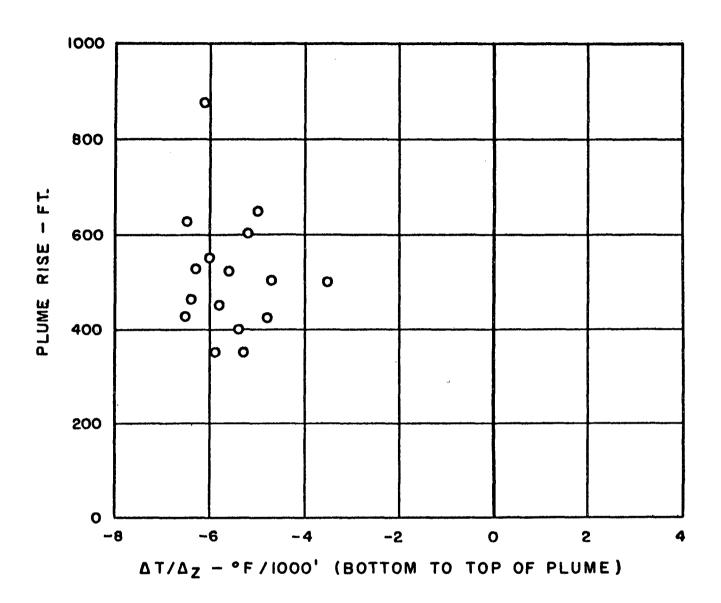


Figure 43. RELATION OF AVERAGE PLUME RISE AT 1/2 AND 1 MILE TO TEMPERATURE GRADIENT - LAPSE CONDITIONS

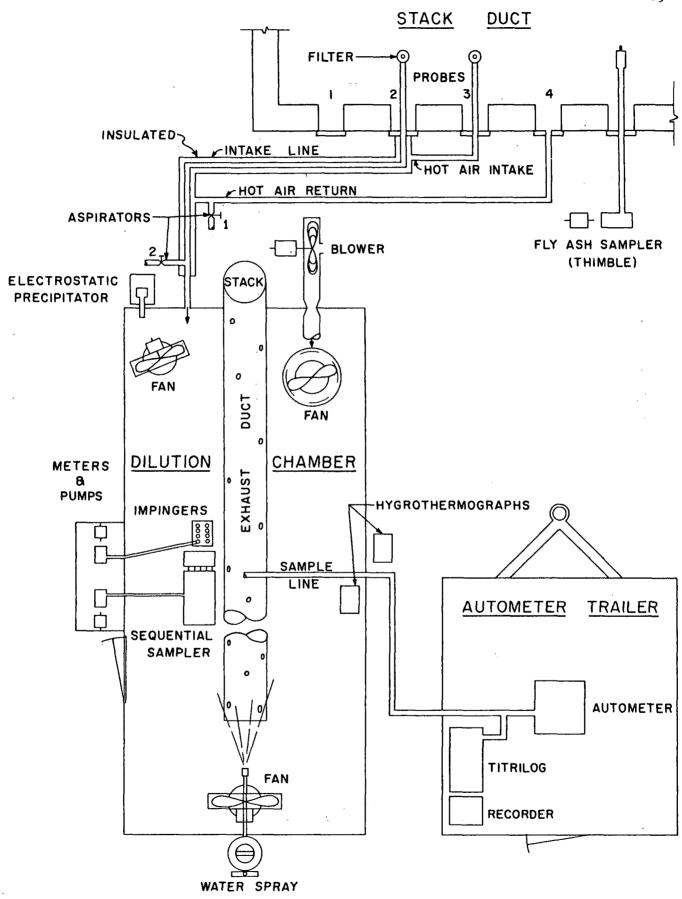
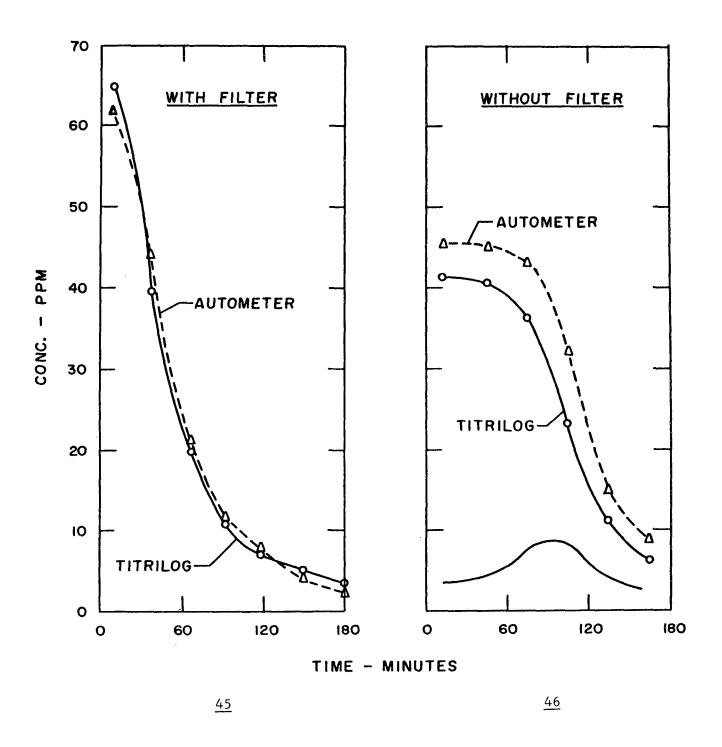


Figure 44. FLUE GAS DILUTION AND SAMPLING FACILITIES



Figures 45 and 46. RELATION OF AUTOMETER AND TITRILOG DATA FROM DILUTION CHAMBER

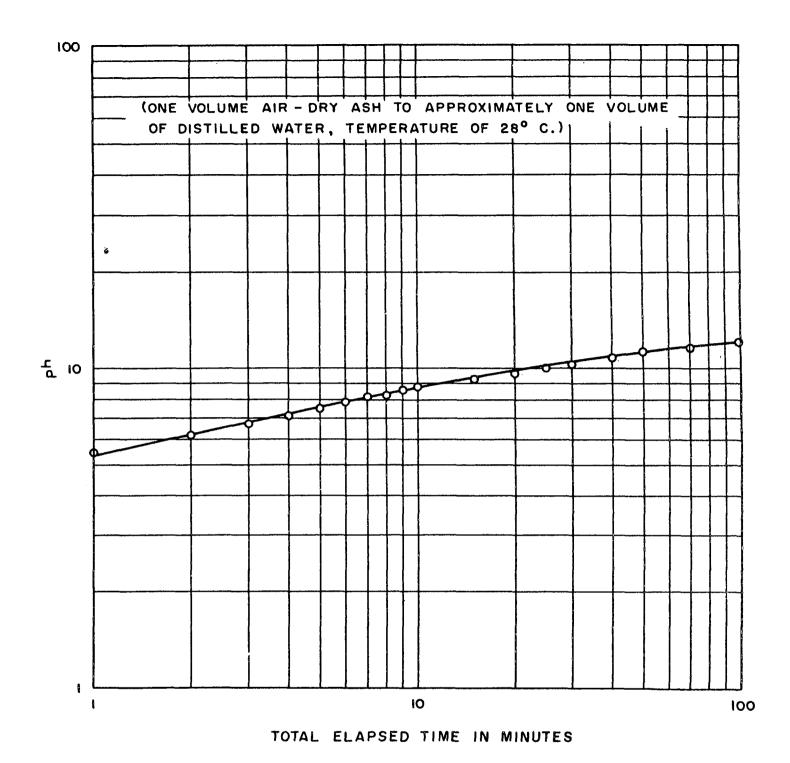
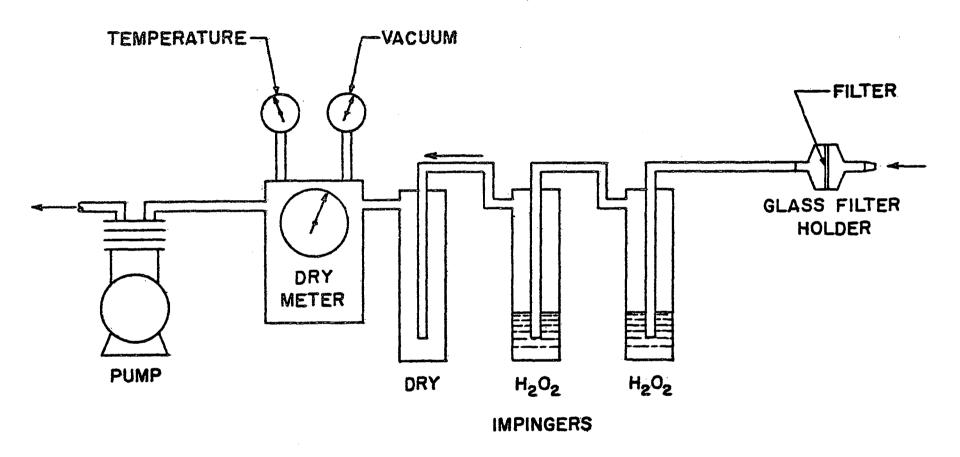


Figure 47. SHIFT IN pH OF FLY ASH WITH TIME



## NOMENCLATURE - DIFFUSION EQUATIONS

Symbol		System of Units
$\mathtt{c}_{\mathtt{y}}$	Diffusion coefficients along y axis	ft. or ft. e
C <sub>z</sub>	Diffusion coefficients along z axis	ft. or ft.
exp	The value e	2.718
m	Stability parameter	dimensionless
<sup>m</sup> y	Stability parameter along y axis	dimensionless
$m_{\mathbf{Z}}$	Stability parameter along z axis	dimensionless
n	Number of stacks (equations 16 and 17)	-
ବ	SO <sub>2</sub> emission rate	$ft.^3/sec.$
u	Wind speed	mph
<u>x</u>	SO <sub>2</sub> concentration	ppm
$\overline{\underline{\mathbf{x}}}_{\mathtt{max}}$	Maximum SO <sub>2</sub> concentration	ppm
x	Distance downwind from source	ft.
<sup>x</sup> g	Distance from centerline along normal distribution curve (Gaussian)	ft.
У	Crosswind distance from centerline of plume	ft.
z	Vertical distance from centerline of plume	ft.
σ	Standard deviation of normal distribution curve (Gaussian)	ft.
$\sigma_{\mathbf{y}}$	Standard deviation along y axis	ft.
$\sigma_{f z}$	Standard deviation along z axis	ft.
θ	Angular difference between plume direction and stack alignment	degrees
Δ	Distance between stacks	ft.