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FURTHER STUDIES
OF JET CONTRAIL NUCLEATION
AND VISIBILITY CHARACTERISTICS

by

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and

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Contract No. F19628-67-C-0283
Project No. 8679
Task No. 867902
Work Unit No. 86790201

FINAL REPORT

June 1967 through May 1968

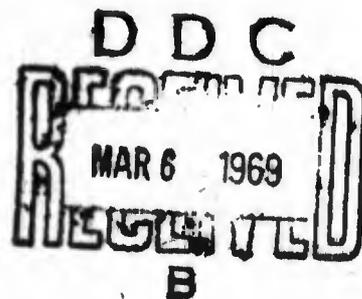
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Contract Monitor: Seymour J. Birstein
Meteorology Laboratory

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Prepared for

AIR FORCE CAMBRIDGE RESEARCH LABORATORIES
OFFICE OF AEROSPACE RESEARCH
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Abstract

This report is a continuation of the final report for contract AF19(628)-5193. The assumptions and assertions of that report are examined, and estimates made of how these may affect the worth and accuracy of numerical predictions of contrail properties. A special type of analytic-experimental analysis is proposed to ferret out nonthermodynamic variables, and is applied to flight test data. Following the analysis, further computer simulation is explored. Additional data lead to bounds on the values of heretofore unspecifiable parameters. Further experimental study is indicated. The data is presented in terms of the dependence of contrail formation on ambient temperature and relative humidity.

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I. Introduction

The current study has been done in fulfillment of Air Force Contract F19628-67-C-0283. Its purpose is the formulation and study in depth, using computer simulation techniques, of mathematical models to describe heterogeneous water nucleation in jet exhaust contrails. In particular, general equations for contrail properties are sought as a function of system variables. Information provided in the final report for contract AF19(628)-5193(1)¹ is the base for which this report is a continuation.

For this work, the authors took the approach of developing:

- (a) an analysis of the assumptions and assertions of contract AF19(628)-5193,
- (b) a set of experimental and statistical, rather than purely theoretical, techniques for quantitative prediction of contrail characteristics, and
- (c) a further theoretical treatment involving contrail simulation by computer including the effects of relative humidity and ambient temperature.

Sections II, III, and IV of this report deal with parts (a), (b), and (c) above, respectively. Conclusions are presented in Section V.

Typical fuels used by jets contain large amounts of hydrogen (C_nH_{2n}). The exhaust gases of jets under certain conditions tend to be super-saturated with respect to water vapor, and cooling of the gases produces water droplets known as contrails. Under normal circumstances, water vapor condenses only on foreign particles, and the fuel exhaust usually contains enough such particles for condensation to take place.

The reason for adding SO_3 to the fuel is the addition to the exhaust of much larger numbers of extremely small hygroscopic nuclei than are normally present. The water vapor formed by the burning of a hydrocarbon fuel will then preferentially condense on these nuclei rather than on the fewer, larger, and less hygroscopic nuclei normally produced in the combustion process. The droplets formed will thus be smaller than those formed in normal contrail formation.

The objective of this study is the determination of those amounts of additive which produce an invisible contrail for a given set of ambient conditions and engine characteristics. Such a contrail contains a large number of particles sufficiently small that the total cannot be seen.

¹ Numbers in square brackets refer to similarly numbered items in the bibliography.

It is an observed fact that contrail visibility requires a minimum density (in space) of particles whose radius exceeds one - quarter micron. Calling such a particle a "large particle", the objective may then be restated as: Under given conditions, find an additive - fuel combination that produces a contrail in which the density (in space) of large particles is less than the critical density.

II. Review of Previous Research

After investigating possible chemical combinations in the jet exhaust due to oxidation of additive, the authors in their final report for contract AF19(628)-5193, describe the formation of droplets. Thermodynamic considerations, involving ambient conditions as parameters, predict the size and rate of formation of droplets, and knowledge of the physical chemistry of additive end products leads to specific numerical results.

Of course, the worth and accuracy of numerical predictions of a complicated physical simulation are no more valid than the worth and accuracy of the many assumptions involved. In AF19(628)-5193, assumptions fall naturally into categories associated with properties of the exhaust wake, the exhaust products, the nucleation process, and the contrail itself.

Description of the exhaust wake has meaning only to the extent that turbulence has not completely dissipated the central core of the wake. Photographs [2] of actual jet flights at altitude 40,000 ft. indicate a distance behind the jet of about 100 to 150 ft. before contrail formation (the range decreases at higher altitudes), with maintenance of the general shape for several hundred additional feet.

Forstall and Shapiro [3] have measured and analytically described the wake temperature distribution over an interval of several hundred feet, but the measurements were taken at ground level; as a result, temperature distribution is not verifiably stated for high altitudes, although the ground-level description is probably adequate after some mathematical adjustments. Accordingly, changes reflecting altitude were made in two parameters of the temperature distribution function employed in the simulation:

- (1) an exponent in the temperature - dispersal relation (increasing which has the numerical effect of increasing turbulence), and
- (2) the distance over which the authors' estimate of exhaust temperature off the centerline remains in effect.

Computation of virtually every property of the contrail process requires knowledge of the temperature distribution, and it can not be overstressed that only crude estimates of it are available. Therefore, in order to provide results in which more confidence might be placed, numerous trial simulations were performed where the two parameters were varied over wide ranges. The results showed almost no dependence of contrail visibility on these parameters. The values 1.0 and 300 ft. for the respective parameters were then used during all other runs, with the exception of an occasional run which "tested" other values. The results however, did not change.

Specification of the exhaust end products and their properties for a given fuel additive provided some problems. Information regarding the exact form of the vapor products and the forms into which the gases condense is incomplete. For both sulfur-type and phosphorus-type additives, information is available about the weight percent of acid (sulfuric or phosphoric) present in the condensate formed in a binary water vapor and SO_3 (or P_2O_5) system, but pressures of one atmosphere are presumed. Further, data on surface tension and density are not at hand for all substances assumed to appear in the condensate, although they are usually available for closely allied compounds; e. g., surface tension is presented for liquid P_2O_5 but not for phosphoric acid by weight percent P_2O_5 .

Working assumptions were then:

- (1) P_2O_5 and SO_3 are the only oxidation states at the high temperatures involved,
- (2) all acids presumed in the condensate at one atmosphere actually appear in the contrail,
- (3) condensate data available applies at high altitudes, and
- (4) physical properties of closely related substance suffice.

There would be considerable difficulty in assessing the extent to which these assumptions affect the results were it not for the fact that independent studies of phosphorus-type and sulfur-type additives produced almost identical results with respect to visibility. It would appear then that knowledge of the exact form of the exhaust end products is not crucial, provided that nucleation definitely occurs and reasonable estimates of condensate properties are available.

The third set of assumptions involves the nucleation process itself. Critical radius size and rate of formation of critical size nuclei are described by formulas derived from thermodynamic considerations, and have been experimentally verified. These formulas, however, require a knowledge of partial pressure of water vapor for each instance of nucleation (at discrete intervals in the simulation), and this, in turn, requires knowledge of how much water has already condensed. Due to the complexity of the problem, described more fully in [1] on pages 17-19, the assumption was made that all nucleation has terminated before condensation begins. This is not unreasonable, considering the small mass of acid relative to condensing water vapor.

The fourth and last group of assumptions, summarized on pages 20-21 of [1], is concerned with formation of the contrail itself from the nucleated particles and excess water vapor. Initial particles grow as they collide and stick, grow as water condenses on them, diminish as they collide and split, and perhaps diminish due to evaporation. Eventually, the collection of particles becomes statistically stable; i. e., during a relatively lengthy time interval, the number of particles of any given size is roughly constant.

For the sake of simplicity, prior research has assumed that the size frequency distribution of particles in an observed contrail was of the type $f(\text{radius}) = \text{constant} \cdot (\text{radius})^{-n}$, but this obviously need not be the case. In fact, just as there is a critical radius size in the nucleation process, there may be one in the condensation - collision process. However, even if the mode of such a distribution were 20 times the minimum radius, this would produce a change over only a few percent of the total size range (extending to at least 450 times the minimum). It is reasonable then to assume that the model $f(\text{radius}) = \text{constant} \cdot (\text{radius})^{-n}$ is qualitatively correct.

While neither the exponent "n" of the distribution nor the maximum size is known with certainty, ranges for each can be established which almost surely include the "correct" values. Tables of visibility (discussed further in Section IV) using these two parameters as variables show a very wide range, so that here is a set of parameters in which small variations can produce major changes in visibility.

Visibility is also substantially affected by relative humidity and ambient temperature. This result of the simulation (supported, of course, by empirical evidence) implies a need to monitor these parameters if dynamic control is to be achieved. However it may be possible to avoid this non-trivial problem if acceptable design values can be established which suppress visibility within normal ranges of these parameters.

III. Experimental Analysis

Much experimental work has been done in attempting to isolate the factors affecting contrail visibility. Among others a series of in-flight trials using a sulphur additive has been instituted at Holloman Air Force Base in New Mexico, data from which (Appendix A, table 1) has been made available. Regardless of the method of SO₃ addition all analyses of the data are based on the use of a hypothetical additive having the formula C₅H₁₁SO₃. The analyses conducted had as their purpose the determination of the two particle size distribution parameters of the theoretical description according to the following process.

Using the thermodynamic variables recorded for each trial, the computer simulation results in a numerical relation between assumed exponent "n" of the size frequency distribution and the corresponding predicted visible fraction of the contrail. Let us assume now that the "actual" value of n in one contrail is the same as that in any other, that it is in fact "universal" over all contrails. While its value is unknown, an estimate can be made and the implications examined to possibly verify or reject the estimate.

Some value of n is now selected, and for each flight trial, a visible fraction of contrail is computed. The difference between this estimate and the observed visible fraction is presumably due to the effects of the non - thermodynamic set of variables; a regression is accordingly performed on each of the two flight - trial sets of differences by the remaining variables. If the value selected for n was truly the "actual" value, then the two regressions would be substantially similar, since they are each fitting the same variables to their true effects. If the selected value is not close to the "actual" value, the regressions will be estimating biased effects and are extremely unlikely to agree substantially. The "actual" value of n can then be best estimated by executing the above process for each n in a range covering the true value, and looking for rough equivalence of two (linear) equations, the accuracy of the result probably depending on the degree of equivalence.

Unfortunately, the above rather elegant scheme requires elimination of several possible shortcomings in the given data, before confidence can be placed in results;

- (a) lack of accurate and repeatable recording of observed visible fraction of contrail,
- (b) distribution exponent probably varies with total mass,
- (c) lack of information on accuracy of readings (of dials, gauges, etc.),
- (d) data unavailable for some variables,
- (e) every potential variable is not included in the data.

It should be clear at this point that analysis of the Holloman data was undertaken with a particular goal in mind, even though the conclusions of the analysis are open to question.

The data in Appendix A consist of 30 observations of in-flight trials. Values of nine variables are recorded with each value of the response (estimated percent of suppression). The entire set was subjected to extensive regression - correlation analysis with virtually no significant prediction capability resulting.

Among the difficulties was the absence of recorded values for additive flow rate (variable nine) when suppression was not 100%. This means that for the recorded cases of variable nine the response had no variation.

Variables 1, 3, 4 and 8 are statistically equivalent, with correlations of over 99% between any pair. Variables 6 and 7 correlate over 99% also. The immediate result is a reduction in the number of available variables to four: variables 1, 2, 5 and 6. These four, it must be noted, do not exhibit correlations near 99% between any two, but are hardly independent nonetheless. Correlations in the group run from 68% (2 and 6) to 90% (5 and 6), and 1 and 5 together correlate 6 by 94%.

The best regression estimate would, therefore, involve variables 1, 2 and 5, and these three variables and their squares were fitted to the thirty observations of percent of contrail suppression. The squares of the three variables contributed nothing to the fit after the variables themselves were used, and were thereafter ignored.

Sixty percent of the variation in the response can be accounted for by variables 1, 2 and 5, using the equation:
(Percent Suppression) = 4048 (Mach Number) + 1.126 (Exhaust Temp)
- 2.916 (Altitude) - 3406.

The variance remaining in the "best fit", after using the measured variables and parameters is sufficient to preclude use of the method outlined above for determining the exponent in the contrail particle size distribution. In fact, further analysis of data of this type is unlikely to prove of value until problems (a), (c), (d) and (e) listed above are adequately resolved, and methods to account for (b) are defined. It thus became necessary to seek some other means of estimating the distribution exponent.

IV. Further Simulation Studies

At this point, when the experimental analysis clearly could go no further, two additional facts were brought out. [5]

- (1) At altitude 60,500 feet, using an additive flow rate of 35 pounds per hour, the contrail formed could be described as having impending visibility; that is, the contrail was at the borderline of the visible and invisible regions.
- (2) A new set of in-flight data had been made available, (Appendix A Table 2).

The additive in this case was 70% chlorosulfonic acid (CSA), HClSO_3 , and 30% sulfur trioxide, SO_3 , by weight, and was injected through a small nozzle at the tailpipe.

Whereas previous theoretical results included a mathematical relationship between extent of visibility, range of particle size, and distribution exponent, the additional fact (1) removes "extent of visibility" from the relationship, and provides a simple method for determining the remaining two parameters.

The given flow (35 pounds per hour) and mix of additive produces 27.334 pounds of SO_3 per hour, which is equivalent (in SO_3 production) to a flow rate of 51.622 pounds per hour of sulphur additive ($\text{C}_5\text{H}_{11}\text{SO}_3$).

A computer program was then written to implement the formula at the top of page 23 in [1]. For that formula, the values for m and R_c were found by running the simulation program for altitude 60,500 feet and 23.36% additive, m_{vis} is 0.018 gms/m^3 , and R_1 is one-quarter micron. The value "23.36" for percent additive is a result of knowing the total fuel flow (221.0 pounds per hour) and the above sulphur additive flow rate (51.622 pounds per hour). Given any value of the distribution exponent, n , a corresponding value of the ratio, M , of largest to smallest particle size can be computed, and pairs of such values satisfying the "impending visibility" criterion are graphed in Appendix B, Figure 1.

The value of fuel flow rate at 60,500 feet is not necessarily that at 60,000 feet (221.0 pounds per hour) but the difference was assumed to be insignificant. Parameters in the simulation may also be affected by using the sulphur additive instead of CSA- SO_3 (such as exhaust temperature or total H_2O mass), but these effects are expected to be quite minor.

Figure 1 of Appendix B, notwithstanding a degree of freedom in the M - n relationship, does indicate a narrow range in which the "true" value of n lies.

The ratio of largest to smallest particle size must be at least as large as the ratio of minimum visible size to typical initial size,

$(0.25 \text{ micron})/(6\text{\AA}) = 417$, and could certainly go as high as twice that. If the "probable" range is taken from 450 to 800 (at 417 itself, visibility is precluded), the corresponding range for n is from 4.00 to 4.52. Note that the vicinity of $n = 4.69$ is a singularity, where M takes on all large values, qualitatively indicating that values of n near 4.69 should be eliminated.

Because the simulation run of the impending visibility case took into account an altitude of 60,500 feet, and all runs for [1] were based on 60,000 feet, the decision was made to study the effect of altitude on the contrail. Sixty simulations were run, for altitudes 45,000, 55,000, and 65,000 feet, for ambient temperatures 217°K and 207°K , and for 10, 15, 20, 25, and 30 percent sulphur additive.

Various results are presented in Table 1 and Figures 2 and 3 of Appendix B, but before conclusions are drawn, it should be noted that engine and ambient conditions (except temperature and pressure) were not changed from their values at 60,000 feet. Resulting inaccuracies should be more pronounced at 45,000 feet than at the other values.

The most obvious conclusion, on examination of the computer output, is that the fraction of SO_3 which does not decompose to SO_2 before nucleation is almost completely a function of altitude. This conclusion, it may be noted, is statistical only, appearing because pressure (the true controlling variable) is an approximately linear function of altitude within the ranges encountered. Figure 2 in Appendix B shows the mean value (over altitude) of the non-decomposing fractions plotted as a function of altitude; the fractions themselves are within 0.1% of the mean in almost all cases.

The next obvious conclusion is that, for a fixed set of exhaust wake temperature distribution parameters, the value of R_c (the critical radius) is almost invariant to altitude and relative humidity changes (see Tables 1 and 2 in Appendix B). Even when one of those parameters, ambient temperature, was varied for part of the set of 60 runs, no significant changes appeared.

A third conclusion is evident after examination of Figure 3 in Appendix B. The graphs show (for a constant value of n) that, as the altitude increases, the effect on visibility of increasing the percentage of additive in the fuel tends to diminish so that at relatively high altitudes, near 60,000 feet, percent additive is not as effective a factor in controlling contrail formation as at lower altitudes. Of course, this result depends somewhat on the assumption of a fixed acid fraction in each nucleated acid-water droplet, but the inaccuracy is trivial unless the additive fraction decreases to below 10%. This conclusion is borne out by both experimental and theoretical studies on contrail formation which show that a contrail is more unstable and more difficult to form at these high altitudes.

The second set of experimental data (Appendix A, Table 2) is possibly but not necessarily characterized by impending visibility. Information is not recorded as to how "visible" each contrail is, and as a first guess, it was assumed that "impending" would describe the visibility of the contrail formed by an agent flow rate five pounds per hour less than that recorded [5].

On Figure 3 in Appendix B the point Q represents observed data (see statement (1), beginning of this section). The data points of Appendix A, Table 2 are each associated with a percent additive and an altitude which, if the above assumption were valid, should produce a cluster of points around Q on the line "large-particle density=0.018 gm/m³", when plotted on Figure 3. An alternative way of exploring the problem is the plotting of Q and data from Table 2 on a graph of percent additive versus altitude; the shape of the curve in the latter case is not known a priori, but should approximate a simple curve, and should pass through Q.

The latter plot is shown in Appendix B, Figure 4; the curve is neither simple nor coincident with Q.

Figure 4 indicates clearly that the assumption of impending visibility for the data in Table 2 is invalid. The five pounds per hour decrement in flow rate serves only to shift the curve in Figure 4 to the left, and in no way affects the shape.

The result is that the data set in Table 2 can not be used as it stands to expand our knowledge of the two parameters, distribution exponent and ratio of largest to smallest size.

The effects of relative humidity on visible particle concentration are shown in Appendix B, Figure 5. Relatively low values of the size distribution exponent are seen to increase the sensitivity of the model to changes in relative humidity. Higher values of relative humidity produce the expected increase in visibility.

V. Conclusions

Each of sections II, III and IV includes its own results and conclusions, and those will not be repeated here. Instead, a summary of the kind of results found in those sections is presented, and a final experimental-theoretical technique is suggested.

The basic conclusion is that those parameters which determine contrail visibility are the parameters "distribution exponent" and "size range ratio" of the contrail particle size frequency along with relative humidity and ambient temperature. A corollary conclusion, unfortunately, is that the variables included in the experimental analyses and in the thermodynamic description can only indicate a probable range for the size distribution parameters. A description of the model used in this study is contained in Appendix C. This description is of sufficient level of detail to indicate the method of generating data without being encumbered with non-essential details. A final technique is now suggested whose results may determine the two significant parameters of the particle size distribution.

Figure 1 of Appendix B indicates the locus of pairs of values of the two parameters subject to condition (1) of section IV. If similar sets of data relating additive flow rates to altitude (at impending visibility) are recorded, graphs similar to Figure 1 can be constructed for each such set. As a result, if

- (a) the two particle size distribution parameters are truly constant,
- (b) information is accurately recorded, and
- (c) the assumed monotonic particle size distribution closely fits the physical distribution,

then all of the graphs should intersect at the true values of the two parameters. If they do not intersect at exactly one point, the spread should indicate how the parameters vary with respect to those variables with recorded values.

In either event, further analysis is reasonable and justified, and study in the indicated direction will probably prove to be the only method leading to predictions of contrail visibility as quantitative functions of system variables.

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- [1] Jungreis, T. , and Cole, P. , "Digital Computer Simulation of Jet Contrail Nucleation and Visibility Characteristics", Final Report of Air Force Contract AF19(628)-5193 (August 1967).
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- [3] Forstall, W. , and Shapiro, A. , "Momentum and Mass Transfer in Coaxial Gas Jets", Journal of Applied Mechanics, December 1950, pp. 399 - 407.
- [4] Handbook of Chemistry and Physics, 46th Edition, p. F114.
- [5] Personal communications with S. J. Birstein of Air Force Cambridge Research Laboratories, Bedford, Massachusetts.

APPENDIX A (Table 1)

If an entry is blank, use the value above it

Run	Flight Number	Altitude	Mach	Air Speed	Tailpipe	Exhaust Gas	Gauge	Outside Air	Saturation	Additive	Percent
		(x1000) Var. 1	No. Var. 2	(Knots) Var. 3	Pres(InHg) Var. 4	Temp(-°C) Var. 5	Temp(-°C) Var. 6	Temp(-°C) Var. 7	Temp(-°C) Var. 8	Flow Rate Var. 9	
5	1B	50.0	.715	169	11.3	634	60.7	80.0	60.2	45	70
	1B		.707	167		632	61.2			45	70
	2B	58.4	.716	139	7.7	662	45.6	66.0	63.7	40	100
	2B				7.5	664	45.5			40	100
	3B	62.5	.725	128	6.4	691	41.4	62.5	65.2	40	100
	3B		.720	127		688	41.9			40	100
	4B	61.5	.717	129	6.6	685	43.0	64.0	65.0	38	100
	4B		.721	130	6.7	684	44.9	66.0		38	100
	5B	60.5	.721	133	7.0	683	44.9	66.0	64.9	38	100
	5B		.726	134		679	42.1	64.0		38	100
	6B	60.5	.721	133	6.9	670	44.4	66.0	64.9	44	100
6B		.726	134	7.0		45.1			44	100	
7	1A	58.4	.716	139	7.8	650	52.4	72.5	63.9	N/A	0
	1A					648	53.4	73.5		N/A	0
	2A	60.5	.716	132	6.7	648	50.3	71.0	64.6	N/A	0
	2A						49.8	70.0		N/A	0
8	1A	58.4	.720	140	7.8	644	54.5	75.0	63.9	44	100
	1A					643	53.4	74.0		44	100
	2A	60.5	.721	133	6.8	648	47.0	68.0	64.6	43	100
	2A						46.7	67.5		43	100
	3A	61.5	.721	130	6.5	650	48.0	69.5	65.4	37	100
	3A						46.1	67.0		37	100
	4A	56.3	.721	147	8.5	632	58.2	78.0	62.9	N/A	0
	4A						57.1	77.0		N/A	0
	5A	54.2	.706	151	9.1	622	54.5	74.0	62.0	N/A	0
	5A									N/A	0
	6A	52.2	.708	159	10.0	620	59.2	78.5	61.1	N/A	0
	6A						58.9	78.0		N/A	0
	7A	50.0	.711	168	11.0	618	58.9	78.0	60.2	N/A	0
	7A						58.4	77.5		N/A	0

APPENDIX A (Table 2)

(100 Percent Contrail Suppression in All Cases)

Altitude (x1000 ft)	Engine Rpm (Percent)	Outside Air Temp. (-°C)	Pressure (millibars)	Additive Flow (pounds/hour)
63.3	100.0	67.5	60.3	30
61.5	100.0	66.0	66.7	30
61.5	100.0	67.0	66.7	30
58.0	100.1	75.0	77.4	44
60.5	100.0	68.0	70.0	43
61.5	99.7	69.5	66.7	37
60.5	100.0	66.5	70.0	45
60.5	100.0	66.5	70.0	40
62.6	99.0	68.0	63.3	35
62.6	99.0	68.5	63.3	40
64.6	99.0	69.0	57.5	35
64.6	99.0	69.0	57.5	42
65.5	99.0	68.0	55.1	39
65.5	99.0	68.0	55.1	41

APPENDIX B (Table 1)

Ambient Temperature 217°K

Altitude (Feet) Additive Fraction	45,000		55,000		65,000	
	Rc	%	Rc	%	Rc	%
10%	6.42	21.2	6.29	17.5	6.16	14.3
15%	6.34	21.3	6.20	17.6	6.03	14.4
20%	6.19	21.4	6.08	17.6	5.99	14.4
25%	6.17	21.5	6.05	17.7	5.95	14.5
30%	6.14	21.6	6.02	17.8	5.91	14.5

Ambient Temperature 207°K

Altitude (Feet) Additive Fraction	45,000		55,000		65,000	
	Rc	%	Rc	%	Rc	%
10%	6.31	21.2	6.27	17.5	6.14	14.3
15%	6.27	21.3	6.10	17.6	6.03	14.4
20%	6.22	21.4	6.07	17.6	5.95	14.4
25%	6.18	21.5	6.04	17.7	5.91	14.5
30%	6.14	21.6	6.00	17.8	5.88	14.5

The left number of each pair is critical radius, in angstroms (10^{-8} cm.)
 The right number of each pair is the percent of SO_3 not decomposing.
 Rc has a mean value of 6.12, and standard deviation of 0.14 (2% of mean).

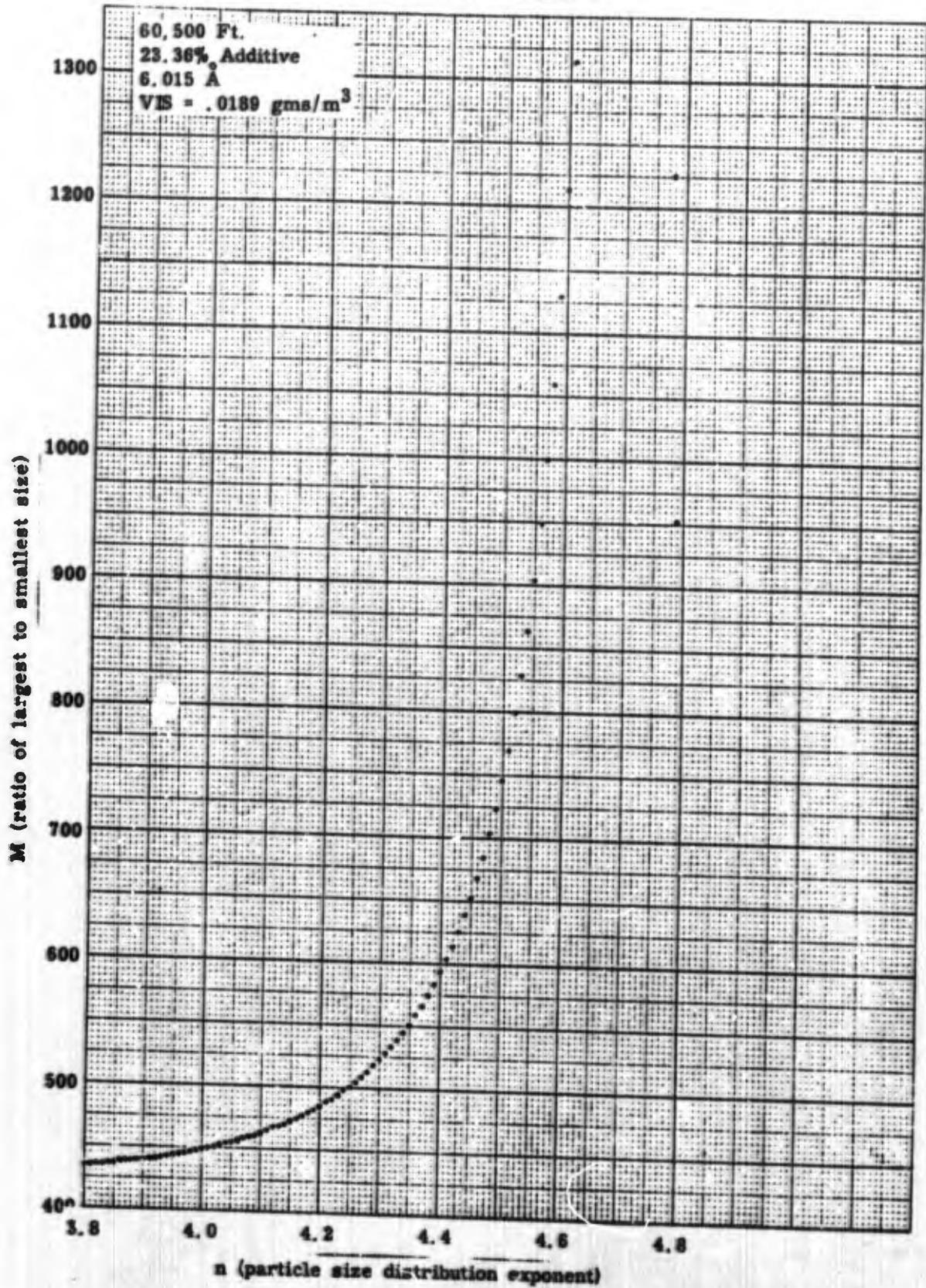
APPENDIX B (Table 2)

Relative Humidity (%)	Ambient Temperature (°K)		
	198 Rc	212 Rc	217 Rc
0	.4776x10 ⁻⁷	.4762x10 ⁻⁷	.4763x10 ⁻⁷
10	.4836x10 ⁻⁷	.4836x10 ⁻⁷	.4896x10 ⁻⁷
20	.4892x10 ⁻⁷	.4892x10 ⁻⁷	.4989x10 ⁻⁷
30	.4943x10 ⁻⁷	.4943x10 ⁻⁷	.5053x10 ⁻⁷
40	.4981x10 ⁻⁷	.4981x10 ⁻⁷	.5106x10 ⁻⁷
50	.5016x10 ⁻⁷	.5016x10 ⁻⁷	.5135x10 ⁻⁷
60	.5048x10 ⁻⁷	.5048x10 ⁻⁷	.5166x10 ⁻⁷
70	.5075x10 ⁻⁷	.5075x10 ⁻⁷	.5204x10 ⁻⁷
80	.5087x10 ⁻⁷	.5087x10 ⁻⁷	.5237x10 ⁻⁷
90	.5108x10 ⁻⁷	.5108x10 ⁻⁷	.5257x10 ⁻⁷
100	.5124x10 ⁻⁷	.5124x10 ⁻⁷	.5287x10 ⁻⁷

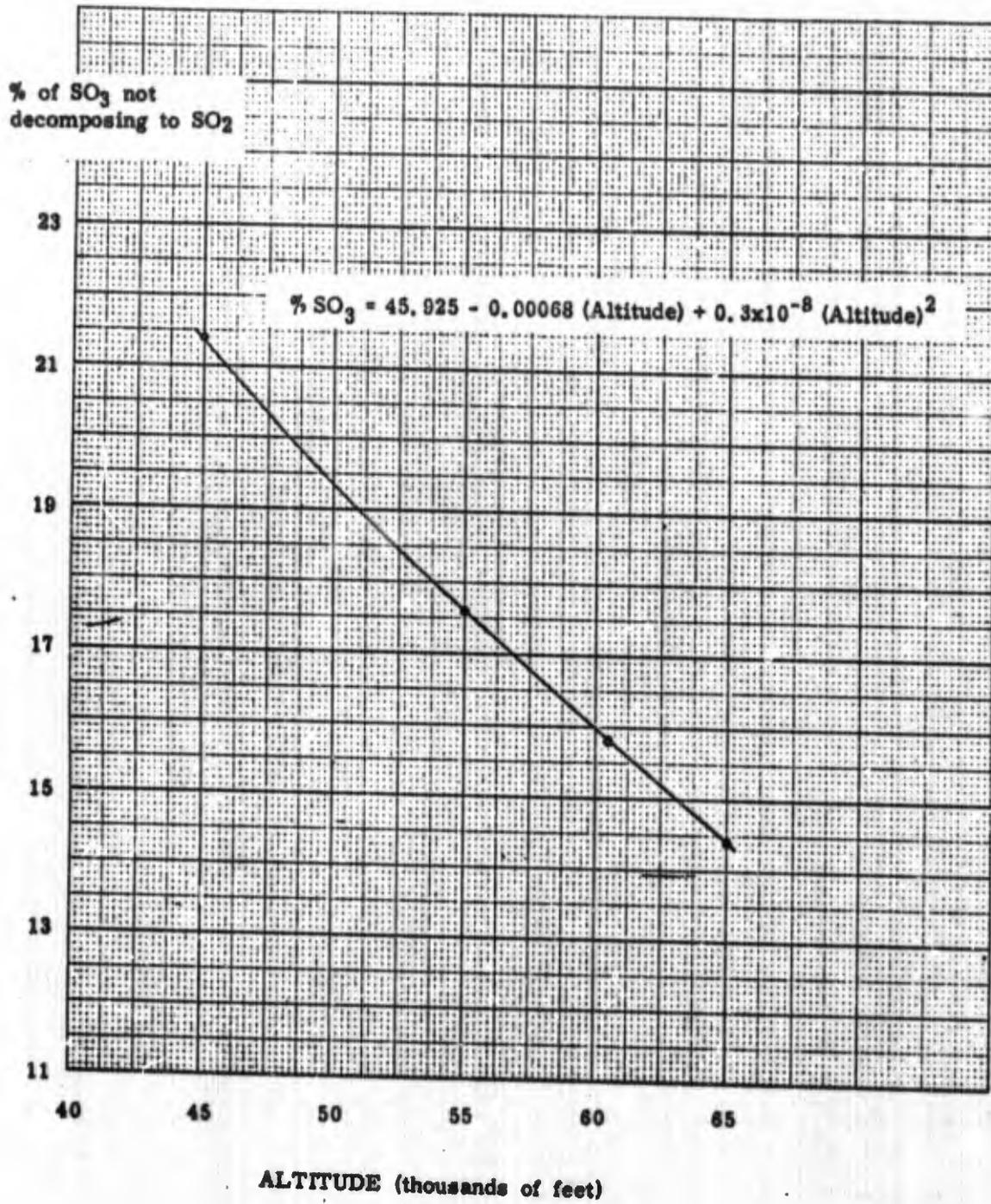
Table of critical radius Rc (cm) vs. relative humidity

At various ambient temperatures

APPENDIX B FIGURE 1

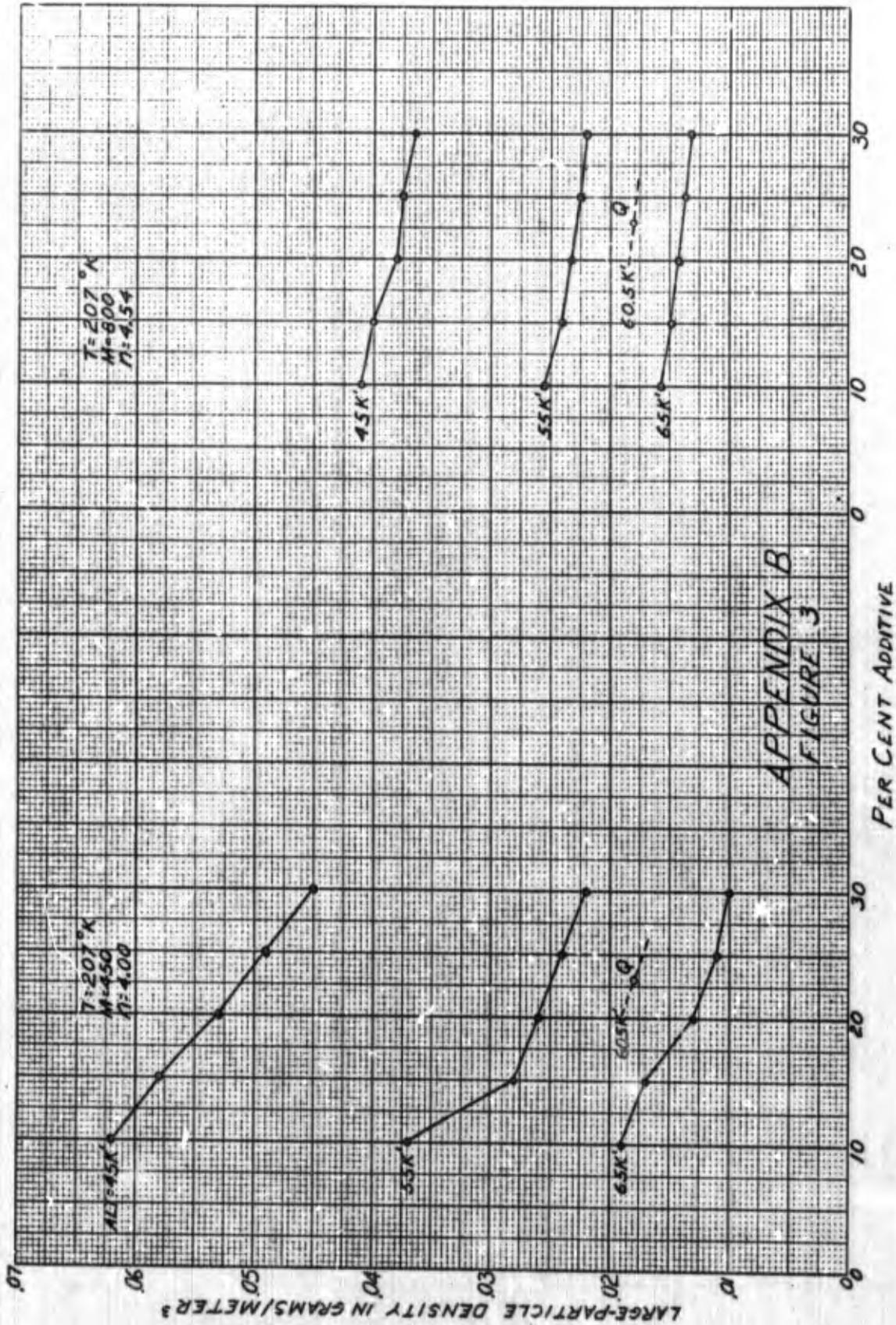


APPENDIX B FIGURE 2

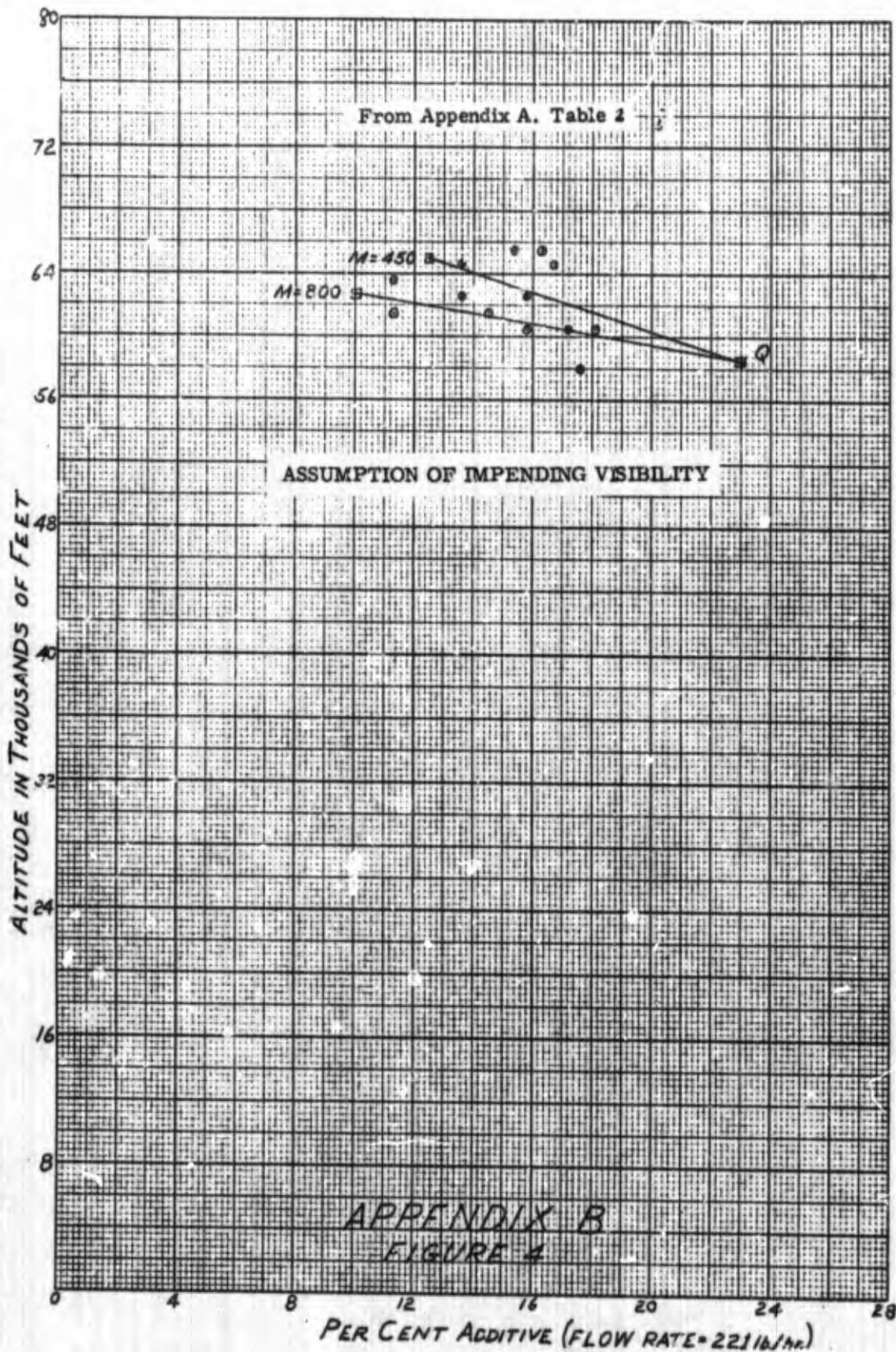


EURENE DIETZGEN CO.
MADE IN U. S. A.

NO. 341-20 DIETZGEN GRAPH PAPER
20 X 20 PER INCH



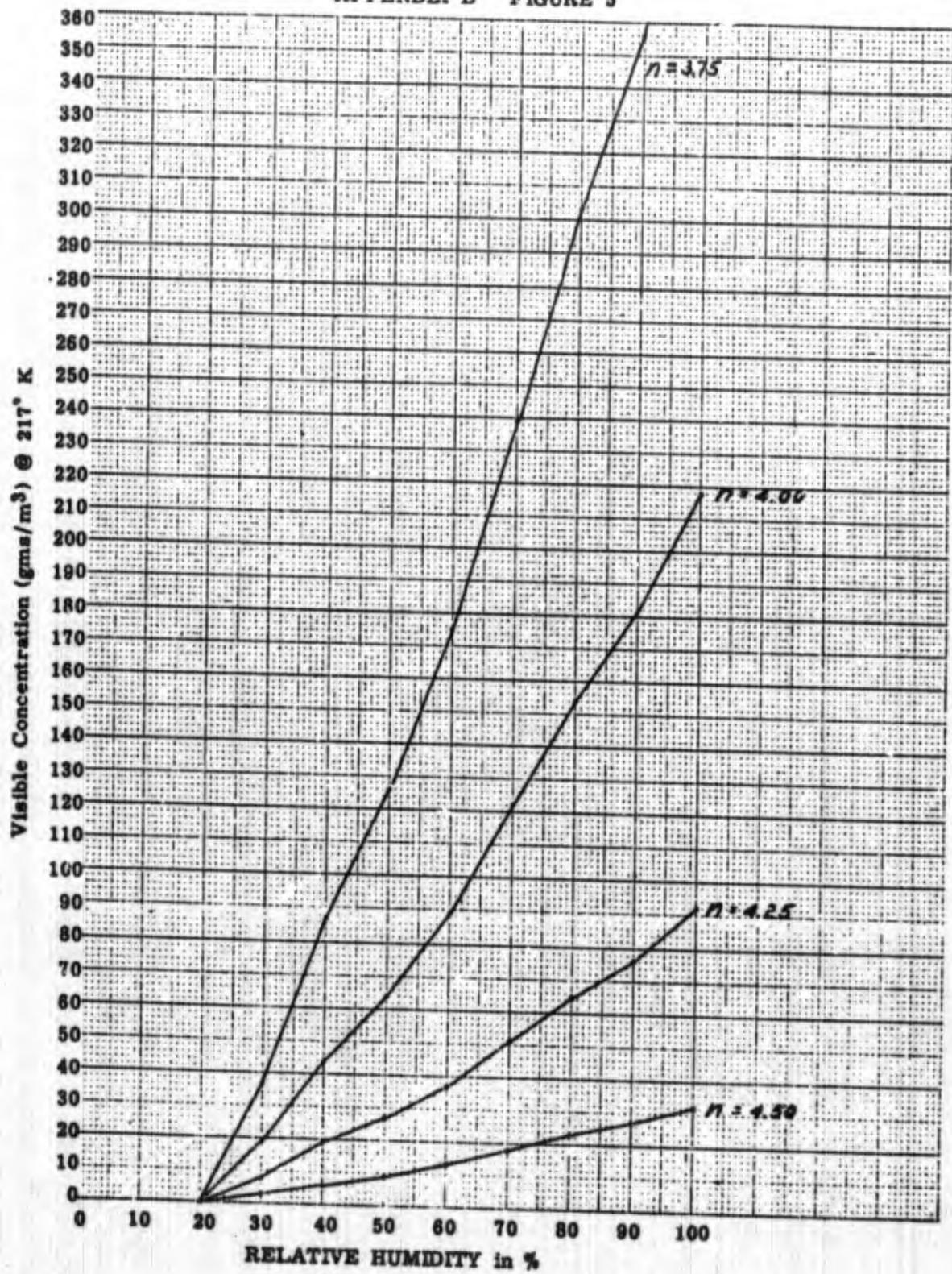
APPENDIX B
FIGURE 3



EUREME DIETZGEN CO.
MADE IN U. S. A.

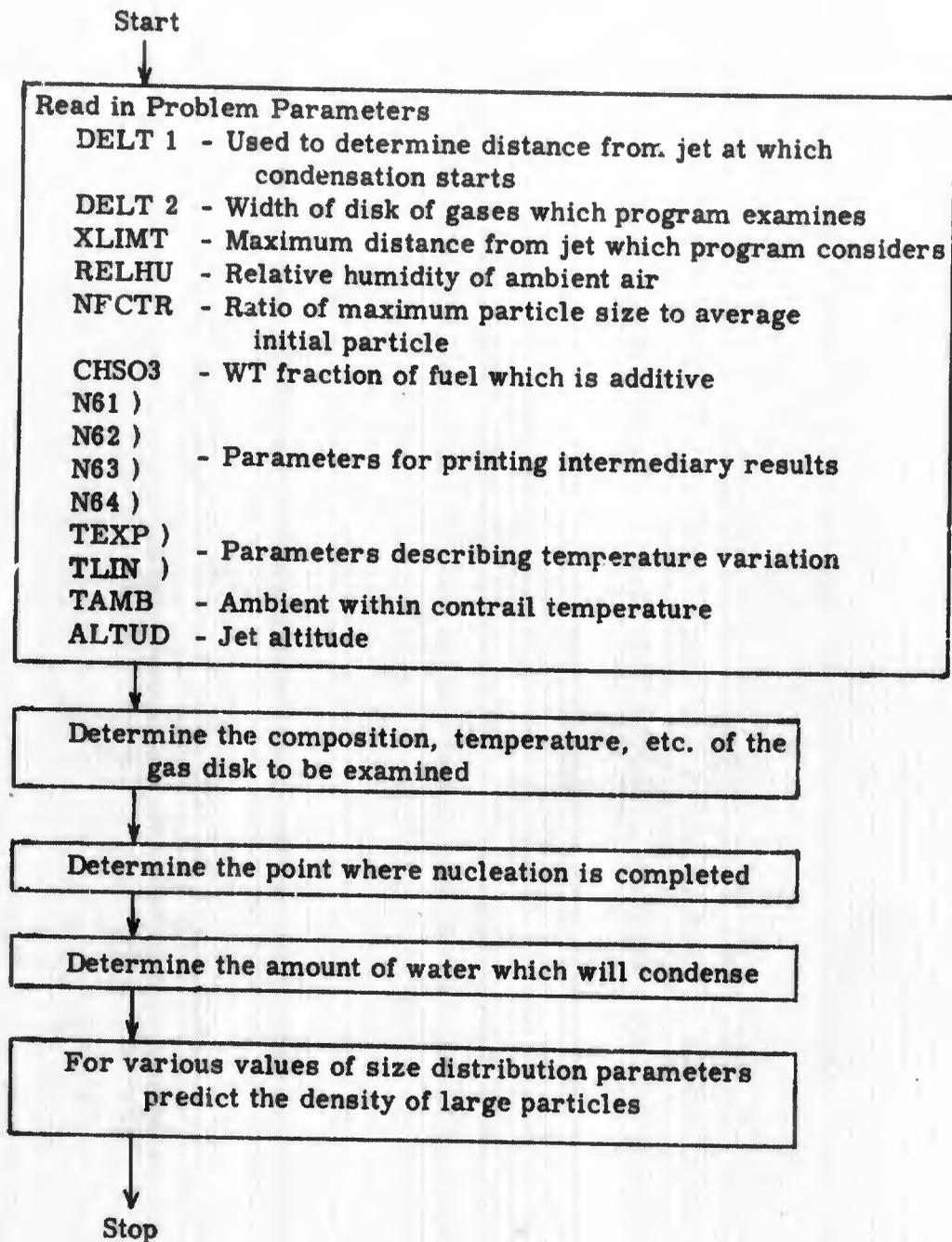
NO. 341-50 DIETZGEN GRAPH PAPER
30 X 30 PER INCH

APPENDIX B FIGURE 5



APPENDIX C FIGURE 1

Simplified Block Diagram of Computer Model



TRAJ P COLE TRAJECTORY
 - - - - - -
 TRAJ - EFN SOURCE STATEMENT - IFN(S) -

MW5 = 99.0775
 MW6 = 151.2057
 MW7 = 1339.685

C
 C ... IF THE DISC IN ITS CURRENT POSITION (DISTANCE FROM JET = X) DOES
 C ... NOT PRODUCE DROPLETS, X IS AUTOMATICALLY INCREMENTED BY DELT1
 C ... DELT2 IS THE WIDTH OF THE DISC OF EXHAUST EXAMINED BY THE PROGRAM
 C ... THE PROGRAM IS ALWAYS TERMINATED IF THE VALUE OF X REACHES XLIMIT
 C

READ (5,1232) DELT1, DELT2, XLIMIT
 1232 FORMAT (F4.1, 1X, F4.1, 1X, F6.1)
 1 CONTINUE

C *****
 C *****
 C *****

READ (5,13) RELHU
 15 FORMAT (F5.1)
 WRITE (6,25) RELHU
 25 FORMAT (20A1 THE RELATIVE HUMIDITY IS, F6.1, 10H PER-CENT.)
 RELHU = RELHU/100.

C *****
 C *****
 C *****

C ... FACTR IS THE FACTOR BY WHICH THE LARGEST PARTICLE'S RADIUS IS
 C ... ASSUMED GREATER THAN THE AVERAGE INITIAL RADIUS

READ (5,1232!) NFCTR
 1232! FORMAT (14)
 FACTR = NFCTR

C
 C ... CHSD3 IS THE WEIGHT FRACTION OF FUEL WHICH IS C5H11SO3
 C
 C ... COLUMNS 51 AND 52 OF THE CHSD3-VALUE CARD ARE EQUAL (PRESUMABLY
 C ... TO 'BLANK') IF PRINTING OF DATA POINT BY POINT IS TO BE SUPPRESSED
 C ... PRINTING WILL OCCUR IF COLUMNS 61-2 DIFFER (ONE MAY BE LEFT BLANK)

C
 C COLUMNS 53 AND 54 ARE EQUAL IF PRINTING OF COEFFICIENTS AND ROOTS
 C OF THE EQUILIBRIUM-RELATED CURIC EQUATION IS TO BE SUPPRESSED
 READ (5,12325) CHSD3, N61, N62, N63, N64

12325 FORMAT(F7.4, 53X, 5A1)

C
 C ... TEXP IS THE EXPONENT OF THE FRACTION DEFINING TEMPERATURE WHICH
 C ... MORE OR LESS DESCRIBES THE MIXING AND VOLUME CHANGE (TURBULENCE)
 C TLINR IS THE DISTANCE, IN FEET, OVER WHICH THE MEAN DISC
 C ABS TEMPERATURE IS ASSUMED TO VARY LINEARLY WITH DISTANCE
 C FROM PIPE AT (L*DIPE/12) TO (T-CENTERLINE + TAMB)/2 AT TLINR
 C BEYOND TLINR THE MEAN IS TAKEN AS (T-CENTERLINE+TAMB)/2

READ (5,1233) TEXP, TLINR
 1233 FORMAT(F5.3, 2X, F5.1)

C ... TAMB IS THE ATMOSPHERIC (AMBIENT) TEMPERATURE AT 60,000 FT
 C ... ALTUD IS JET ALTITUDE IN FEET
 C IF NO ALTITUDE IS SPECIFIED, 60,000 FT WILL BE ASSUMED.

READ (5,1234) TAMB, ALTUD
 1234! FORMAT (F7.2, 2X, F5.0)
 IF (ALTUD) 6000, 6000, 6000

6000 ALTUD = 60000.0
 6010 CONTINUE

TRAJ P COLE TRAJECTORY
 TRAJ - EFN SOURCE STATEMENT - (FN(S) -

C
 C ... AIRIN IS THE AIR INTAKE IN LBS/SEC (NOT INCLUDING 0.06 FOR FUEL)
 AIRIN = 3.05
 C ... TPIPE IN DEGREES KELVIN
 TPIPE = 1300
 C ... PPIPE IN ATMOSPHERES (EXIT PRESSURE)
 PPIPE = 0.25
 C
 C ... DPIPE IS THE TAILPIPE DIAMETER IN INCHES
 DPIPE = 11.0
 C ... VPIPE IS IN FEET PER SECOND
 VPIPE = $R * TPIPE * AIRIN * LB2GM * 576.0 /$
 $1 * (4 * L * (PPPIPE * 1.01325E 6) * PI * DPIPE ** 2 * FT2CC)$
 C ... XTIME IS THE TIME REQUIRED FOR THE FORMATION OF THE DISC
 XTIME = $DEL T2 / VPIPE$
 C ... THRST IS THE JET THRUST IN POUNDS .
 THRST = 1700
 C ... FUEL IS THE WEIGHT OF FUEL BURNED PER HOUR PER POUND OF THRUST
 FUEL = 1.3
 C ... TEMP IS THE NUMBER OF GRAMS OF FUEL BURNED DURING DISC FORMATION
 TEMP = $FUEL * THRST * LB2GM * XTIME / 3600.0$
 C
 C ... PAMB IS AMBIENT PRESSURE IN ATMOSPHERES, ALTITUDE IN FEET
 PAMB = $10.0 * (-0.64528 - 0.68133 * (-1.10191 + ALTUD / 32000.0)) * C.986923$
 C
 C ... MOLS1 IS MOLES OF AIR IN THE DISC
 C ... MOLS2 IS MOLES OF H2O IN THE DISC
 C ... MOLS3 IS MOLES OF SO3 IN THE DISC
 C ... MOLS4 IS MOLES OF SO2 IN THE DISC
 C ... MOLS5 IS MOLES OF O-2 IN THE DISC
 C ... MOLS6 IS TOTAL INERT MOLES IN THE DISC (CO2 PLUS INERT AIR)
 MOLS1 = $AIRIN * LB2GM * XTIME / MW1$
 MOLS2 = $TEMP * (CHSO3 * 5.5 / MW6 + (1.0 - CHSO3) * 98.5 / MW7)$
 C *****
 C *****
 C *****
 C ... VPSAT IS VAPOR PRESSURE (IN MILLIBARS) AT SATURATION.
 (A MILLIBAR IS 1,000 DYNES PER SQUARE CM.)
 VPSAT = $10. ** (-7.90298 * (373.16 / TAMB - 1.0) + 5.02808 * ALG10(373.16 / TAMB - 1.0) - 1.3816E - 7) * (10. ** (11.344 * (1.0 - TAMB / 373.16)) - 1.0) + (8.1328E - 3) * 2(10. ** (-3.49149 * (373.16 / TAMB - 1.0) - 1.0) + ALG10(1013.246))$
 C ... ABSHU IS ABSOLUTE HUMIDITY AT SATURATION IN GRAMS PER CC
 ABSHU = $VPSAT * 142 * 1000.0 / (R * TAMB)$
 C ... SATHU IS NO. OF GRAMS OF WATER PER GRAM OF AIR AT SATURATION.
 SATHU = $ABSHU * R * TAMB * CGSAT / (PAMB * MW1)$
 SHAMB = SATHU
 MOLS1 = $MOLS1 - LB2GM * AIRIN * RELHU * SATHU * XTIME / MW2$
 MOLS2 = $MOLS2 + LB2GM * AIRIN * RELHU * SATHU * XTIME / MW2$
 C *****
 C *****
 C *****
 C ... MOLS3 = $TEMP * CHSO3 / MW6$
 MOLS4 = 0.
 OMOLS5 = $0.13957 * MOLS1 - 149.25 * TEMP * (1.0 - CHSO3) / MW7$
 1 - $7.75 * TEMP * CHSO3 / MW6$
 OMOLS6 = $0.313 * MOLS1 + TEMP * (1.0 - CHSO3) * 100.0 / MW7$

```

TRAJ          P COLE          TRAJECTORY
TRAJ          - EFN  SOURCE STATEMENT - IFN(S) -

1          + TEMP*CHSC3*5.0/MW6
C ... MOLS  IS TOTAL MOLES IN DISC (MOLS1 INDIRECTLY INCL IN MOLS5,6)
C
C          MOLS = MOLS2 + MOLS3 + MOLS4 + MOLS5 + MOLS6
C
C ... FRCT3, FRCT4, AND FRCT5 ARE THE FRACTIONS OF THE TOTAL MOLES WHICH
C          ARE SO3, SO2, AND O-2, RESPECTIVELY.
C          FRCT3 = MOLS3 / MOLS
C          FRCT4 = MOLS4 / MOLS
C          FRCT5 = MOLS5 / MOLS
C          APROX = 0.5 * FRCT3
C
C ... AVLBW IS THE INITIAL NO. OF MOLES OF H2O AVAILABLE FOR NUCLEATION
C          AVLBW = MOLS2
C          AVLBL IS THE NO. OF MOLES OF SO3 AVAILABLE FOR NUCLEATION OR DECAY
C          AVLBL = MOLS3
C          AVLBR IS THE NUMBER OF MOLES AVAILABLE FOR NUCLEATION AFTER DECAY
C          AVLBR = AVLBL
C ... VAMB IS FORWARD VELOCITY OF JET IN KNOTS
C          VAMB = 315.0
C          L = 0.0 + 12.0 * 1.6878099 * VAMB / VPIPE
C ... THIS VALUE OF X IS WHERE THE CALCULATIONS START (X IN FEET)
C          START = L * VPIPE / 12.0
C          X = START
C
C          WRITE (6,12342) TAMB, ALTUD
12342 FORMAT (1HJTA4H = ,F7.2, 10H AT AN ALTITUDE OF, F7.0)
C          WRITE (6,12343) DELT1, DELT2, XLIMT
12343 FORMAT (1HJUCLT1 = ,F4.1,10H DELT2 = ,F4.1,12H X-LIMIT = ,F6.1)
C          WRITE (6,12345) TEXP, TLINR
12345 FORMAT (25H TEMPERATURE EXPONENT IS ,F6.3,/60H DISTANCE OVER WHICH
C          IN THE TEMPERATURE FUNCTION IS LINEAR IN T-AMB IS ,F6.1)
C ... DIAM2 AND DIAM3 ARE THE DIAMETERS OF H2O AND SO3 MOLECULES IN CMS
C ... DIAMETERS ARE BASED ON LIQUID VOLUMES. SPEC GRV OF SO3 IS 1.95
C          DIAM2 = 3.3314E-8
C          DIAM3 = 5.3684E-8
C ... MASS2 AND MASS3 ARE THE MASSES, IN GRAMS, OF H2O AND SO3 MOLECULES
C          MASS2 = 0.299133E-22
C          MASS3 = 1.329400E-22
C ... VOL40 IS INITIAL VOLUME IN CUBIC FEET
C          VOLMO = DELT2 * PI * (DPIPE/24.0)**2
C          WRITE (6,12350)
12350 FORMAT(1J7H
C          IS SO3          VPIPE          TEMP          MOLES H2O          MOLE
C          PAMB )
C          WRITE (6,12352) CHSO3, TEMP, MOLS2, MOLS3, VPIPE, L, PAMB
12352 FORMAT (1H CHSO3 = ,F7.4,6E16.7)
C          WRITE (6,12354)
12354 FORMAT (121H0 EXIT TIME TCTAL MOLES SO3 FRACTION SO2 FR
C          IACTION O-2 FRACTION INITIAL X INITIAL VOLUME EXIT TEMP (K
C          2))
C          WRITE (6,12355) XTIME, MOLS, FRCT3, FRCT4, FRCT5, X, VOLMO, TPIPE
12355 FORMAT (1H ,7E15.8,F11.2)
C          IF ( N61 - 462 ) 6030, 6035, 6030
6030 WRITE (6,12356)
12356 FORMAT (110H)
1          P-ZE4J          ALPHA          TEMPERATURE          VOLUME          Z
          TEMP4          MWBAR)

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TRAJ - P COLE TRAJECTORY
 TRAJ - EFN SOURCE STATEMENT - IFN(S) -

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C ... J      IS THE INDEX OF THE LATEST POINT OF SIGNIFICANT NUCLEATION
6035 J      = J
C ... SAVE(J,1) IS THE DISTANCE X OF PCINT J FROM THE TAILPIPE, IN FEET
C ... SAVE(J,2) IS THE AVERAGE MOLECULAR WEIGHT OF THE NUCLEATED DROPLET
C ... SAVE(J,3) IS THE NUCLEATION RATE AT DISTANCE X
C ... SAVE(J,4) IS THE CRITICAL RADIUS
C ... SAVE(J,5) IS THE NUMBER OF NEW NUCLEI CREATED AT DISTANCE X
C ... SAVE(J,6) IS THE TEMPERATURE AT DISTANCE X
C ... SAVE(J,7) IS MOLES OF H2O AVAILABLE WHEN THE DISC REACHES POINT X
C ... SAVE(J,8) IS MOLES OF P2O5 AVAILABLE WHEN THE DISC REACHES POINT X
C ... SAVE(J,9) IS THE DENSITY OF THE NUCLEATED DROPLET
C ... TOT1 IS THE TOTAL NUMBER OF PARTICLES NUCLEATED FROM XMIN TO XMAX
C ... TOT2 IS THE SUM OF THE RADII OF ALL PARTICLES NUCLEATED
C ... TOT3 IS THE SUM OF THE RADIUS-SQUARED OF ALL NUCLEATED PARTICLES
C ... TOT4 IS THE SUM OF THE RADIUS-CUBED OF ALL PARTICLES NUCLEATED
      TOT1 = J.0
      TOT2 = J.0
      TOT3 = J.0
      TOT4 = J.0
C ... UZPCT = THE PERCENT OF THE ORIGINAL SO3 ALREADY USED IN NUCLEATION
      UZPCT = J.0
C ... DELTX IS 14 FEET AND IS DELT1 BEFORE NUCLEATION, DELT2 SUBSEQUENTLY
      DELTX = DELT1
      GO TO 7011
6999 J      = J + 1
7000 X      = X + DELTX
C ... IF X IS GREATER THAN XLINT FEET, TERMINATE NUCLEATION PROCESS
7001 IF( X - XLINT ) 7010, 7010, 7075
C
C ... BETWEEN START AND TLINR, THE MEAN DISC TEMPERATURE VARIES
C ... LINEARLY FROM TPIPE=T-CENTERLINE AT X = START TO
C ... (1/2)*(T-CENTERLINE+TAMB) AT TLINR
7010 IF( X - TLINR ) 7012, 7013, 7013
7012 COEF1 = -J.0 / (TLINR - START)
      COEF2 = (TLINR - 0.5*START) / (TLINR - START)
      COEF3 = COEF1 * X + COEF2
      TABS = COEF3 * (TEMPX(X) - TAMB) + TAMB
      GO TO 7014
7013 TABS = (TEMPX(X) + TAMB) / 2.0
7014 TCENT = TABS - 273.16
C *****
C *****
C *****
      RDISK=1.5 + .0385*X
      VOLNE=JELT2*PI*(RDISK**2)
      NNAVE = 1.0/(MOLS2/(VOLNE*FT2CC) + .03466798)
      WTAIR=4*NAVE*(PAMB*VOLNE*FT2CC/(R*TABS*CGSAT)-MOLS1)
      MOLS1 = MOLS1 + WTAIR/NNAVE
      MOLS2 = MOLS2 + SHAMB*RELMU*WTAIR/NW2
      MOLS5=MOLS5+.20946*WTAIR/31.9988
      MOLS6=MOLS6+.78384*WTAIR/28.0134+.00033*WTAIR/44.0995+.00934*
      WTAIR/39.948
      VPSAT= 10.**(-7.90298*(373.16/TABS-1.0)+5.02636*ALOG10( 373.16/TAB
      IS)-(-1.3416E-7)*(10.**((11.344*(1.0-TABS/373.16))-1.0))+(-8.1328E-3)*
      2(10.**(-3.49149*(373.16/TABS-1.0))-1.0)+ALOG10(1213.246))
      ABSHU = VPSAT*AWZ*1000.0/(R*TABS)
    
```

TRAJ P COLE TRAJECTORY
 TRAJ - EFN SOURCE STATEMENT - IFN(S) -

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SATHU=ABS(HU**R*TABS*CGSAT/(PAMB*PWAVE)
AVLBW = 4JLS2-SATHU*WTAIR/MW2
IF (AVLBW .LT. 0.) AVLBW = 0.0
C *****
C *****
C *****
C
MOLS = 4JLS2 + MOLS3 + MOLS4 + MOLS5 + MOLS6
C FRCT3, FRCT4, AND FRCT5 ARE THE FRACTIONS OF THE TOTAL MOLES WHICH
C ... ARE S3, S2, AND O-2, RESPECTIVELY
FRCT3 = 4JLS3 / MOLS
FRCT4 = 4JLS4 / MOLS
FRCT5 = 4JLS5 / MOLS
C THE EQUILIBRIUM RELATION LEADS TO A CUBIC EQUATION FOR THE
C INCREMENT OF MOLES OF SO3 DECAYED. GIVEN A FIRST ESTIMATE, THE
C ROOT MAY BE ACCURATELY FOUND USING NEWTON'S METHOD. THE FOUR
C COEFFICIENTS OF THE CUBIC ARE CUBIC, QUAD1, QUAD2, QUAD3.
C THE ESTIMATE IN THE FIRST INTERVAL IS 0.5*FRCT3, AND IN ALL
C SUBSEQUENT INTERVALS IS 0.01*FRCT3. IF THE ROOT IS NOT POSITIVE
C THE ESTIMATE FOR THE NEXT INTERVAL IS SET TO ZERO.
C
C ... THE EQUILIBRIUM CONSTANT IS EQCON(TABS) = EXP(4956.0/TABS - 4.678)
C ... THE CONSTANT RELATES PSO3, PSC2, PO2 (FOR P IN ATMOSPHERES) BY
C ... EQCON(TABS) = PSO3 / ( PSC2 * SORT(PO2) ) (AT EQUILIBRIUM)
EQCON = EXP( 4956.0 / TABS - 4.678 )
CONST = PA48 * EQCON * EQCON
CUBIC = CONST - 1.0
QUAD1 = 2.0 * ( CONST*(FRCT4*FRCT5) + FRCT3 - 1.0 )
QUAD2 = CONST * FRCT4 * 14. FRCT5+FRCT4) - FRCT3 * (FRCT3 - 4.0 )
QUAD3 = 2.0 * CONST * FRCT -2 * FRCT5 - 2.0) * FRCT3**2
C
IF( N63 - N64 ) 861, 862, 861
861 CONTINUE
WRITE (6,12357) APROX, CUBIC, QUAD1, QUAD2,QUAD3
12357 FORMAT (9H4APROX = ,E13.6,20H CUBIC, QUAD1-3 ARE,4E16.8)
862 CONTINUE
C
IF( CUBIC ) 5003, 5001, 5003
C THE COEFFICIENT OF THE CUBIC TERM IS ZERO--DEGENERATE CASE.
5001 ROOT = 0.0
CUBIC = QUAD1
QUAD1 = QUAD2
QUAD2 = QUAD3
C WHEN 'CUBIC' IS ZERO, QUAD1 IS ALWAYS NEGATIVE, AND QUAD2 POSITIVE
IF( N63 - N64 ) 863, 864, 863
863 CONTINUE
WRITE (6,11177)
11177 FORMAT ( 35H DEGENERATE CUBIC EQUATION--THE REDUCED QUADRATIC EQUA
TION ALWAYS HAS A POSITIVE ROOT.)
864 CONTINUE
GO TO 5007
C
5003 TEST =SIGN(1.,CUBIC)+SIGN(1.,QUAD1)+SIGN(1.,QUAD2)+SIGN(1.,QUAD3)
C IF ALL COEFFICIENTS ARE NON-NEGATIVE, NO POSITIVE ROOT IS POSSIBLE
IF( TEST - 4. ) 5005, 5030, 5030
5005 F = QUAD3 + APROX * (QUAD2 + APROX * (QUAD1 + APROX * CUBIC))

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TRAJ      P COLE      TRAJECTORY
TRAJ      - EFN      SOURCE STATEMENT - IFN(S) -

FPRIM = QUAD2 + APROX * ( 2.0*QUAD1 + 3.0*CUBIC*APROX )
ROOT = APRJA - F / FPRIM
IFI N63 - N64 ) 865, 866, 865
865 CONTINUE
WRITE (6,12353) ROOT
12353 FORMAT (9H ROOT = ,E13.6)
866 CONTINUE
IFI ABS( (RJJT-APROX)/ROOT ) - 0.00005 ) 5007, 5006, 5006
5006 APROX = RJJT
GO TO 5005
5007 QUAD1 = CUBIC
QUAD2 = QUAD1 + ROOT * CUBIC
QUAD3 = QUAD2 + ROOT * (QUAD1 + ROOT*CUBIC)
C THE ROOT 'RJJT' IS ALGEBRAICALLY REMOVED FROM THE CUBIC EQUATION
C DISCR IS THE DISCRIMINANT OF THE RESULTING QUADRATIC EXPRESSION
DISCR = QUAD2**2 - 4.0 * QUAD1 * QUAD3
IFI DISCR ) 5008, 5009, 5009
5008 IF( ROOT ) 5030, 5030, 5020
C
5009 ROOT2 = ( -QUAD2 + SORT( DISCR ) ) / (2.0 * QUAD1)
ROOT3 = ( -QUAD2 - SORT( DISCR ) ) / (2.0 * QUAD1)
IFI N63 - N64 ) 867, 868, 867
867 CONTINUE
WRITE (6,11188) ROOT2, ROOT3
11188 FORMAT (9H ROOT2 = ,E14.7,11H ROOT3 = ,E14.7)
868 CONTINUE
C THE NEXT TEN CARDS FERRET OUT THE SMALLEST POSITIVE ROOT
C THE TWO REMAINING ROOTS ARE NOT REQUIRED AND ARE IGNORED
IFI ROOT2 ) 5011, 5011, 5010
5010 IF( ROOT3 ) 5013, 5013, 5014
5011 IF( ROOT3 ) 5020, 5020, 5012
5012 ROOT2 = ROOT3
5013 IF( ROOT ) 5019, 5019, 5018
5014 IF( ROOT ) 5015, 5015, 5016
5015 ROOT = ROOT3
5016 IF( ROOT3 - ROOT2 ) 5017, 5018, 5018
5017 ROOT2 = ROOT3
5018 IF( ROOT2 - ROOT ) 5019, 5020, 5020
5019 ROOT = ROOT2
5020 IF( ROOT - ROOT3 ) 5025, 5025, 5030
C CHANGE IS THE CHANGE IN MOLES OF SO3 DUE TO DECAY AT THIS STEP
5025 CHNGE = ROOT * MOLS
MOLS3 = MOLS3 - CHNGE
MOLS4 = MOLS4 + CHNGE
MOLS5 = MOLS5 + CHNGE / 2.0
MOLS = MOLS + CHNGE / 2.0
AVLBR = AVLBR - CHNGE
GO TO 5035
C
5030 IF( N63 - N64 ) 869, 870, 869
869 CONTINUE
WRITE (6,12359)
12359 FORMAT (38H NO CHANGE IN EQUILIBRIUM AT THIS STEP)
870 CONTINUE
5035 CONTINUE
IFI N63 - N64 ) 871, 872, 871

```

TRAJ P COLE TRAJECTORY
 - EFN SOURCE STATEMENT - IFN(S) -

071 CONTINUE
 WRITE (6,12365) MOLS3, MOLS4, MOLS5, MOLS
 12365 FORMAT (354 AFTER TAKING INTO ACCOUNT PREVIOUS NULLATION (IF ANY)
 1, AND DECAY THIS STEP (IF ANY),/,21H MOLES S03 = ,E14.7,
 215H MOLES S02 = ,E14.7,15H MOLES C-2 = ,E14.7, 17H TOTAL MOL
 3ES = ,E15.8)

872 CONTINUE

```

C
C *****
C *****
C *****
    FAKE = TCENT
    IF (TCENT .LT. 0.) TCENT = 0.0
C *****
C *****
C *****
C ... GAMMA IS SURFACE TENSION IN ERGS / CM SQUARE, KELVIN TEMPERATURE
    GAMMA = 54.022 + 0.4086*SQRT(TCENT) - 0.059017*TCENT
    TCENT = FAKE
C ... ALPHA IS THE PERCENT OF THE TOTAL NUMBER OF MOLECULES OF H2O AND
C ... S03 IN THE EXHAUST WHICH ARE H2O. ALPHA IS LESS THAN ONE.
    ALPHA = MOLS2 / (MOLS2 + MOLS3)
    BETA = 1.0 - ALPHA
C ... VOLME IS THE VOLUME OF THE DISC AT DISTANCE X FROM THE TAILPIPE
C ... VOLME IS DEPENDENT ON THE TOTAL NUMBER OF MOLES AT P-AMB AND T-ABS
    VOLME = MOLS * R * TABS * CGSAT / (PAMB * FT2CC)
C ... MPV2 AND MPV3 ARE THE NO. OF MOLECULES OF H2O AND S03 PER CUBIC CM
    MPV2 = 4JLS2 * N / (VOLME*FT2CC)
    MPV3 = 4JLS3 * N / (VOLME*FT2CC)
C ... Z IS THE COLLISION FREQUENCY OF WATER AND S03 MOLECULES
    0Z = 2.0 * (2.0 + 4.0*ALPHA*BETA) * (ALPHA*DIAM2+BETA*DIAM3)**2
    ) * MPV2**(2.0*ALPHA) * MPV3**(2.0*BETA)
    2 * SQRT( PI * K * TABS
    3 * ( (ALPHA/MASS2) + (BETA/MASS3) ) )
C
C ... WTPCA IS THE WEIGHT PERCENT IN SOLUTION OF H2SO4 ACID
    WTPCA = 0.95
    WTPCA = 0.95
    WTPCA = 0.95
C
C ... PZERO IS THE SATURATION PRESSURE IN ATMOSPHERES AT ABS TEMP TABS
    A = WTPCA*(0.55490+WTPCA*(-3.14591+WTPCA*3.58243)) + 8.89743
    B = 4TPCA * ( -2361.98 + WTPCA * 3736.88 ) + 2560.87
    PZERO = ( 10.0** ( A - B/TABS ) ) / 760.0
C ... MWBAR IS THE AVERAGE MOLECULAR WEIGHT OF THE CONDENSATE
    MWBAR = MW2 * MW5 / ( MW5 - WTPCA * (MW5-MW2) )
C ... CNST1 IS THE FACTOR CHANGING GRAM-MOLES TO PRESSURE (ATMOSPHERES)
    CNST1 = PAMB / MOLS
C ... DNSTY IS THE DENSITY OF THE NEW CONDENSATE, IN GRAMS PER CUBIC CM
    DNSTY = 1.00070 + WTPCA*(0.80185+0.08819*WTPCA) - 0.0006447*TCENT
C ... RC IS THE MINIMUM RADIUS OF STABLE NUCLEI AT ABS TEMP TABS--CM
    ORC = 2.0 * GAMMA * MWBAR / ( R * TABS * DNSTY *
    1 ALG( CNST1*(MOLS2+MOLS3)/PZERO ) )
    VOLNC = 4.1887902 * ( RC )**3
    TEMP4 = 4.0 * PI * GAMMA * RC*RC / (3.0 * K * TABS)
    IF ( N61 - N62 ) 7016, 7019, 7016
    
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TRAJ P COLE TRAJECTORY
 TRAJ - FFN SOURCE STATEMENT - IFN(S) -

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7016 IF ( J - 10 ) 7018, 7018, 7019
7018 WRITE (5,1238) A, TABS, VOLME, Z, PZERO, ALPHA, TEMP4, MWBAR
1238 FORMAT ( 14, 2HX=, 4E13.5, 4E15.7 )
C ... EXPMN IS THE MINIMUM EXPONENT OF I AS A POWER OF TEN (G.T. 4)
7019 EXPMN = 10.)
C FOR NUCLEATION TEMP4 SHOULD BE L.T. OR EQUAL TO THE FRACTION BELOW
IF( TEMP4 - (23.6-EXPMN)/0.434 ) 7020, 7020, 7020
C ... I IS THE NUCLEATION RATE IN PARTICLES/CC/SEC AT ABS TEMP TABS
7020 I = Z * EXP(-TEMP4)
IF( J - 1 ) 7021, 7021, 7022
C XMIN IS THE VALUE OF X WHERE SIGNIFICANT NUCLEATION FIRST OCCURS
7021 XMIN = X
7022 CONTINUE
C
C ... DELTT IS THE TIME NEEDED TO TRAVERSE DELT2 AT CONSTANT VELOCITY
DELTT = DELT2 / (VAMB + L*PIPE*(VPIPE-VAMB)/(12.0*X))
C CNONS IS THE INCREMENT OF MASS OF CONDENSATE IN DELT2, IN GRAMS
CNONS = I * VOLME*FT2CC * DELTT * VCLNC * DNSTY
C IN A SOLUTION CONSISTING OF ONLY H2O AND H2SO4, THE WEIGHT PERCENT
C OF H2SO4 = THE WEIGHT PERCENT OF SO3 MULTIPLIED BY THE FACTOR
C 1.225015 = 1 + MW(H2O)/MW(SO3) THE RECIPROCAL IS 0.8163159
C WTPCC IS THE WEIGHT PERCENT OF SO3 IN THE CONDENSATE
WTPCC = WTPCA * 0.8163159
C ... UZDUP IS THE NUMBER OF SO3 MOLES NECESSARY IN DELT2 TO FORM
C ... I PARTICLES PER CC, AT TEMPERATURE T-ABS, OF RADIUS RC
UZDUP = CNONS * WTPCC / MW3
C IF THE AMOUNT THAT CONDENSES IS MORE THAN THE AMOUNT LEFT,
C ALTER THE AMOUNT USED TO REFLECT THIS BUT FORCE A TERMINATION
IF( UZDUP - MOLS3 ) 7024, 7023, 7023
7023 DELTT = DELTT * MOLS3 / UZDUP
CNONS = CNONS * MOLS3 / UZDUP
UZDUP = MOLS3 * 1.0001
7024 CONTINUE
IF ( N61 - N62 ) 7025, 7029, 7025
7025 IF( J - 10 ) 7027, 7027, 7029
7027 WRITE (5,1239) CNONS, DELTT, UZDUP
1239 OFORMAT ( 14, 11HCONDENSED =, E13.5, 5X, 8HDELTA T=, E13.5,
1 5X, 8HUSED UP=, E13.5 )
IF( J - 10 ) 7029, 7028, 7029
7028 WRITE (5,12395) CHSO3
12395 FORMAT (11+M) X MWBAR I RC
1 NUMBER TEMPERATURE MOLES H2O MOLES SO3 DENSITY./,
2104H VOLUME
3 ( BEFORE NUCLEATION ),/, 9H CHSO3 = ,F7.4)
C
C ... A POINT IS DESCRIBED AS HAVING ACNTRIVIAL NUCLEATION IF IT CAN USE
C ONE-TWENTIETH PERCENT OF THE SO3 AVAILABLE IN DELT2 AT CONSTANT I
7029 IF( UZDUP - J.0005*AVLBR ) 7048, 7030, 7030
C ... THE IDEA IS THAT IF A POINT HAS A NUCLEATION RATE THAT IS BOTH
C ... ABOVE TEN**EXPMN AND ENOUGH TO USE 0.05 PERCENT OF THE
C ... INITIAL DISTRIBUTION OF SO3, IT HAS SIGNIFICANT NUCLEATION
C ... IF THE AMOUNT USED IS LESS THAN 0.05 PERCENT, AND J IS NOT 1,
C ... INDICATING PREVIOUS SIGNIFICANT NUCLEATION, NO ERROR EXISTS
C ... AN UPPER BOUND OF XLIMIT FT IS CHOSEN AS A PRECAUTION THAT THE
C ... CURRENT CASE WILL HAVE NO SIGNIFICANT NUCLEATION AT ALL
C

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	TRAJ	P CODE	TRAJECTORY
	TRAJ	EFN	SOURCE STATEMENT - IFN(S) -
7030			DELTX = DELT2 DELUZ = JZDUP / AVL8L NEWJ = J - ((J-1)/10) * 10 SAVE(NEWJ,1) = X SAVE(NEWJ,2) = MWBAR SAVE(NEWJ,3) = I SAVE(NEWJ,4) = RC SAVE(NEWJ,5) = I * VOLME*FT2CC * DELTT SAVE(NEWJ,6) = TABS SAVE(NEWJ,7) = MOLS2 SAVE(NEWJ,8) = MOLS3 SAVE(NEWJ,9) = DNSTY SAVE(NEWJ,10) = VULME UZPCT = UZPCT + 100.C*DELUZ MOLS2 = MOLS2 - CNDNS * (1.0-WTPCC) / MW2 MOLS3 = MOLS3 - UZDUP TOT1 = TJT1 + SAVE(NEWJ,5) TOT2 = TOT2 + SAVE(NEWJ,5)*RC TOT3 = TJT3 + SAVE(NEWJ,5)*RC*RC TOT4 = TJT4 + SAVE(NEWJ,5)*RC*RC*RC IF (N61 - 162) 7031, 7034, 7031 IF (NEWJ - 10) 7034, 7033, 7034 7033 WRITE (6,12+0) ((SAVE(JL,KL),KL=1,10),JL=1,NEWJ,5) 1240 FORMAT (14 , 9E13.6, /, 7H , E13.6) 7034 IF (MOLS3) 7051, 7051, 7035 7035 IF (MOLS2) 7040, 7040, 7036 7040 IF (X-XLIMIT) 7036, 7051, 7051 7036 GO TO 5999 C IF J IS NOT 1, THE INCREMENT IN PERCENT OF SO3 USED HAS DECREASED C TO LESS THAN .0005 OF INITIAL SO3. IT IS ASSUMED TO REMAIN LESS 7048 IF (J - 1) 7000, 7000, 7049 7049 J = J - 1 X = X - DELTX IF(N61 - 162) 17049, 7051, 17049 17049 IF(NEWJ - 10) 7050, 7051, 7050 7050 WRITE (6,12+0) ((SAVE(JL,KL),KL=1,10),JL=1,NEWJ,5) C C ... JMAX IS THE MAXIMUM J WITH SIGNIFICANT NUCLEATION 7051 JMAX = J XMAX = X SQ = J4X SO = SJ * DELT2 R1BAR = (TOT2/TOT1) R2BAR = (TJT3/TOT1)**0.5 R3BAR = (TJT4/TOT1)**0.333333 SURFC = 4.J * PI * TCT3 / TOT1 VOLM = .J * PI * TOT4 / (3.0*TCT1) C THE VISIBILITY CRITERION IS 0.025 GRAMS/CUBIC METER OF C PARTICLES OF RADIUS AT LEAST 1/4 * 10**(-4) CM C ... VISBL IS THE MINIMUM VISIBILITY RADIUS, IN CENTIMETERS VISBL = 0.00025 C ... FACTR IS THE FACTOR BY WHICH THE LARGEST PARTICLE'S RADIUS IS C ASSUMED GREATER THAN THE AVERAGE INITIAL RADIUS C ... RMAX IS THE MAXIMUM ALLOWED RADIUS IN THE SIZE DISTRIBUTION RMAX = FACTR * R1BAR SHMRG = FACTR * R1BAR / VISBL

TRAJ - P COLE TRAJECTORY
 TRAJ - EFN SOURCE STATEMENT - (FN(S) -

GREBL = RIBAR / VISBL

C
 C DENSE IS THE AVERAGE CONDENSATE DENSITY AFTER ALL REMAINING WATER
 C VAPOR HAS CONDENSED ON THE 'INITIAL' PARTICLES. THE VOLUME USED
 C TO DETERMINE THE DENSITY IS THE SUM OF THE VOLUME OF THE INITIAL
 C PARTICLES AND THE VOLUME OF THE REMAINING WATER TAKEN AS 1 CC 'GM
 C OF VAPOR, WITH NO PROVISION MADE FOR DECREASE IN VOLUME DUE TO
 C CHEMICAL COMBINATION. THE MASS IS THE SUM OF THE INITIAL MASSES
 C OF H2O AND SO3 AS COMPUTED FROM EXHAUST VELOCITY AND DISC WIDTH.
 C $DENSE = (MW2 * AVLW + MW3 * AVLBR) /$
 C $1 (4.0 * PI * TOT4 / 3.0 + MW2 * PCLS2)$

C ... CONC1 IS THE CONTRAIL DROPLET MASS CONCENTRATION (GRAMS/CU METER)
 C $CONC1 = 100000.0 * (AVLBR * MW3 + AVLW * MW2) / (VOLUME * FT2CC)$

WRITE (6,1070)

1070 FORMAT (14H0* - * - * - *)
 WRITE (6,1071) SQ, XMIN, XMAX, UZPCT

1071 FORMAT (5HIN 4, F6.1, 27H FT INTERVAL EXTENDING FROM, F6.1,
 13H TO, F6.1, 18H FT BEHIND THE JET, F6.1, 39H PERCENT OF THE SO3 M
 MASS HAS CONDENSED.)

C UZOVR IS THE PERCENT OF THE NON-DECAYED SO3 THAT HAS CONDENSED
 C $UZOVR = UZPCT * AVLBL / AVLBR$
 WRITE (6,10712) UZOVR

10712 FORMAT (72H AFTER TAKING INTO ACCUNT THE DECAYED PORTION OF THE O
 ORIGINAL SO3 MASS, F6.1, 39H PERCENT OF THE SO3 MASS HAS CONDENSED/
 2.)

C
 WRITE (6,10715) CONC1

10715 FORMAT (101H0 ASSUMING THAT ALL REMAINING WATER CONDENSES ON THE DR
 IOPLETS, THE MAXIMUM POSSIBLE MASS-CONCENTRATION, /, 69H OF THE
 2 CONTRAIL, TAKING INTO ACCUNT PARTICLES OF ALL SIZES, IS, F7.3,
 320H GRAMS / CUBIC METER)

C
 WRITE (6,1072) TOT1

1072 FORMAT (82H0 THE TOTAL NUMBER OF PARTICLES NUCLEATED (BEFORE AGGLO
 MERATION TO LARGER SIZES) IS, E13.6, 1H.)
 WRITE (6,1073) RIBAR, R2BAR, SURFC, R3BAR, VOLM

1073 FORMAT (33H) FOR THE SET OF INITIAL DROPLETS, /, 27H THE AVER
 AGE RADIUS IS, 6X, E12.5, 4H CM, /, 33H THE AVERAGE SURFACE AREA
 2 IS, E12.5, 36H SQ CM (CORRESPONDING TO A RADIUS OF, E12.5, 9H CM), AND
 3, /, 27H THE AVERAGE VOLUME IS, 6X, E12.5, 36H CC (CORRESPONDIN
 4G TO A RADIUS OF, E12.5, 4H CM)

C
 WRITE (6,1074) RIBAR, RMAX, NFCTR, SHMRG

1074 FORMAT (100H) FOR THE SET OF DROPLETS RESULTING FROM COLLISION, AGG
 LOMERATION, AND CONDENSATION (OF WATER VAPOR), /, 26H THE RA
 2DII RANGE FROM, E11.4, 6H CM TO, E11.4, 16H CM, A FACTOR OF, 15,
 31H, /, 31H AND THE MAXIMUM RADIUS IS, F6.2, 25H TIMES THE VIS
 4IBLE LIMIT., /, 113H THE FREQUENCY DISTRIBUTION OF DROPLET RAD
 SIUS IS ASSUMED TO BE OF THE FORM $F(R) = CONSTANT * RADIUS^{-(N)}$
 6)

IF (RMAX - VISBL) 7775, 7775, 7055

7055 WRITE (6,1079) XMAX, CHSO3

1079 FORMAT (/, 21X, 12H THE EXPCNT, 11X, 7H VISIBLE, 13X, 15H FRACTION OF
 1 THE, 8X, 13H (EVALUATED AT, F6.1, 19H FT BEHIND JET), /, 21X, 60H

TRAJ P COLE TRAJECTORY
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2N OF THE CONCENTRATION TOTAL CONCENTRATION, /,
 379H C5H11S33 FRACTION DISTRIBUTION (GM/GJ METER)
 4 THAT IS VISIBLE, /, 7H IS, F7.4)

```

C
7057 DO 7073 IVDX=1,17,1
      XINDX = IVDX
      EXPON = (XINDX-9.0) / 4.0
C
IF EXPONENT IS ZERO, N = 4 SO USE LOGS (SEE DERIVATION)
IFI EXPON: ) 7061, 7060, 7061
7060 FRCTN = ALUG( SHMRG ) / ALCG( FACTR )
      GO TO 7063
7061 TERM = SHMRG**EXPON
      FRCTN = ( TERM - 1.0 ) / ( TERM - GREBL**EXPON )
7063 CONC2 = FRCTN * CONC1
      EXPRL = 4.0 - EXPON
      WRITE (6,1)EXPRL, CONC2, FRCTN
1085 FORMAT ( 23X, F7.4, 12X, F10.6,15X, F8.5)
7073 CONTINUE
C ***
C ***
C ***
      DELTX = 10.0
      X = X + DELTX
      IF (X-XLIM) 7080, 7080, 1
7080 IF (X-TLINR) 7082, 7084, 7084
7082 COEF1 = -.5/(TLINR-START)
      COEF2 = (TLINR-.5*START)/(TLINR-START)
      COEF3 = COEF1*X+COEF2
      TABS = COEF3*(TEMPX(X)-TAMB)+TAMB
      GO TO 7085
7084 TABS = (TEMPX(X)+TAMB)/2.0
7086 TCENT = TABS-273.16
      RDISK = 1.0 + .0385*X
      VOLME = DELT2*PI*(RDISK**2)
      MWAVE = 1.0/(4JLS2/(VOLME*FT2CC) + .03466098)
      WTAIR=MWAVE*(PAMB*VOLME*FT2CC/(R*TABS*CGSAT)-MOLS)
      MOLS1 = 4JLS1 + WTAIR/MWAVE
      MOLS2 = 4JLS2 + SHAMB*RELHU*WTAIR/MW2
      MOLS5=MOLS5+.20946*WTAIR/31.9988
      MOLS6=MOLS6+.78084*WTAIR/28.0134+.00033*WTAIR/44.00995+.00934*
      WTAIR/39.946
      VPSAT= 10.**( -7.90298*(373.16/TABS-1.0)+5.02808*ALUG10( 373.16/TAB
      1S)-(-1.3316E-7)*(10.**(11.344*(1.0-TABS/373.16))-1.0)+(8.1328E-31)*
      2(10.**( -3.49149*(373.16/TABS-1.0))-1.0)+ALUG10(1313.246))
      ABSHU = VPSAT*MW2*(100.0/(R*TABS)
      SATHU=ABSHU*R*TABS*CGSAT/(PAMB*PHAVE)
      AVLBW = MOLS2-SATHU*WTAIR/MW2
      IF (AVLBW .LT. 0.) AVLBW = 0.0
      MOLS = 4JLS2+MOLS3+MOLS4+MOLS5+MOLS6
      CONC1=100000.*(AVLBR*MW3+AVLBW*MW2)/(VOLME*FT2CC)
      WRITE (6,1)700) X,TABS
10700 FORMAT (8H1 X = ,F6.2,13H, AND TABS = ,F6.2)
      WRITE (6,1)713) CONC1
      WRITE (6,1)79) X,CHSC3
      GO TO 7057
C ***

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TRAJ P COLE TRAJECTORY
- EFN SOURCE STATEMENT - IFN(S) -

C ***

C ***

C ... IF J IS NOT 1, THERE WAS NO NUCLEATION BY XLIMIT FEET--ERROR

7075 IF(J - 1) 7049, 7777, 7049

7775 WRITE (6,1775)

1775 FORMAT (12I10THE FACTOR BY WHICH THE LARGEST RADIUS EXCEEDS THE AVERAGE INITIAL RADIUS IS SMALL ENOUGH THAT THE CONTRAIL IS INVISIBLE)

GO TO 1

7777 WRITE (6,1777) CHS03

1777 FORMAT (531 X GREATER THAN X-LIMIT -- NO NUCLEATION FOR CHS03 = ,
1F7.4)

GO TO 1

END

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13. ABSTRACT This report is a continuation of the final report for contract AF19 (628)-5193. The assumptions and assertions of that report are examined, and estimates made of how these may affect the worth and accuracy of numerical predications of contrail properties. A special type of analytic-experimental analysis is proposed to ferret out nonthermodynamic variables, and is applied to flight test data. Following the analysis, further computer simulation is explored. Additional data lead to bounds on the values of heretofore unspecified parameters. Further experimental study is indicated. The data is presented in terms of the dependence of contrail formation on ambient temperature and relative humidity.()			

14.	KEY WORDS	LINK A		LINK B		LINK C	
		ROLE	WT	ROLE	WT	ROLE	WT
	Fuel Additive	1	1				
	Jet Exhaust	1, 5	3				
	Particle Size Distribution	2	3				
	Contrail	2, 8	3				
	Visibility	4, 8	3				
	Theory	8	2				
	Digital Computer	10	1				
	Mathematics	10	1				
	Thermodynamics	10	1				
	Experiment	8	2				