

The Third International Cloud Condensation Nuclei Workshop

*Reno Workshop Objectives,
Accomplishments,
Instrument Descriptions
and Review Papers*

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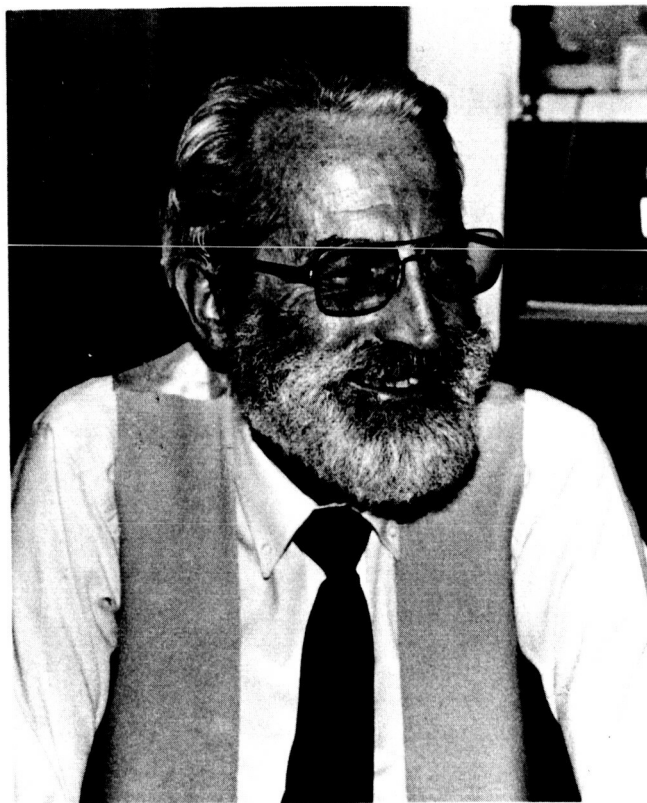
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DEDICATION

It is with great sorrow that we recall the untimely loss of our good friend and colleague, Professor Roger Lincoln Steele.

A man of great talent and undeniable style, Professor Steele contributed extensively to Workshops such as this one, from his scientific involvement in the First International Workshop at Lannemezan, France in 1967, to his responsibilities as Chairman of the Organizing/Steering Committee during the Second Workshop held in Ft. Collins, Colorado in 1970.

Professor Steele will remain in our minds as an individual of uncommon strength and unique character, and it is to his memory that we dedicate these Proceedings.



ROGER LINCOLN STEELE

February 12, 1924 - November 10, 1980

CCN WORKSHOP PARTICIPANTS

Standing, left to right:

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Kneeling, left to right:

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ABSTRACT

The Third International Cloud Condensation Nuclei Workshop was held at the Desert Research Institute, Reno, Nevada, October 6-17, 1980. The goals of the Workshop were to intercompare CCN measurement technology and to perform a limited number of experiments of fundamental scientific interest. A total of 39 scientists representing 20 institutions were in attendance. Twenty-five instruments were tested, including size characterization devices and two Aitken counters. The test aerosols were supplied to the instruments by an on-line generation system, thereby eliminating the need for storage bags. Some of the main conclusions reached during the two-week Workshop were as follows:

(1) Test aerosols of pure soluble salts, both monodisperse and polydisperse, can be provided with stability in output concentration to about $\pm 3\%$ per hour;

(2) Of nine static diffusion chambers (SDC), the five best units (averaged) agreed to within 20% of the NRL mobility analyzer and to within 10% at 1% supersaturation;

(3) Four of the five continuous flow diffusion (CFD) chambers agreed with each other to within about 15% at 0.7% supersaturation and about 20% at 0.3% supersaturation;

(4) The best CFD's and SDC's agreed to within about 15%;

(5) Two of four isothermal haze chambers agreed with each other to within about 40%;

(6) Analysis of the results showed that most instruments' estimation of the CCN spectral slope, k , and the known dry aerosol size distribution slope, β , confirmed the theoretical relationship $k = 2/3 \beta$.

THIRD INTERNATIONAL CLOUD CONDENSATION NUCLEI WORKSHOP

TABLE OF CONTENTS

<u>SECTION</u>		<u>PAGE</u>
	<i>DEDICATION</i>	
	WORKSHOP PARTICIPANTS	-i-
	ABSTRACT	-ii-
I.	INTRODUCTION	1
II.	OBJECTIVES	2
III.	WELCOMING REMARKS	
	H.W. Georgii	3
	Sean A. Twomey	3
	Vincent J. Schaefer	4
IV.	WORKSHOP PROGRAM, PROCEDURES AND FACILITIES	6
	List of Participants	8
	List of Experiments	10
V.	INSTRUMENT DESCRIPTIONS	11
	● Contribution to CCN Workshop Report from University of Wyoming Group	11
	David C. Rogers and Marcia K. Politovich	
	● Description and Discussion of the NRL TGDCC	14
	W.A. Hoppe1 and T.A. Wojciechowski	
	● The CCN Counter of the I.O.P.G. of Puy de Dome: Main Characteristics and Results of Measurements	17
	Roger Serpolay	
	● UMR Dual Mode CCN Counter	20
	D.J. Alofs and M.B. Trueblood	
	● A Cloud Condensation Nucleus Spectrometer Designed for Airborne Measurements	23
	Lawrence F. Radke, Stephen K. Domonkos and Peter V. Hobbs	
	● Static Diffusion Cloud Chamber	26
	Greg Ayers	
	● An Airborne Isothermal Haze Chamber	28
	Edward E. Hindman	
	● A Description of the UK Meteorological Office CCN Counter	33
	M. Kitchen and E. Stirland	
	● Cloud Condensation Nucleus Counter by Impactor Sampling Technique	35
	Takeshi Ohtake	
	● Diffusion Tube	37
	R. Leatch and W.J. Megaw	
	● An Automatic Light Scattering CCN Counter	40
	G. Garland Lala	

<u>SECTION</u>	INSTRUMENT DESCRIPTIONS (Continued)	<u>PAGE</u>
	● Description of the NRL Isothermal Haze Chamber	42
	William A. Hoppel	
	● Modified Mee Industries Static Thermal Gradient Diffusion Cloud Chamber	44
	Randolph D. Borys	
	● Simultaneous Operation of Three CCN Counters and an Isothermal Haze Chamber	46
	James G. Hudson, C.F. Rogers and G. Keyser	
	● Steady Generation of Aerosols with an Improved Constant Output Atomizer	54
	Jack Y. Dea and Ulrich Katz	
	● A. Gagin - Contribution not submitted	
	● T. Mee - Contribution not submitted	
VI.	SUMMARY - REVIEW PAPERS	
	● CCN Comparison of Static Diffusion Chambers	57
	J.E. Jiusto, R.E. Ruskin and A. Gagin	
	● CCN-supersaturation Spectra Slopes (k)	64
	James E. Jiusto and G. Garland Lala	
	● Measurement of the Aerosol Size Distribution with NRL's Mobility Analyzer	69
	William A. Hoppel	
	● Performance of the Continuous Flow Diffusion Chambers.....	79
	James G. Hudson and Darryl J. Alofs	
	● Review of Isothermal Haze Chamber Performance	85
	J.W. Fitzgerald, C. F. Rogers and J.G. Hudson	
	● Aerosol Generation and Distribution System for the Third International CCN Workshop	93
	Ulrich Katz and Jack Y. Dea	
	● Elemental Composition of Aerosols in Fourteen Experiments of the CCN Workshop	97
	William H. Mach and Richard R. Hucek	
	● Comparison of Two Aitken Counters and with Cloud Condensation Nuclei Counters at the 1980 International CCN Workshop	99
	C.F. Rogers and R.L. McKenzie	
	● Comparing Dry Aerosol Measurements	106
	David C. Rogers	
APPENDIX A	TABULATED SIZE DISTRIBUTIONS	
APPENDIX B	WORKSHOP DATA FILE - Computer Printouts and Graphs	

SECTION I. INTRODUCTION

In 1967 the First International Workshop on Condensation and Ice Nuclei was convened under the guidance of Prof. Henri Dessens at his Centre de Recherches Atmospheriques in Lannemezan, France. The Workshop provided a unique opportunity for scientists from all parts of the world to exchange ideas on measurement techniques and to compare instruments for observing ice and condensation nuclei.

A number of important lessons were learned from the first Workshop; principal among them was the need for improved methods of particle generation and characterization, and also the requirement to provide a continuous and stable source of nuclei to the instruments. Solutions to these problems were provided by the combined efforts of several scientists, and at the Second International Workshop, held at Colorado State University in Ft. Collins, Colorado, in August 1970, substantial progress was made in generating, storing and delivering aerosols in the desired size ranges to the more than 25 CCN and Ice Nucleus measuring devices.

In retrospect, the Ft. Collins Workshop was an enormous success, providing researchers an opportunity to learn about the strengths and limitations of the then current instruments used for measuring atmospheric nuclei. At the conclusion of the Workshop, it was determined that, since the needs of CCN and IN counters are so vastly different in terms of aerosol concentrations, separate workshops would be preferable in the future for comparing these two major types of instrumentation. Indeed, this has been the case and, in 1975, an independent Ice Nucleus Workshop was held at the University of Wyoming in Laramie, Wyoming. The objective of that Workshop was to measure and compare methods of ice nucleus measurement; additional useful experiences were gained in methods of generating and delivering ice-forming nuclei to the instrument.

Ten years were to pass before the present Workshop was convened at the Desert Research Institute (DRI) in Reno, Nevada. In the intervening period, progress has been made in instrument design technology, and several new measurement concepts have emerged. Notable among them have been the various types of continuous flow diffusion (CFD) chambers, the isothermal "haze" chambers which are able to operate at extremely low supersaturations, the highly efficient CFD spectrometers and the automated static diffusion chambers. Detailed discussions of each of these types of CCN instruments are provided within these proceedings.

The advances in CCN counter technology have been more than matched by the growth and improvements in aerosol size characterization technology. As just one example, electrical aerosol analyzers are now an off-the-shelf item in use in laboratories throughout the world, whereas at the time of the Ft. Collins Workshop, only a few groups possessed prototype devices.

In view of the many recent innovations in CCN counters, as well as the vast improvements that have been made in particle characterization technology, it was the unanimous decision of the Ad Hoc Commission of the Nucleation Committee, International Commission on Cloud Physics (ICCP), that a high quality instrument comparison workshop be conducted. With the support of the National Science Foundation and the National Aeronautics and Space Administration, such a Workshop was conducted at DRI, Reno, Nevada, October 6-17, 1980, with W. Kocmond and J. Jiusto serving as sponsor Principal Investigators. It is the purpose of these proceedings to document the results and activities of the two-week Reno Workshop.

SECTION II. OBJECTIVES

In the broadest sense, the Third International CCN Workshop had two principal objectives:

- (1) To intercompare CCN measurement technology over a wide range of instrument operating conditions, and
- (2) to perform one or perhaps two fundamental cloud physics experiments of interest to the scientific community.

Because of the strong desire to check each instrument's detection limits and response characteristics under varying operating conditions, most experiments were directed toward this goal. Deliberations by a Workshop Steering Committee (Gagin, Jiusto, Kassner, Kocmond, Megaw, Ruskin and Radke) in conjunction with DRI representatives produced several recommendations with regard to the scientific details of the experiments that were performed. The principal recommendations were:

- (1) that the composition of the test aerosols should include NaCl, AgI and $(\text{NH}_4)_2\text{SO}_4$;
- (2) aerosol concentrations should be in the range of a few hundred to a few thousand per cm^3 active at 0.1 to 1.0% supersaturation;
- (3) ambient air monitoring should be a part of each day's runs;
- (4) on-line continuous generation of aerosols with high stability ($\pm 2\%$) would be preferred over use of a storage bag to supply aerosols to the instruments;

- (5) aerosol samples should be forced through a duct, rather than drawn by suction in order to avoid possible contamination from negative line pressures; and
- (6) both polydisperse and monodisperse aerosols should be generated for the intended instrument checks.

In addition to these recommendations, the consensus held that a few steep sloped aerosol distributions should be produced for tests of instrument responsiveness and also one or two experiments of very high ($> 2000 \text{ cc}^{-1}$ active at 1.0%) and very low ($< 200 \text{ cc}^{-1}$ active at 1% supersaturation) aerosol concentrations should be performed to test each instrument's limits of detection.

With such a broad range of instrument calibration checks, there was less opportunity to conduct specific experiments of fundamental scientific interest. A number of suggestions were considered which included e.g. tests of Kohler theory for a range of particle sizes; comparisons of experiment and theory regarding condensation on particles of limited wettability; checks of temperature dependence of CCN activity in terms of S_c ; and attempts to identify conditions under which particle multiplication may occur.

Partial answers to several of these questions were obtained from the results of experiments performed during the course of the Workshop. Details regarding the findings can be found in the individual summary papers.

SECTION III. WELCOMING REMARKS

Welcoming Address

by

H.W. Georgii

Institut für Meteorologie und Geophysik
Frankfurt, Germany

Ladies and Gentlemen:

It is a privilege for me to welcome you to the Third International Workshop sponsored by the Committee on Nucleation of the International Commission on Cloud Physics and supported by the National Science Foundation and NASA. We are here as the guests of the Desert Research Institute of the University of Nevada, and my special thanks and gratitude go to the host, Professor Warren Kocmond and his associates who provided these beautiful facilities for us. To all of you, I want to express our gratitude. This workshop is mainly devoted to the measurement of cloud condensation nuclei. As a matter of fact, the Second International Workshop, held during August 1970 at Fort Collins had to deal, among other goals, with the following subjects:

(1) Survey of the state-of-the-art in the field of measurement of CCN, (2) compare the operating characteristics of various types of CCN counters, and (3) humidity activation characteristics in various types of CCN counters. While the main effort of the 1970 Workshop, and also of the 1975 Workshop held at the University of Wyoming in Laramie, was devoted to the measurement of ice nuclei, six CCN counters were also tested at Ft. Collins. In the meantime, during the last decade, the importance of CCN for cloud and precipitation physics became more evident. We therefore found it appropriate to devote this Workshop predominantly to the measurement of cloud condensation nuclei.

We became aware in the meantime that only aerosol particles activated at a supersaturation of about 0.5% are of interest to the cloud physicist. We have improved our knowledge on the distribution of CCN in the troposphere and its relation to the atmospheric aerosol in general. This is important, since concentration and composition of cloud nuclei influence directly the average size of cloud drops, the number concentration of cloud drops, the colloidal stability of clouds and the optical density of clouds. The importance of sulfate containing particles as potential cloud nuclei was emphasized during the last ten years by many investigators, and it was confirmed that over oceans the sea-salt particles are only a small fraction of the total maritime cloud condensation nuclei. It could be shown that also over the oceans a large fraction of cloud nuclei is composed of ammonium sulfate or sulfuric acid.

Not long ago, it was assumed that Aitken nuclei are of little importance as cloud nuclei. According to more recent observations, we have to assume that a certain fraction of Aitken nuclei is activated as cloud nuclei. However, the results are still somewhat controversial. A large concen-

tration of Aitken particles does not always lead to a large number of CCN. While measurements in the plumes of large cities in the United States showed an increase of CCN downwind of pollution sources, this was not observed in Israel. In general, it can be said that the concentrations of cloud nuclei over the continent range from 100 to 1000/cc while over the oceans they range from some tens to a few hundred/cc. These values are in good agreement with the drop concentrations in continental or maritime clouds. From this point of view, it appears that the cloud nuclei counters used in these investigations detect the right fraction of the atmospheric aerosol activated in the process of cloud formation. One problem, still unsolved, is the possible long-term trend of the cloud nuclei concentration on a global scale. One major source of cloud nuclei is probably the emission of reactive gases and the subsequent formation of secondary nuclei by gas to particle conversion. A long-term increase of the cloud nuclei population will certainly influence the efficiency of the rain-forming process.

During the 1970 Workshop, five CCN counters of the thermal diffusion principle showed satisfactory results and good agreement with natural aerosols. The agreement among the instruments was less satisfactory for artificial aerosols. In the meantime, more sophisticated instruments have been developed and we are looking forward with interest to the experimental phase of this Workshop. Cloud nuclei have become a more and more important fraction of the atmospheric aerosols. I therefore believe that this Workshop is very timely and provides the necessary international platform to study and to discuss the progress which had been made during the last ten years and to give the necessary directives to researchers in this field.

Welcoming and Keynote Address

by

Sean A. Twomey*
University of Arizona
Tucson, Arizona

A. Historical Perspectives

Early workers (e.g., P. Squires) were aware that there were many more condensation nuclei than cloud droplets; the realization came that it was important to make measurements at low supersaturations rather than with Aitken counters. Discussions in the early 1950's turned to the problem of detecting and counting the small cloud droplets that would be produced at small, cloud-like supersaturations.

* This summary of Dr. Twomey's Welcoming and Keynote remarks has been composed based upon notes provided by the author.

The creation of small supersaturations under isothermal conditions (rather than by adiabatic expansion) seemed highly desirable, leading to the conception of the chemical diffusion chamber. At that time (middle 1950's), Wieland in Switzerland had constructed a thermal gradient diffusion chamber and used it to nucleate and grow droplets at small supersaturations. Detection and counting in Wieland's chamber was accomplished by examination of a sticky layer on the chamber floor, into which droplets fell and were hopefully preserved. Twomey introduced photographic detection, and soon the thermal gradient diffusion chamber, with large diameter-to-height ratio, cooled at the base (for convective stability and to avoid transient supersaturations), became the standard device. Photography became the standard detection method, although problems such as nominally "fast" film being sometimes slower than nominally "slower," but less grainy films, remained to be sorted out. Enough devices were in operation that an early comparison workshop could be held at NRL's Chesapeake Bay Annex in 1965.

Whatever the detection method, it will have a threshold - a minimum detectable size of droplet, and it is evidently vital to determine or estimate that, since otherwise one may be counting inactivated (haze) droplets, or not counting all activated droplets. At very low supersaturations an inactivated droplet may be several microns radius and a clear distinction between "haze" and "cloud" (in terms of size) no longer exists. Under such conditions, a simple counting procedure is hardly sufficient and some method of sizing is required, essentially calling for a different technology.

At about the same time, NRL's airborne CCN studies began, and soon investigators were able to compare CCN counts to cloud droplet concentrations. The comparison results were, in general, satisfactory.

B. Points Concerning the Sizes and Composition of CCN

An aspect of typical cumulative CCN distributions fortunate for cloud physics is that there is found a convenient and simple proportionality between critical supersaturation and numbers of activated cloud droplets:

$$N = CS^k,$$

where k is typically less than unity and often around 0.5 or so. The parameter k could equally well have been much larger, or the simple power law could have been replaced by some less tractable functional relationship. At very low critical supersaturations, k does tend to increase to greater than unity, creating a more difficult situation where the number of droplets activated is critically dependent upon small variations in the ambient supersaturation, and hence on the detailed time evolution of temperature before and during the condensation processes.

The sizes of natural CCN have been shown by several experiments to be close to the minimum size that Köhler's theory (of the critical supersaturation of pure soluble electrolytes) would allow. Again, it is fortunate that most CCN of interest seem to be soluble compounds, not simply insoluble, wettable particles of various contact angles.

It has long seemed that the most likely candidates for these soluble compounds were sodium chloride, ammonium sulfate, and sulfuric acid. Ammonium sulfate continues to be regarded as a predominant constituent.

C. Gaps in the Understanding of CCN

The sources of CCN are not completely understood, although gas-to-particle processes must be a major contributor. The speaker described observations of CCN production occurring over several-hour periods in Arizona, as well as diurnal cycles in the CCN count at the Robertson site in Australia. Related observations by J. Hudson and J. Jiusto were mentioned. CCN are apparently rather transient in nature, with a lifetime of no more than a couple of days. Indeed they cannot exhibit a diurnal cycle if their life expectancy is much longer.

The region of the size spectrum between Aitken nuclei and CCN is another unknown; the ratio of Aitken particle concentrations to CCN concentrations is highly variable, and it seems unlikely that any simple extrapolation or interpolation to connect the two will be sufficient. And to completely understand the progenitors of CCN, it is important to obtain information on the size range from Aitken nuclei on down to molecular clusters, although this region may not be directly relevant to CCN.

The opposite end of the spectrum - from CCN up to "giant" nuclei - is sometimes of influence in cloud droplet growth calculations. It is certainly the critical range in slow condensation processes such as many fogs may be.

A critical instrumental shortcoming is the minimum detectable size of nuclei; some aerosol workers quote $0.01 \mu\text{m}$ as the minimum sensitive size for particular expansion counters, too large to be helpful in resolving the problems just mentioned.

Welcoming Address

by

Vincent J. Schaefer
State University of New York
Albany, NY

I am very pleased to have the opportunity to participate in this Workshop on instruments for measuring cloud condensation nuclei. I am particularly intrigued to see the number of young scientists now involved in this interesting and important aspect of cloud physics.

About 25 years ago, Ted Rich of the General Electric Company and I (then Director of Research of the Munital Foundation) gathered most of the persons interested in atmospheric nuclei and held a three-day conference on this fascinating subject. Only about 20 scientists and engineers could be found in the United States. During this conference, we identified most of the problems that are still with us, but our techniques then were somewhat primitive when compared with the sophisticated electronic equipment now being used in the laboratory, as well as in the field and on aircraft.

I was intrigued with the comments of Dr. Georgii, which in his absence were read by Dr. Dieter Stein. I am indeed sorry to learn that due to illness Dr. Georgii will not be attending this Workshop. I had looked forward to seeing him and hope he is well on the road to recovery. Several statements in his remarks were particularly interesting to me, since I have spent a great deal of time in many parts of the world in an attempt to establish the aerosol concentration patterns in a wide variety of environments. My findings agree completely with the patterns mentioned by Dr. Georgii.

Using the portable and highly reliable Gardner counter with which most of you are familiar, I have found consistent patterns in the concentration of both condensation and cloud condensation nuclei in very clean as well as in very polluted air.

It is well recognized that the Gardner counter provides a very good indication of the concentration of Aitken nuclei when used to provide supersaturations in excess of 300 percent. I have found that this instrument can also be used to provide consistent, as well as semi-quantitative measurements of the number of cloud condensation nuclei in an air sample. If instead of using 20 to 27 inches of mercury vacuum only one scale division is used, the Gardner provides a rather good measurement of the concentration of particles active as water condensing nuclei (CCN) at 1 percent supersaturation.

There are several reasons why this method of using this instrument is frowned upon! Ted Rich who, as you know, invented the so-called Gardner counter was the first to tell me that such a procedure was highly irregular and meaningless. He cited intricate relationships in the life cycle of the cloud droplets forming on nuclei at various supersaturations which he had measured during the development of his instrument.

Despite this disapproval, I have made many thousands of measurements using this instrument in its low vacuum mode as well as high and median settings. I have found that the results provide an extremely consistent measurement of the concentration of cloud condensation nuclei in a particular type of environment. Thus in particle-free air I read zero particles. In very clean air, such as the stratosphere, mid-ocean, the Fiji Islands, mountain summits, caverns, deep forests, and similar places far from man's influence, my measurements of cloud condensation nuclei (CCN) range from 0 to 150 CCN cm^{-3} when the Aitken particle concentration ranges from 200 to 1000 particles per cubic centimeter.

At the other extreme, in cities, the plumes of heavy industries, large airports, superhighways, vehicular tunnels, and the like, the values I find range from 800 to 3000 cloud condensation nuclei and 50,000 to 300,000 Aitken nuclei per cubic centimeter.

I have classified the natural and anthropogenic aerosol concentration measured mostly in the Northern Hemisphere and have summarized these findings in three volumes which have recently been published by our Research Center and which I'll be glad to send to participants of this Workshop while the supply lasts.

I hasten to add that what I have just said does not in any way suggest that the Workshop which is starting with this meeting is not extremely important. If progress is to be made in our cloud physics studies, it is extremely important that all measuring instruments used to establish scientific facts such as the concentration of atmospheric particles should agree with each other. Only then will we be able to communicate with each other in a meaningful way.

As most of you who have worked in the atmosphere know, the concentration of airborne particles varies over a considerable range, no matter where it is measured. It is necessary when establishing an aerosol "climatology" at a given geographic location to determine the range in concentration over a period of time, its diurnal pattern and the effects that appear with wind direction and atmospheric stability. It is also necessary to measure seasonal variations.

On the west coast of Ireland, for example, with a southerly or southeasterly flow the particle concentration pattern shows the characteristic of continental air (1000 CCN cm^{-3} /40,000 AN cm^{-3}) while with a northerly or northwesterly flow it is Polar Maritime (100 CCN cm^{-3} /800 AN cm^{-3}). Even in a region noted for high pollution levels such as the Los Angeles Basin, a strong persistent flow of oceanic air will drive the Basin pollution through the mountain passes and even over the crest of the Sierra to affect the air quality of Las Vegas, Phoenix, Flagstaff and more distant places. When this occurs, the particulate levels which could have been 3000 CCN cm^{-3} /300,000 AN cm^{-3} can drop to 500 CCN cm^{-3} /10,000 AN cm^{-3} . When this happens, the visibility is greatly improved and it is possible to see what a delightful place the Basin would be if its population was reduced by an order of magnitude!

I hope when this Workshop is ended and we collectively gain a higher appreciation of the performance of our instruments that extensive field measurements can be mounted so as to establish confidence in the meaning of our data as they relate to atmospheric visibility, the genesis and nature of storms, the stability of clouds and the formation of precipitation, as well as the basic causes of such specific occurrences as acid rain.

I look forward to participating in this Workshop and look forward to meeting you all and seeing your equipment in satisfactory operation.

Have fun!

SECTION IV. WORKSHOP PROGRAM, PROCEDURES AND FACILITIES

In developing a plan for the Third International CCN Workshop, initial efforts were carried out by an Ad Hoc Commission of the Nucleation Committee of the International Committee on Cloud Physics (ICCP); this Ad Hoc Commission, appointed by Professor Georgii, Chairman of the Nucleation Committee, included J. Kassner, J. Megaw, L. Radke, K. Whitby, and J. Jiusto, Chairman. The Ad Hoc Commission returned the recommendation that the Desert Research Institute's offer to host the meeting should be accepted and that a Steering Committee composed of A. Gagin, J. Jiusto, J. Kassner, W. Kocmond, J. Megaw, L. Radke, and R. Ruskin should proceed with the planning of the specific logistical and scientific details of the meeting.

The Steering Committee functioned until the end of the Workshop on October 17, 1980, as the principal scientific decision-making body. A group completely local to DRI, the "Local Arrangements Committee," handled logistical matters such as participant travel arrangements and lodging, the shipping of instruments to and from Reno, and the creation of suitable laboratory space. Individual members of the DRI staff took responsibility for technical matters such as design of an aerosol generation system which would meet the goals set by the Steering Committee, determinations of instrument power supply and heat rejection requirements, and tentative placement of instruments along the aerosol supply duct.

In the meantime, correspondence between the offices of the Principal Investigators and interested individuals, together with announcements in the Bulletin of the American Meteorological Society and the *Journal de Recherches Atmospheriques*, resulted in a tentative list of over 40 participants, later to be somewhat reduced to the actual attendees as shown in Table 1.

As the effort proceeded into the summer months of 1980, it was decided that some members of the Steering Committee could meet during the International Cloud Physics Conference, Clermont-Ferrand, France (July 1980). Three Steering Committee members who attended this meeting held discussions with a contingent of three DRI representatives; out of the talks came a number of important recommendations that were incorporated into the Workshop Objectives (see Section II).

At DRI, U. Katz took primary responsibility for the aerosol generation and distribution system (Figure 1); a more thorough description of the system can be found in Section V. Aerosol sizing, also described in Section V, was "officially" provided by the Naval Research Laboratory at the request of the Steering Committee.

The final physical and logistical arrangements were handled by the Local Arrangements Committee, mainly through Mrs. Jo Janowski and the staff of the DRI machine shop. Under their supervision, the large components of the aerosol delivery system were installed, provision was made for both 50 Hz

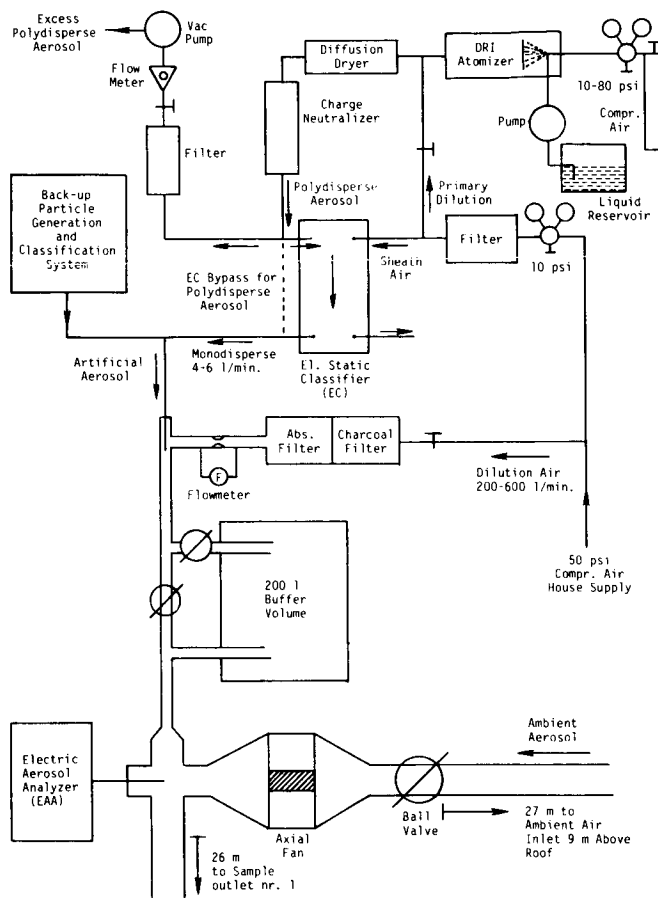


Figure 1. Schematic of Aerosol Generation System

and 60 Hz electricity totalling over 80 kw delivered to outlets spaced along the entire length of the aerosol duct, and 12 new laboratory benches were built. Travel and lodging arrangements were finalized for 22 participants from the USA and 7 from overseas. A total of 17 instruments of various types had to be transported to DRI; in the case of instruments from overseas, customs inspections and clearances were involved.

With most participants and instruments in Reno by October 6, the Workshop opened with welcoming remarks by Prof. W.C. Kocmond, Prof. H.-W. Georgii (represented by Mr. D. Stein) and Dr. V. Schaefer. A keynote address was given by Dr. S. Twomey. Two weeks of intense activity then began.

The Steering Committee met daily to plan the test aerosols to be used; generally their deliberations were conducted at the beginning of each day, while instrument operators readied their devices and monitored outside ambient aerosol. Table 2

summarizes the 30 experiments actually performed. The specifications for each test aerosol varied day-to-day, in response partly to earlier suggestions originating in the Steering Committee or coming from individual participants:

(a) High CCN concentrations (relevant to the atmosphere in areas of high pollution, volcanic plumes, etc.) to establish the practical upper limit of applicability of various instruments;

(b) Low CCN concentrations (e.g., less than 100 cm^{-3} at 1%), to check instrument performance in clean maritime or polar environments;

(c) A bimodal size distribution (two monodisperse, soluble aerosols of different nominal sizes), to check the ability of instruments to resolve a correspondingly bimodal CCN activity spectrum.

With "feedback" from participants, these suggestions were incorporated into the rather tight experiment schedule (suggestion "a" is reflected in experiments 9 and 24, "b" in 4 and 23, and "c" in 10). Additional suggestions during the Workshop led to experiment 21, a test of the ability of instruments to give a consistent and accurate reading over a prolonged period of time, CCN concentration being held constant; experiment 25, a test of the "zero" of instruments sampling particle-free air, and experiment 29, a test of the possible CCN activity of a hydrophobic aerosol, paraffin wax.

Daily activities included frequent discussions between members of the Steering Committee and the larger body of participants. Steering Committee members assisted with the presentation of collected data after each experiment; in addition, participants presented informal talks on their equipment on a daily basis.

At the end of the Workshop, the Steering Committee completed its function by requesting all participants to supply a description of their instruments together with comments on performance during these experiments; those reports are assembled in Section V. In addition, various individuals were asked to submit reports on special interest topics, such as reviews of the CCN counters by generic type; those reports are to be found in Section VI.

The final action of the assembled group was to provide verbal and written response to the Steering Committee's request for criticisms and suggestions for improvements. The following points resulted:

(a) Although previously debated, the question of whether or not to standardize data-taking such that all CCN counters would measure CCN concentrations at a few, given "set-point" supersaturations should be reviewed again. The practice at this Workshop was to avoid specific, standard settings (because of different instrument requirements) and to provide as complete a spectrum as possible. "Set-points" would simplify data comparisons but, of course, not all instruments are designed to operate at the same supersaturations.

(b) It was intended that the results of each experiment would be quickly entered onto computer file. In practice, this effort, which ideally could have provided data listings and computer graphics in short turn-around time, was hampered by difficulties in design and planning as well as

equipment problems. Future Workshops would benefit from rapid turn-around of the data, but better control of the experimenter input is also needed.

(c) Power and aerosol supply were in reasonably good shape; ambient heat rejection (air conditioning) and physical space had less margin of comfort but were satisfactory. Noise level was high at times; one participant suggested that a vacuum manifold (replacing numerous individual pumps) would alleviate this problem.

In summary, most participants felt that the Workshop proceeded quite smoothly and that the objectives set forth by the Ad Hoc Commission of the Nucleation Committee were successfully achieved. In making this determination, special mention must be made of the close cooperation between participants, Steering Committee and DRI hosts. Several social events were held during the 12 day Workshop, including an open house hosted by Dr. Cliff Murino, President of DRI. The strong sense of friendship and good times that develop in a working environment such as this will always be remembered.

TABLE 1. CCN WORKSHOP PARTICIPANTS

<u>Name/Affiliation</u>	<u>Instrument</u>	<u>Duct No.</u>	<u>Name/Affiliation</u>	<u>Instrument</u>	<u>Duct No.</u>
Dr. Jeffrey B. Anderson Space Science Lab NASA-MSFC Alabama 35812			Dr. James G. Hudson Desert Research Institute University of Nevada System P.O. Box 60220 Reno, NV 89506	CFDCC 3SS CFDCC IHC	14,16,18
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Dr. W.A. Hoppel, Code 4320 Atmospheric Physics Branch Naval Research Laboratory Washington, DC 20375	Aero. Sizing	12			
Mr. Richard Hucek Florida State University Dept. of Meteorology Tallahassee, FL 32306	CFDCC Impactor	22, 23			

<u>Name/Affiliation</u>	<u>Instrument</u>	<u>Duct No.</u>	<u>Name/Affiliation</u>	<u>Instrument</u>	<u>Duct No.</u>
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Dr. W.J. Megaw Department of Physics York University 4700 Keele Street Downsview, Toronto CANADA M3J1P3	Diffusion Tube	7	Dr. V.J. Schaefer State University of New York Albany, NY 12222		
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Ms. Marsha Politovitch University of Wyoming Dept. of Atmos. Science Box 3038, Univ. Station Laramie, WY 82071	STGDCC Aerosol Sizing	1,2	Dr. Sean Twomey Inst. of Atmospheric Science University of Arizona Tucson, AZ 95721		
Dr. Lawrence F. Radke Dept. of Atmos. Sciences University of Washington Seattle, WA 98195	4SS CFDCC	20	Mr. C.H. Wilson NASA Langley Research Center MS 404B Hampton, VA 23665		
Dr. C. Fred Rogers Desert Research Institute University of Nevada System P.O. Box 60220 Reno, NV 89506	DRI-NASA CFDCC	15	Dr. T. Wojciechowski Naval Research Laboratory Atmospheric Physics Branch Washington, DC 20375	STGDCC	17

Table 2. List of Experiments

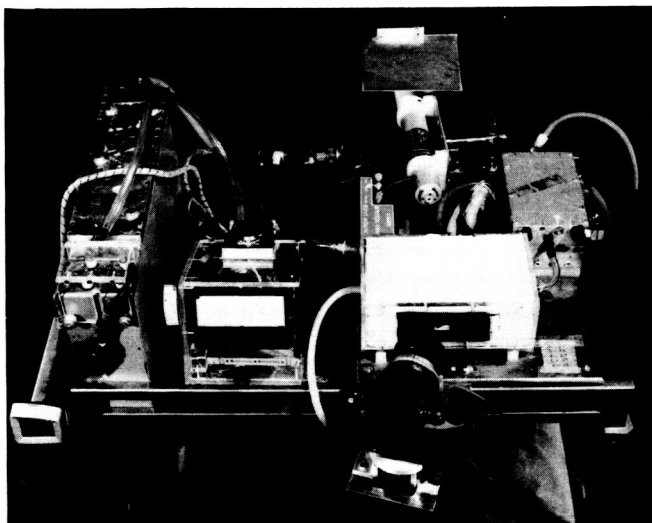
No.	Date	Aerosol	
		A=Ambient; M=Monodisperse; P=Polydisperse	
0	Tues 7 Oct	AM	M - $(\text{NH}_4)_2\text{SO}_4$
1	Tues	PM	P - NaCl - oscillating concentration
2	Tues	PM	P - NaCl - higher concentration
3	Wed 8 Oct	AM	A - quite fluctuating
4	Wed	AM	M - NaCl - low concentration
5	Wed	PM	M - NaCl - medium concentration
6	Wed	PM	A
7	Thurs 9 Oct	AM	A - aborted - duct blockage
8	Thurs	AM	M - NaCl - slight drift down
9	Thurs	AM	M - NaCl - higher concentration
10	Thurs	PM	Bimodal - NaCl - "flat k"
11	Thurs	PM	A
12	Fri 10 Oct	AM	A
13	Fri	AM	P - $(\text{NH}_4)_2\text{SO}_4$
14	Fri	PM	P - $(\text{NH}_4)_2\text{SO}_4$
15	Fri	PM	M - $(\text{NH}_4)_2\text{SO}_4$
16	Fri	PM	A
17	Mon 13 Oct	AM	A
18	Mon	AM	M - $(\text{NH}_4)_2\text{SO}_4$
19	Mon	AM	M - $(\text{NH}_4)_2\text{SO}_4$
20	Mon	PM	M - $(\text{NH}_4)_2\text{SO}_4$
21	Mon	PM	M - $(\text{NH}_4)_2\text{SO}_4$ - time variations
22	Tues 14 Oct	AM	P - $(\text{NH}_4)_2\text{SO}_4$ - medium concentration
23	Tues	PM	P - $(\text{NH}_4)_2\text{SO}_4$ - low concentration
24	Tues	PM	P - $(\text{NH}_4)_2\text{SO}_4$ - high concentration
25	Wed 5 Oct	AM	Filtered air - noise check
26	Wed	AM	A
27	Wed	PM	P - AgI, "insoluble"
28	Wed	PM	M - AgI, "insoluble"
29	Wed	PM	P - paraffin, hydrophobic

SECTION V. INSTRUMENT DESCRIPTIONS

CONTRIBUTION TO CCN WORKSHOP REPORT FROM UNIVERSITY OF WYOMING GROUP

David C. Rogers and Marcia K. Politovich

University of Wyoming
Laramie, Wyoming



1. APPARATUS

The University of Wyoming's CCN counter is a static, horizontal, parallel plate thermal gradient diffusion chamber of rather conventional design. Its intended use is primarily for field measurements, hence the small physical size and straightforward simplicity of operation. The plate separation is 0.9 cm, and the inside chamber dimensions are 8.5 cm x 10.0 cm; aspect ratio is 9.4:1. Activated CCN grow to visible size droplets which are photographed with a 35 mm camera (Nikon F, Micro-Nikkor-P lens f3.5, 55 mm plus M2 extension tube, Tri-X film developed at ASA 1600). Illumination is provided by a Helium-Neon laser (0.6328 μm wavelength, multimode 5 mw) oriented at an angle of 23° from the forward direction of the camera's optical axis. The multimode character of the laser provides a flat-top intensity profile which serves to reduce uncertainties about the size of the illuminated volume. The angle of 23° in the forward direction was chosen to take advantage of the first broad peak in the Mie scattering function for water droplets which are in the size range 3 to 7 μm diameter. The circular laser beam is 0.18 cm in diameter and is centered in the chamber to illuminate the middle 20% of the vertical distance. A fixed width on the film is used to define the horizontal dimension of the sample volume; this volume is .034 cm^3 , in the form of an elongated cylinder.

The temperature difference between the top and bottom plates is measured by precision thermistors which are flush-mounted between the aluminum plates and the surface wicking material (blotting paper).

This temperature measurement is displayed to $\pm 0.1^\circ\text{C}$ and is also used by an electronic circuit to control the temperature difference. The measurement is compared with a value selected by the operator, and the difference is used to control the current to thermoelectric modules which cool the bottom plate. The range of temperature differences normally used extends from 3 to 7°C; this results in a range of supersaturations of approximately 0.3 to 2%.

Sample air passes through a temperature preconditioning chamber just before entering the CCN chamber. The preconditioning chamber is maintained slightly warmer than the top plate of the CCN chamber. In this manner, transient supersaturations are minimized. Air samples are brought into the chamber under suction and, after thorough flushing, the chamber outlet is closed. A time delay of several seconds between this closing time and the photography allows droplets to grow large enough to be photographed but not so large that they fall out of the illuminated volume; this delay is controlled and decreases with larger supersaturations.

Earlier calibration experiments using monodisperse latex particles determined the photographic minimum detectable particle size is less than 0.7 μm diameter. Aerosol losses in the entrance region have been measured as negligible by comparing size distributions of various aerosols before and after passing through the chamber. Finally, earlier comparisons with theory and other CCN chambers have been performed for monodisperse and polydisperse salt aerosols as well as natural aerosols. Comparisons between the University of Wyoming's (UW) chamber and that of the Desert Research Institute (DRI) were made during March 1978. These comparisons are briefly summarized in Tables 1 and 2, and Figure 1.

TABLE 1. CFD-SDL COMPARISONS

Aerosol	Super-saturation(%)	Concentration (no. cm^{-3})		
		SDL	CFD	CFD/SDL
Room air	.28	95	168	1.77
	.36	207	288	1.39
	.50	164	219	1.34
	.65	243	352	1.43
	1.02	1310	3596	2.75
Outside air	.35	320	241	0.75
Polydisperse NaCl	1.00	327	497	1.52
Polydisperse AgI	1.00	371	613	1.65
		368	544	1.48
Monodisperse AgI (0.2 μm dia.)	0.2	485	750	1.55
	1.0	485	810	1.67
CFD/SDL avg. $1.57 \pm .47$				
Omitting 2 extremes, avg. $1.53 \pm .14$				

TABLE 2. CRITICAL SUPERSATURATION

Aerosol	Critical Supersaturation(%)		
	SDL	CFD	Theory
NaCl .03 μm diameter	.36	.32	.35
NaCl .05 μm diameter	.75	.68	.75
AgI-NH ₄ I complex (0.2 μm dia.)	<0.2	<.25	.07

SDL = Static Diffusion Liquid CCN Chamber (UW)
 CFD = Continuous Flow Diffusion CCN Chamber (DRI)

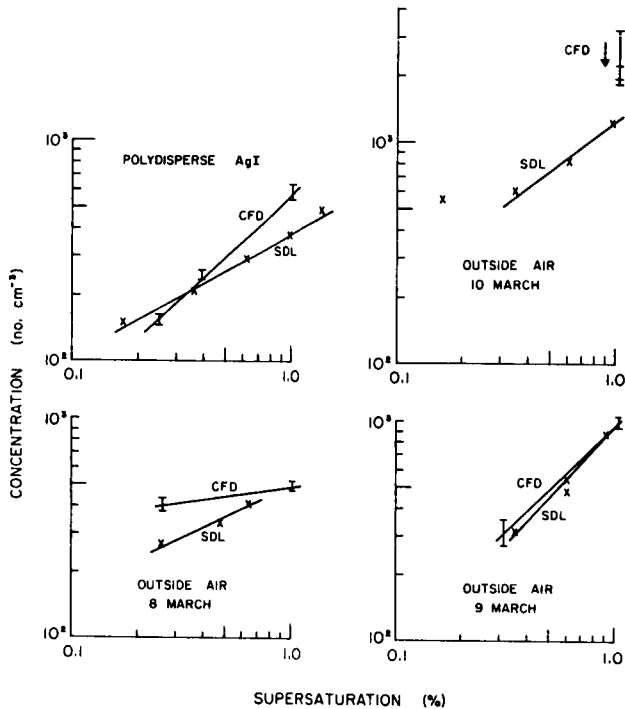


Figure 1. CCN spectra measured in Laramie, March 1978. SDL = Static Diffusion Liquid CCN Chamber (UW); CFD = Continuous Flow Diffusion CCN Chamber (DRI).

2. WORKSHOP PARTICIPATION

Our main interests in participating in the International CCN Workshop were to compare the University of Wyoming's CCN chamber with the others present and to discuss with and learn from the participants various aspects of CCN measurement techniques, their current thinking, and advances in the ten years since the last workshop.

Unfortunately, problems with the electronics in our device arose during the Workshop and prevented us from operating the chamber above about 0.3% supersaturation, so our comparison experiments were attenuated.

3. DATA USAGE

The emphasis of our work with the CCN counter has been in the application of CCN data obtained from it, rather than development of the instrument itself. We have kept the design simple and the size small to enable us to transport it to remote sites for in situ sampling. We include here several examples to illustrate the manner in which the data are used.

Figure 2 shows a C-K plot of measurements obtained at and near our cloud observation facility at Elk Mountain in southeastern Wyoming. These measurements were taken to characterize winter CCN populations in southeastern Wyoming, particularly near Elk Mountain. The data suggest a trend of increasing C-values later in the spring while retaining similar K-values. There was significant snow cover in and upwind of the sampling location during January and February. In later months, snow cover was sparse if present at all.

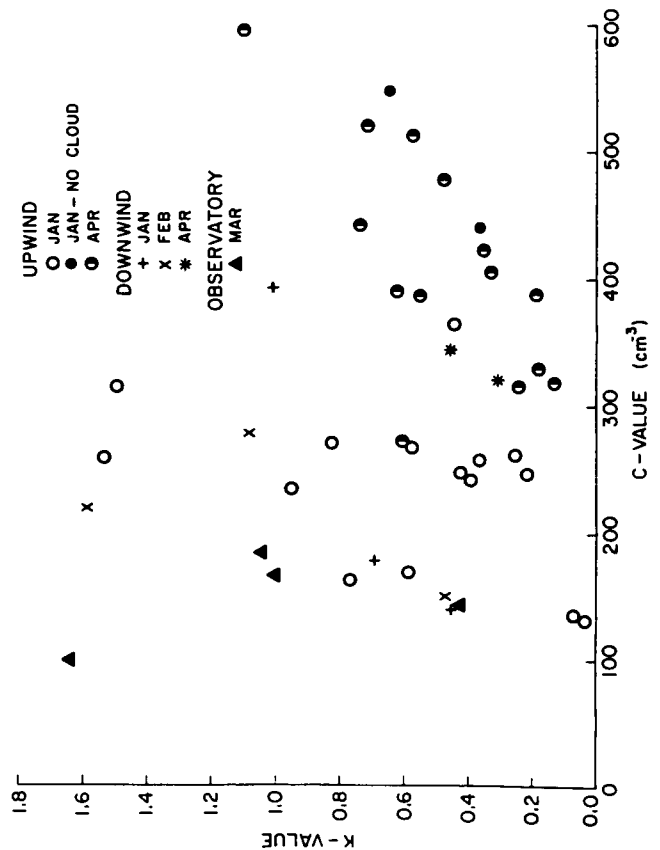


Figure 2. C-K plot of the CCN spectra measured during the winter of 1978-79 near Elk Mountain. Sample months and locations are indicated.

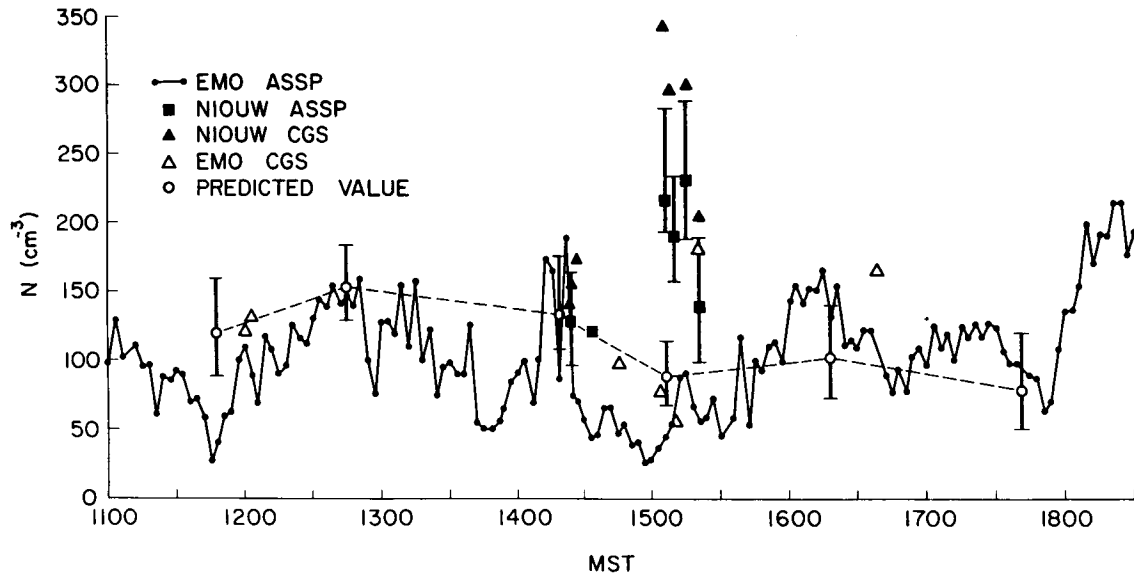


Figure 3. Droplet measurements obtained in an orographic cloud which formed over Elk Mountain on 16 January 1979. Data sources are indicated (ASSP is the Axially Scattering Spectrometer Probe, Particle Measuring Systems, Boulder, CO, and CGS is the University of Wyoming's soot-coated impactor slide droplet sampler). CCN predictions represent application of Twomey's (1959) relationship between CCN spectrum parameters and droplet concentrations to measurements taken upwind of Elk Mountain. (O) represents an updraft of 50 cm/s, (I) represents the range of updrafts 25-100 cm/s. A 20-minute time lag was added to CCN data to account for transport time from the CCN sampling site to the droplet measuring site.

An important use of our CCN data has been in comparisons of droplet concentrations derived from the upwind CCN spectra (using Twomey's, 1959, equation¹ and Young's, 1979, cloud model²) with in situ measurements from the observatory and instrumented aircraft in the Elk Mountain cap clouds. Figure 3 shows the results of such a comparison on 16 January 1979. The CCN predictions track the actual measurements from the observatory well. Droplet concentrations measured by the aircraft (NIOUW) were higher than those from the observatory (EMO) on this day, which we attribute to variations in the vertical structure of aerosol concentrations in the boundary layer, which were measured by the aircraft, rather than substantial increases in updraft speeds at that level.

These figures are from a paper by M.K. Politovich which is being prepared for publication.

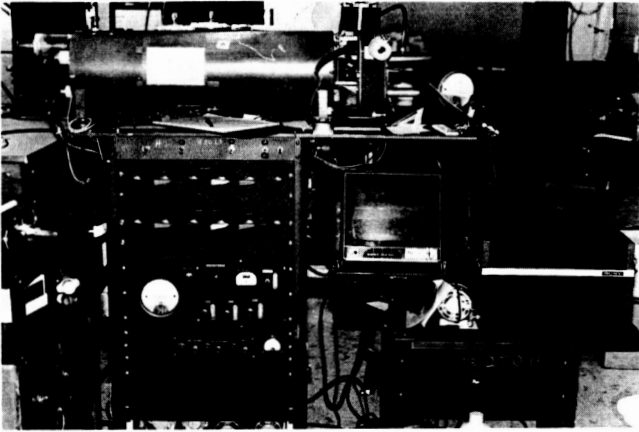
¹Twomey, S., 1959: The nuclei of natural cloud formation, Part II: The supersaturation in natural clouds and the variations in cloud droplet concentration. *Geophysics Pure. Appl.*, 43, 243-249.

²Young, K.C., 1974: A numerical simulation of wintertime, orographic precipitation. Part I: Description of model microphysics and numerical techniques. *J. Atmos. Sci.*, 31, 1735-1748.

DESCRIPTION AND DISCUSSION OF THE NRL TGDC

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The NRL thermal gradient diffusion cloud chamber (TGDC) is identical to that described in the Proceedings of the Second International Workshop on Condensation and Ice Nuclei (Grant, 1971). The chamber consists of two plates 7.5 cm in diameter separated by 1.25 cm and covered with saturated filter paper. The cylindrical wall separating the plates is glass. The top plate is at room temperature and the bottom plate is cooled with a thermoelectric cooler. The temperature difference is measured with several sets of thermocouples. A high intensity mercury arc lamp with appropriate lens and collimating slits which are imaged in the cloud chamber defines an illuminated volume 0.15 cm thick and 0.3 cm high by 1.5 cm long. This illuminated volume is viewed at 90° scattering angle by an 8 mm camera and a video camera system.

The video system permits immediate playback and stopframe counting of the cloud droplet concentration in the chamber. The 12-inch monitor screen is marked off to identify the viewing dimensions for ease in counting the droplet images.

At the Workshop, samples for a given experiment were admitted to the TGDC directly from the sampling duct which was slightly pressurized. The count was recorded both on video tape and on a number of frames of an 8 mm movie film. The 8 mm Bolex camera is operated manually while viewing the chamber through the camera. The first frame is taken one to two seconds after closing the valve and about 10 to 15 frames are taken manually until it is clear that the maximum particle count has passed. The photographic results were not available at the Workshop. The video data were read immediately after each experiment. The method for obtaining the count supplied at the Workshop is as follows: two (or three) successive samples were recorded at each supersaturation. The recordings from these samples were played back and visually

examined to obtain the succession of frames for each sample where the maximum count occurred. Several of these frames were counted and the maximum count obtained. The maximum counts from two (or three) samples were averaged to obtain the data submitted at the Workshop.

The photographic recordings have not yet been analyzed but it has always been our experience in the past with atmospheric aerosols that the two methods track very well but that the photographic count is always about 10 to 15% higher. We attribute this to the smaller detectable size obtained with the photographic method. The minimum detectable size for our photographic system is estimated to be about a half micron whereas the minimum detectable size for the video system is estimated to be just under one micron.

There are definite limits on the supersaturation range for which the results of the TGDC are valid. The range of validity is usually given as 0.2 to 1%. However, even within this range there can be significant errors depending upon the nuclei spectrum (size distribution) being measured. The accuracy limitations of TGDC's in general and our chamber in particular were investigated by Hoppel and Wojciechowski (1976).

The smaller particles which have critical supersaturations about the same as the maximum supersaturation in the TGDC are not nucleated until the chamber has reached its steady state value and then grow more slowly than the larger particles which are nucleated before the chamber has reached equilibrium. The object is to find a period of time when the less active particles have grown to minimum detectable size and the larger (more active) particles have not yet started to fall out. A time which satisfies both criteria may not exist for all size distributions. If such a period of time does exist, then it should evidence itself by a plateau in the curve of number detected versus time. A well-defined plateau was not found to exist in the data on natural continental aerosols presented by Hoppel and Wojciechowski (1976). Alofs and Carstens (1976) did a numerical simulation of the TGDC which predicted large errors depending upon the nuclei distribution and minimum detectable size.

Another source of uncertainty in the TGDC is statistical in nature and has to do with the fact that the number of particles in a small volume will deviate from the true macroscopic mean with a standard deviation given by the square root of the mean. The typical number of particles in our viewing volume varies from about 20 for nuclei concentrations of 300 cm^{-3} to 200 for nuclei concentrations of 3000 cm^{-3} . At the lower end there is therefore a standard deviation (in a large number of measurements) due to real natural variations of

over 20%. This uncertainty, of course, can be reduced by averaging more measurements from the same macroscopic sample of air.

For a monodisperse nuclei sample the plateau in number versus time should be much more pronounced than in the case of a steep size distribution as is usually the case for continental aerosols. At the Workshop, we had a unique opportunity to sample nearly monodisperse aerosols as well as polydisperse and natural aerosols. The Workshop data, therefore, offered a unique opportunity to look for a plateau in number as a function of time.

Figures 1 through 4 show the results of counting every tenth frame (every third of a second) on the videotape starting shortly after the motion in the chamber subsided. There has been no attempt to synchronize the starting points to the same time after the valve was closed. Therefore, there may be a maximum of a half second offset in the plotted times from one sample to the next. The supersaturation in all four figures was about 0.72%.

Figures 1 and 2 are for monodisperse NaCl and $(\text{NH}_4)_2\text{SO}_4$ with two and three runs, respectively. Both of these figures evidence more of a plateau type behavior than we have seen in those ambient samples which we have previously examined in detail. We assume that the variations from one frame to the next after the maximum is caused by the unequal rates at which particles fall into or out of the sensitive volume. Any difference in the level of the plateau from one sample to the next

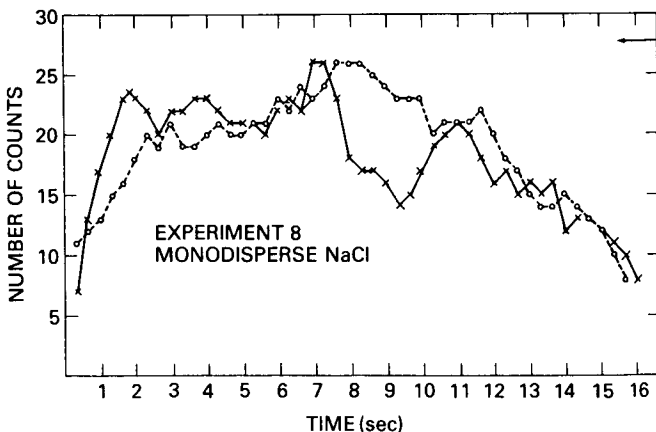


Figure 1

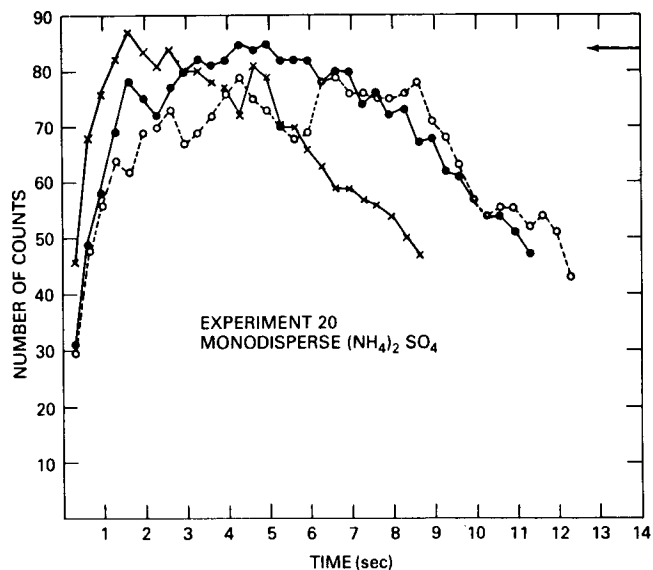


Figure 2

would be due to natural fluctuations discussed above and should have a standard deviation equal to the square root of the number of particles counted.

Figures 3 and 4 are for polydisperse $(\text{NH}_4)_2\text{SO}_4$ and ambient air, respectively. Here the curves are peaked more strongly as would be expected with little evidence of a plateau. The arrows along the right side of the figures indicate the values obtained at the Workshop by the method of analysis indicated earlier.

For many years, NRL has used the standard method cited earlier for determining the CCN concentration from the video recording. This procedure of averaging the maximum count obtained on several successive recordings at the same supersaturation results in concentrations which are somewhat higher than concentrations calculated from an average across the plateau. We have persisted in analyzing our data in this manner for several reasons: (1) the time required to count enough frames to define a plateau (or lack of one) is prohibitive on a routine basis; (2) comparison of results using our photographic system and video system gives values for the video system which are 10 to 15% lower than those obtained with the photographic system, which has a smaller minimum detectable size; (3) most importantly, if there is no plateau then the maximum value should be closest to the correct value; and (4) our results over many years are internally consistent since we have not changed this procedure.

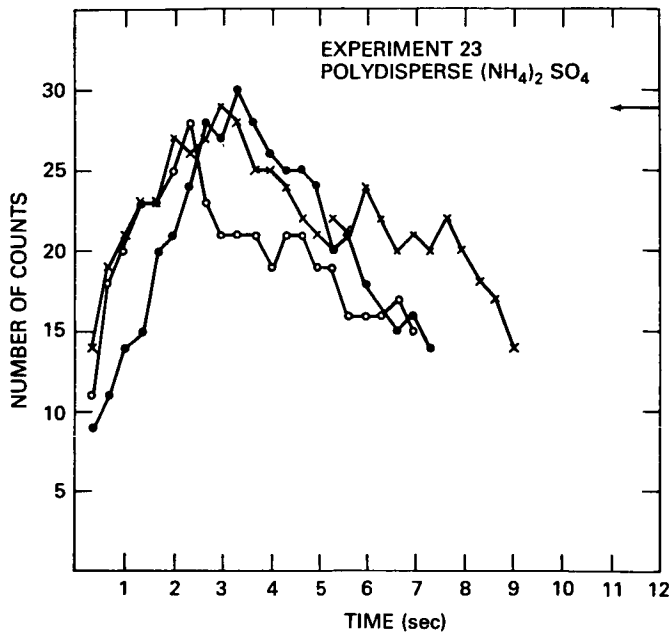


Figure 3

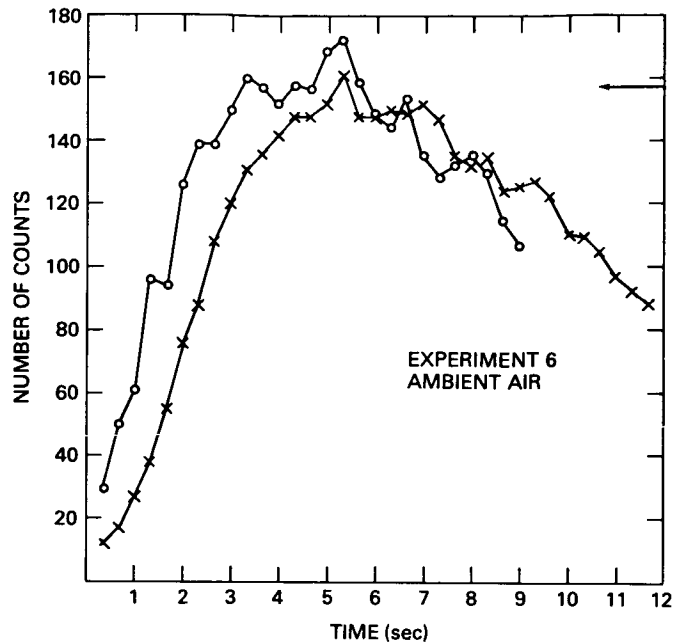


Figure 4

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ACKNOWLEDGEMENTS

This work was partly supported by the Naval Air Systems Command.

THE CCN COUNTER OF THE I.O.P.G. OF PUY DE DOME:
MAIN CHARACTERISTICS AND RESULTS OF MEASUREMENTS

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1. DESCRIPTION OF THE INSTRUMENT

The experimental device used by the I.O.P.G. of Puy de Dome for counting CCN during the Workshop is a cylindrical static thermal diffusion chamber with cylindrical plates, 7.0 cm in diameter and 2.1 in thickness (volume $V \approx 80$ cc). The system includes a multimode laser beam (power > 5 mW), illuminating a part of the median volume of the chamber in which the maximal supersaturation is developing. A video set (TV camera + monitor) displays a motion picture of the droplets generated by the CCN and observed in a volume $V = 7.5 \cdot 10^{-3}$ cc. For obtaining a better signal-to-noise ratio (a better contrast), the TV tube was selected so that its current-wavelength characteristic was in good agreement with the (He, Ne) laser radiation. The TV image is recorded on a video tape recorder, thus allowing analysis shortly after the time of measurement.

Instead of counting by sight, a device called "Image Processing Unit", originally provided for counting in real time, is used. The upper plate is electrically heated and the lower plate is at ambient temperature. Water vapor is supplied by a wet mineral paper filter. The desired temperature difference ΔT (limited at 5°C) between the horizontal plates is displayed and then regulated by an automatic control system, which includes a series of eight differential thermocouples imbedded in the plates. This mounting has the advantage of not only amplifying the voltage but also of taking into account an average temperature on each plate.

The air sample under analysis is introduced in the chamber at a flow rate of 1.0 l/m , and a section of the air intake tube is heated, allowing air sample introduction into the chamber at a temperature of about 3°C above the lower plate tempera-

ture. The supersaturation range covered by this equipment is generally between 0.1 and 1.5% and the time required for obtaining a spectrum in this supersaturation range is approximately 30 to 40 minutes. But in conditions of ambient temperature above 25°C (as occurred during the Workshop), it is not possible to exceed a supersaturation of 0.8% because of the ΔT limitation (see above). Because the device cannot use another greater value of observed sample volume, the accuracy of the measurement is poor at the low extremity of supersaturation range (about 50%).

2. SELECTIVE DATA ANALYSIS

2.1 Global Survey

With such a device, 20 supersaturation spectra (plus one point for Experiment No. 13) corresponding to our complete participation in the Workshop have been outlined and classified according to their shape and position in regard to the series of others established for each experiment.

The following main characteristics of our spectra, appearing in Table 1 with the same capital letter, have been distinguished like that: (A) similar to and well placed among the others in the series; (B) same type of concavity as for the majority in the series; (C) close to the upper (or lower) extremes of the series; (D) smaller CCN concentrations at low supersaturation and/or greater CCN concentrations at higher supersaturation; (E) CCN concentrations abnormally high along the most part of the spectra, with some of them being higher than those obtained with Pollak or TSI counters; (F) lack of plateau; and (G) type of concavity differing from that of the majority.

For each supersaturation spectrum denoted by the number of the experiment, one, two or three, but no more, of these main characteristics have been summarized in Table 1 by a cross in the corresponding square. On Line C, we distinguish respectively by an "s" or "i" index, the fact that the spectrum is close to the upper or lower envelope of the series. In comparing the measurement sheet and the TSI recorded graph corresponding to Experiment No. 16, it can be observed that the abnormal concavity of the spectrum might correspond to a measurement carried out during a peak of aerosol discharge (cross in bracket, Line G).

Through this analysis, Table 1 appears to be divided into two parts: (A, B, C) - grouping the characteristics which reveals a behaviour not far from a "supposed mean" behaviour for a CCN counter; and (D, E, F, G) - grouping the characteristics which reveals an anomaly of behaviour. It is obvious that the percentage of crosses shown in the

first group (A,B,C) is higher than that of the other group, i.e., 62% against 38%.

Characterization	Experiment n°																											
	4	6	8	9	10	11	12	13	14	15	16	17	18	19	20	22	23	24	26	27	28							
A	X																											
B		X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
C			X					X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
D																												
E																												
F																												
G																												

Table 1

Another more realistic analysis consists of dividing the supersaturation spectra in three categories:

1. The crosses pertaining to a given spectrum are shown in the upper part of Table 1. This situation, corresponding to a good behaviour of the device, concerns 9 spectra, i.e., 43% of the whole.

2. The crosses pertaining to a given spectrum are shown in the lower part of Table 1. This situation, corresponding to a frankly bad behaviour of the device, concerns only 3 spectra, i.e., 14% of the whole spectra.

3. The crosses pertaining to a given spectrum are distributed on both parts of Table 1. This situation corresponds to an intermediate behaviour and concerns the remaining spectra, i.e., 43%.

Moreover, it is possible to point out that: (a) the number of spectra for which the crosses are found either on Line C or D is relatively high, i.e., 13 out of 21; and (b) on Line C, the number of crosses affected by the "s" index (i.e., 5) is higher than the number of crosses affected by the "i" index (i.e., 2). Both factors denote a trend in the device to overestimate the CCN concentration, especially in the range of high supersaturation.

2.2 Individual Comparison

Comparisons have been made between our results and spectra obtained using similar equipment (i.e., static diffusion chamber) on corresponding experiments. At times, the spectra were in close agreement, such as: NRL (n° 6-11-12-17); CSIRO (n° 9-10-14-17-28); SUNY (n° 9-14); Hebrew University (n° 9-14-17-18-24). However, the comparison also revealed differences in the spectra, with the concentration ratio reaching 2 to 3 with NRL (n° 9); CSIRO (n° 4-6-8-20); CSU (n° 4-23); SUNY (n° 6-8-9-10) and Hebrew University (n° 10-28). The discrepancy with the Wyoming equipment was still larger (n° 11-12-14-22-23). Surprisingly, for about 62% of the experiments, a number of our spectra were found in reasonable agreement with those resulting from measurements with the DRI continuous flow diffusion chamber (n° 9-12-14-17-18-19-22-27-28) or the Missouri-Rolla haze chamber (N° 6-10-11-12-13-18-19-22-27-28).

2.3 Stability of the Measurements

In Experiment No. 21, which tested the repeatability of the measurements, two runs were carried out continuously during approximately 16 minutes and 30 minutes at supersaturations 0.50% and 0.26%,

respectively. At the same time, the aerosol to be analyzed was delivered at a stable concentration level of 860 cc⁻¹ (measured with the TSI equipment) or 1000 cc⁻¹ (measured with the CCN Pollak counter). After calculating the corresponding standard deviation, the average concentration values were 1210 ± 105 at 0.50% supersaturation for the 1st run and 190 ± 70 at 0.26% supersaturation for the 2nd run.

Two features are emerging from these results: (1) a sensible overestimate of the concentration; and (2) a standard deviation which reaches an acceptable percentage (9%) of the concentration values at mean or high supersaturation, while it is not acceptable (37%) at low supersaturation. This is mainly due to the fact that the examined volume cannot be adjusted to discrete supersaturation ranges.

3. CONCLUSIONS

In performing the data analysis of the measurements achieved during the Workshop with our device, a global approach was preferred rather than an individual analysis, in order to illustrate some main characteristics in the behaviour of the device with respect to a "mean behaviour" resulting from a general survey of all the equipments involved in each experiment. In this regard, our device seems to have a behaviour not unlike a "mean behaviour", although it tends generally to overestimate the CCN concentrations measured near the high supersaturations and sometimes underestimates the concentrations close to 0.1% or 0.2% of supersaturation. Despite the fact that it belongs to a type of static diffusion chamber, it shows, however, similar spectra to those obtained with other types of chambers (continuous flow diffusion chamber and haze chamber).

In the spring of 1980, at the site selected by WMO for a possible Precipitation Enhancement Project in the area of Valladolid, Spain, the CSIRO device and our static thermal diffusion chamber were placed side by side for the purpose of analyzing the same natural air sampling. Although at that time, the air intake of our chamber was not yet equipped with a heater, the main difference between the two devices was the way in which the ΔT between the plates was achieved. In fact, in the CSIRO device the lower horizontal plate is cooled. The comparison of the results displayed CCN concentrations from twice to three times higher with our device, so that our CCN measurements on the P.E.P. site in 1979 and 1980 were questioned.*

Such discrepancies between these devices were again found in some Workshop experiments; however, it was possible to observe that, in a number of other experiments at the least equivalent, the corresponding spectra were rather close to each other (see Section 2.2). Neither the difference in the methods used to obtain ΔT nor the differences in the geometry of the chambers suffice to explain such variable results. Nevertheless, it is obvious that the comparisons made during meetings of a workshop type are able to greatly improve our knowledge of the behaviour of the device involved. As

* Personal communication of Dr. Warren King from CSIRO.

for our equipment, it seems that the best way to try to reduce its trend to overestimate the CCN concentrations should consist in cooling the lower plate instead of heating the upper one and in searching for a better diameter-to-depth ratio in order to improve the stability inside the chamber.

4. ACKNOWLEDGEMENTS

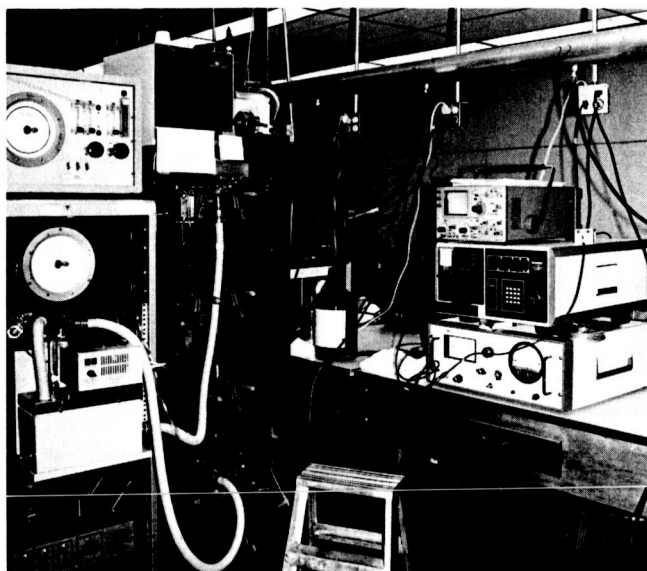
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We are grateful to F. Romeuf from C.R.A. of Lannemezan, for his help in the construction of our static diffusion chamber, and to G. Michel and M. Rahon for having built all the complementary accessories. The Image Processing Unit was designed and built and then calibrated respectively by Miss M. Martin and J.E. Ourtau. The contribution of R. Cordesses and V. Kromer to the maintenance of this unit was also appreciated. Much of the preparation of our journey and that of the present paper was in the capable hands of Mrs. P. Boudinhon.

UMR DUAL MODE CCN COUNTER
(MODES: CFD PLUS HAZE)

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1. INSTRUMENT DESCRIPTION

This instrument and its performance characteristics have recently been described in detail (Alofs, 1978; Alofs, *et al.*, 1979). Therefore only a brief description of it is given here.

The chamber consists of two vertical plates 100 cm long in the vertical direction and 13 cm wide, with a 0.8 cm spacing between the plates. The sample flows downward between the plates in a small diameter stream surrounded by filtered air. The sample flow into the chamber is usually 0.008 liters/min, determined by the pressure drop through a capillary tubing 0.25 mm diameter. This flow branches off of a larger delivery flow of 5 liters/min, to avoid diffusional losses.

The droplets formed on the CCN are counted and sized using a Climet Model 201 optical particle counter (OPC) with an 8-channel pulse height analyzer (Climet Model 210). An advantage of the Climet over the Royco 225 is that the response curves increase monotonically with size (no dips to produce multiple sizes for a given pulse height). The Cooke and Kerker (1978) calculations are used to correct for index of refraction. A sheath air inlet was constructed for the Climet OPC, similar in design to that used in the Royco 225. The rate of flow into the Climet OPC is 0.35 to 1.0 liters/min.

The haze mode is used for the nuclei active at supersaturations (S) from 0.0133 to 0.173%. In this mode the two plates are kept at the same

temperature (25°C). The values of S are determined from the drop sizes, using the relation $S = .08/d$, with S in % and with the drop diameter d in μm . For nuclei with $.068 < S < .173$, the total chamber flow (sample plus filtered air) is 1 liter/min, giving a residence time of 39 sec. For lower S , the chamber flow is 0.35 liter/min and the time is 110 sec. In either case, all of the flow is drawn into the OPC.

In the CFD mode the supersaturation is determined by the temperature difference between the two plates. This temperature difference is controlled by water baths, and is measured with a mercury-glass thermometer immersed sequentially in each water bath. Separate experiments show that the temperature difference at the plate surfaces (measured with thermocouples on the air side of the filter paper) equals the bath temperature difference to within 5%.

About 40 minutes is required to obtain a spectrum (5 values of S in the CFD mode, plus 7 values of S in the haze mode). About half of this time is used to adjust the temperatures and flows; the rest is used in actual counting of the nuclei. Generally the measurement begins in the CFD mode, with the hot plate at 25°C and the cold plate at 20°C. The cold plate temperature is raised in steps (the baths heat very quickly, 1000 watts of heating versus about 50 watts of cooling). The chamber flow rate is usually reduced as S is decreased. The time for flushing the chamber is quite short in the CFD mode, but amounts to about 5 minutes in the haze mode at 0.35 liters/min.

2. PERFORMANCE AT THE WORKSHOP

The analysis that follows was performed based on the computer printouts of the data, as supplied to us in February, 1981.

Consider first the monodisperse sodium chloride and ammonium sulfate experiments. Let S_{CU} denote the critical supersaturation determined from the size given by U. Katz. On a plot of CCN count versus S , these aerosols show a plateau. At a count of 50% of the plateau value, let the corresponding S be defined as the measured critical supersaturation, S_C . Define the parameter X by $X = (S_C - S_{CU})/S_{CU}$. Table 1 shows the average (\bar{X}) and standard deviation (σ) of X for eight instruments that we judged to be giving above average performance. There are two groups of experiments, with $S_{CU} > 0.1$ shown on the top 6 rows, and $S_{CU} < 0.1$ shown on the bottom 4 rows. In each group the instruments are arranged according to σ , lowest σ on top.

In the top group of Table 1, it can be seen that our instrument gave $\bar{X} = +0.0296$ and $\sigma = 0.107$

INSTRUMENT	EXPERIMENTS	S _{CU} RANGE	X = (S _C - S _{CU}) / S _{CU}	
			AVERAGE, \bar{X}	STD. DEV., σ
#18, DRI-CFD Hudson-Squires type	4,5,10,15, 18,20	>0.1	-0.0565	0.103
#21, UMR CFD mode	4,5,10,15, 18,20	>0.1	+0.0296	0.107
#17, NRL Twomey type	4,5,10,18	>0.1	-0.171	0.125
#12, NRL elect. class.	4,5,10,15, 18,20	>0.1	+0.0169	0.156
#15, DRI-CFD NASA	10,15,18, 20	>0.1	+0.0067	0.158
#10, SUNY static, scattering	5,10,18	>0.1	-0.0019	0.315

#21, UMR haze mode	8,9,10,19	<0.1	+0.109	0.0453
#12, NRL, elect. class.	8,9,10,19	<0.1	+0.0932	0.0897
#14, DRI haze	8,9,10,19	<0.1	+0.221	0.190
#11, NRL haze	8,9,10,19	<0.1	+0.184	0.321

TABLE 1. Critical Supersaturation (S_C) of Monodisperse Aerosols. S_{CU} = Value from U. Katz Electrical Classifier.

which is about the same performance as we have reported in the literature (Alfons, *et al.*, 1979). Except for instruments #17 ($\bar{X} = -0.171$) and #10 ($\sigma = 0.315$), the other instruments (#18, #12, #15) gave performance of similar quality, with $|\bar{X}| < 0.06$ and $\sigma < 0.16$.

Now consider the bottom 4 rows of Table 1; that is, experiments with S_C < 0.1. A very good performance is indicated by our instrument in the haze mode ($\bar{X} = +0.109$, $\sigma = 0.0453$). The NRL electrical classifier showed somewhat higher σ (0.089) and the haze chambers of DRI and NRL showed considerably higher σ . These instruments (#14 and #11) both gave $\bar{X} \approx 0.4$ for experiments #8 and #9, and $|\bar{X}| < .05$ for experiments #10 and #19. These instruments also both use Royco Model 225 optical counters, which have a multiple value response function for water drops in the range 1 to 2 μm diameter (Cooke and Kerker, 1975). Experiments #8 and #9 give haze drops of about 2 μm diameter, which is within the ambiguous size range of the Royco and probably explains the decrease in sizing accuracy.

Table 2 shows another type of comparison. The concentration ratio (R = ours \div other) was computed for S = 0.03, 0.1, 0.3, and 1.0. The average (\bar{R}) and standard deviation (σ) of R were then computed over the available set of experiments for each instrument.

In Table 2, consider first the Hudson-Squires type CFD of DRI. At S = 0.3, $\bar{R} = 1.04$ and at S = 1, $\bar{R} = 1.18$. Thus our counts were 4-18% higher than theirs. This is pretty good agreement in our opinion. However, $\sigma/\bar{R} = 0.275$ at S = 0.3, which is definitely higher than we expected in view of the 1% agreement that Hudson and Squires (1976) obtained with a pair of their CFD's. At S = 1, the value of σ/\bar{R} is considerably lower (0.132), and is the lowest value of σ/\bar{R} in Table 2.

In the haze mode, our instrument compares well with the NRL haze chamber ($\bar{R} = .974$ at S = .03, $\bar{R} = 1.067$ at S = .1) but gives considerably higher concentration than the DRI haze chamber ($\bar{R} = 2.2$) and lower concentration than the NRL electrical classifier. The standard deviations at S = .03 and S = 0.1 are higher than in Table 1, but are still not too bad considering the problems involved in sizing water drops with optical counters.

OTHER INSTRUMENT	SUPERSATURATION (%) EXPERIMENTS	Average Ratio, \bar{R} , Univ. of Mo. (#21) \div Other				Relative Std. Deviation σ/\bar{R}			
		.03	.1	.3	1.	.03	.1	.3	1.
#18, DRI-CFD Hudson-Squires type	5,8,9,10,11,12,13, 14,15,16,18,19,20, 22,23,26,27,28	--	--	1.04	1.18	--	--	0.275	0.132
#17, NRL Twomey type	1,2,3,4,5,8,9,10, 11,12,14,15,18,19, 20,22,23,26,27	--	--	0.943	0.937	--	--	0.258	0.215
#10, SUNY static, scattering	5,8,9,10,12,13,14, 15,18,19,20,22,23, 26,27,28	--	--	0.885	1.27	--	--	0.221	0.200
#12, NRL elect. class.	1,2,4,5,8,9,10,13, 14,15,18,19,20,22, 23,24	0.678	0.632	0.899	0.915	0.302	0.418	0.284	0.209
#11, NRL, haze	1,2,3,6,8,9,10,11, 12,14,15,19,20,22, 23,24,26	0.974	1.067	--	--	0.600	0.382	--	--
#14, DRI haze	8,9,10,11,12,14,15, 16,17,19,20,22,23, 24,26	2.17	2.18	--	--	0.722	0.333	--	--

TABLE 2. Ratio, University of Missouri \div Other Instrument

3. ACKNOWLEDGEMENTS

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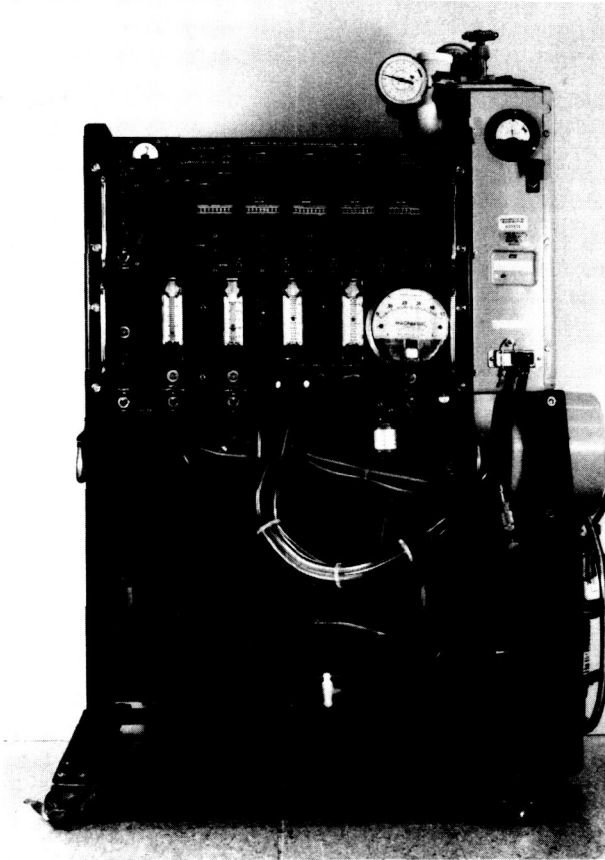
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A CLOUD CONDENSATION NUCLEUS SPECTROMETER
DESIGNED FOR AIRBORNE MEASUREMENTS

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Abstract

A portable, vertical plate cloud condensation nucleus spectrometer, suitable for mounting aboard a small aircraft, is described. This instrument, which incorporates several unique design features, is automated and can provide rapid, simultaneous measurements of the concentrations of cloud condensation nuclei at four supersaturations.

1. INTRODUCTION

In 1970, at the Second International Workshop on Condensation and Ice Nuclei, all but one of the cloud condensation nucleus (CCN) counters were horizontal static-diffusion chambers operating in a manner little changed from those used in the 1950's [1,2]. Only one of the static chambers was fully automated [3]. All of the counters had significant limitations in time resolution and all required an operating time of several minutes or more to measure the supersaturation spectrum.

Recently, variations on the continuous flow CCN counters developed in the 1960's [4,5,6] have produced a number of automated and semi-automated CCN counters which use single particle optical counters as detectors. These new devices offer the potential for both significantly improved accuracy and time resolution in CCN measurements, although most have provided only limited improvements in quickly measuring the CCN activity spectrum.

The CCN spectrometer to be described in this paper is a simple extension of the continuous flow diffusion chamber (CFDC) operated in the vertical mode, much like the instruments described in [7], [8] and [9]. The principal new design features of our instrument are its rapid time response and ability to rapidly measure the CCN activity spectrum. Also, the instrument is small enough to be mounted and operated in a research aircraft of modest size.

2. PHYSICAL LAYOUT AND AIRFLOW

The activity spectrum of CCN in an air sample is measured rapidly by simultaneously operating the spectrometer at four supersaturations. The CCN spectrometer consists of four essentially independent CFDC's in a parallel array (Fig. 1). Each CFDC is 61 cm long, 10 cm wide, and consists of two vertical parallel plates 1 cm apart, maintained at different temperatures. Each CFDC has a single particle optical counter located at the center of its base.

The instrument operates in the following sequential manner:

i) An air sample is drawn through the duct at the top of the CFDC's.

ii) A fraction of the airflow exits the sample duct through a symmetrical, streamlined slit at the bottom of the duct. The rest of the sample flow is discarded.

iii) As the air enters the top of one of the CFDC's, it is sheathed with two temperature-conditioned, particle-free "curtains" of air (called the "sheath flows").

iv) The three airstreams travel in laminar flow, with negligible intermixing, down each of the CFDC's where, after 8 cm of travel, they encounter a saturated felt on the cold plate. At a further distance downstream, the warm plate is also covered with wet felt. This results in the airstream becoming supersaturated and CCN are activated to form droplets. The total length of felt on the

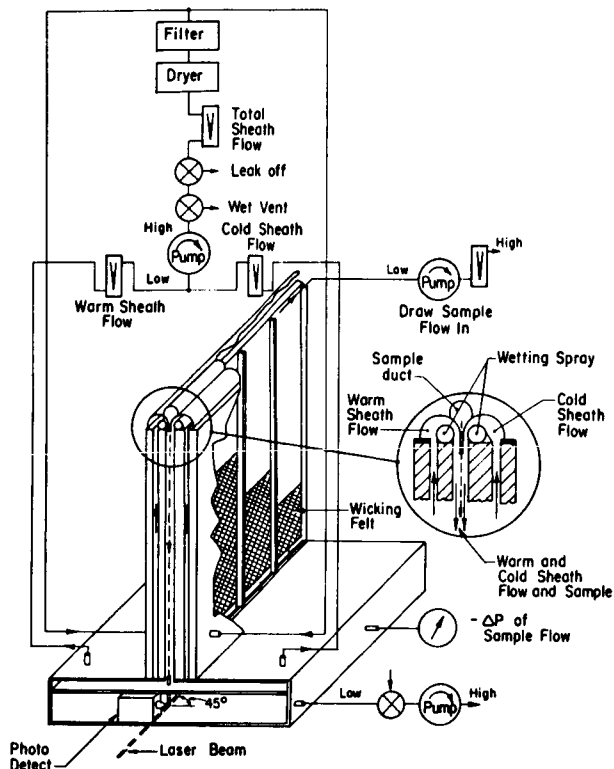


Figure 1. Schematic showing the airflow and optics of the University of Washington's CCN Spectrometer.

warm plate is roughly inversely proportional to the supersaturation in the chamber.

v) As the airstreams leave each of the CFDC's they are split along the vertical, central plane of the sample air, with half of the total flow being removed on each side at the base of the CFDC. A small fraction of the airflow is removed as the stream is divided by a small tube which extends up into the airflow. The droplets in this fraction of the airflow are subsequently counted by optical counters (see §4).

vi) The position of the airstream with respect to the inlet to the optical counters can be precisely adjusted by changing the flow rates in the sheath flows; this allows the airflow to be exactly centered on the inlet to the optical counters.

vii) The two halves of the total airflow are then filtered, dried and reinserted on the back sides of the warm and cold plates. These two airstreams travel up the plates and are in thermal equilibrium with the plates; when they arrive at the top of the CFDC they form the warm and cold sheath flows.

3. SUPERSATURATION CONTROL

A novel method is used for providing four distinct temperature differences between the warm and cold plates of the four CFDC's. As shown in Fig. 2, a large aluminum plate is cooled nearly uniformly by a small mechanical refrigerator. The warm plate, which is made of copper, is connected to the cold plate by a brass end plate and at the

opposite end it is heated by the hot refrigerator gases. The copper plate is vertically segmented between each CFDC with a thin insulator. The four copper segments stabilize at four distinct, and nearly uniform, temperatures. From measurements of these temperatures, and the temperature of the cold plate, the supersaturation in each of the CFDC's is computed by a microprocessor. Note that the supersaturations in all four CFDC's are adjusted by a single control. The four supersaturations normally achieved are approximately 0.2%, 0.5%, 1.0% and 1.5%*.

4. OPTICAL DETECTION AND DATA PROCESSING

The CCN activated in the CFDC's are detected as droplets in an optical box (Fig. 2). The optical box contains a three mW He-Ne laser and four identical photo-detectors placed beneath each of the sample outlet tubes from the four CFDC's. The streams of droplets pass through the laser beam and are viewed by the photo-detectors at a forward scattering angle of $45^\circ \pm 5^\circ$. The photometers readily detect all droplets greater than $0.5 \mu\text{m}$ in diameter. The available growth times in the CFDC's (5-20s) are more than adequate to grow the activated CCN into droplets that are much larger than any unactivated haze droplets. Thus, by appropriate sensitivity adjustments, activated droplets are detected but the unactivated haze droplets are not.

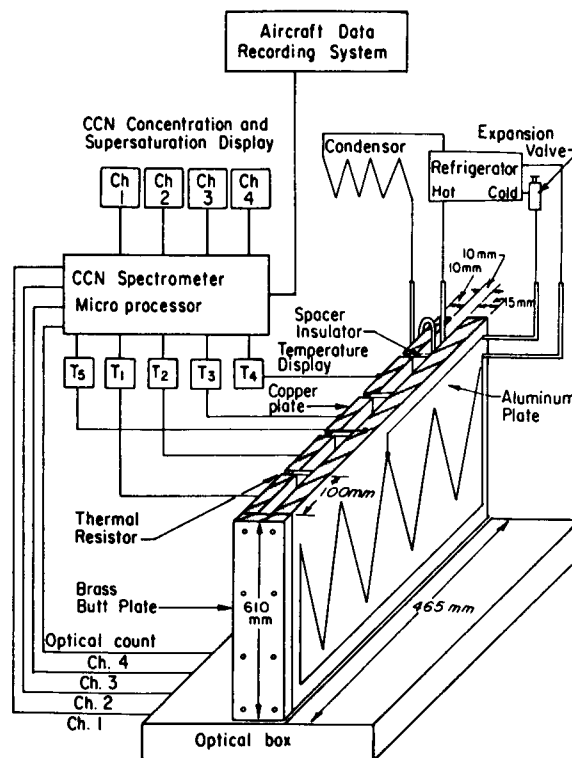


Figure 2. Schematic of thermal and electrical components of the CCN Spectrometer.

* During the CCN Workshop the range of supersaturations achieved was limited by a malfunction of the refrigerator unit.

A microprocessor is programmed to process the data in several ways. It can display the supersaturation and CCN concentrations measured in each of the four CFDC's on command from the aircraft's central computer. Alternatively, the microprocessor can delay output and accumulate data until certain statistical counting criteria are met.

5. CONCLUSIONS

We have designed and built a compact and reasonably simple CCN spectrometer for airborne use. In its first full-scale testing, at the Third International Measurement Workshop on CCN, it compared well with both conventional CCN counters and large continuous flow CCN counters designed for use on the ground. This instrument is currently undergoing final laboratory testing and modification before flight testing.

6. ACKNOWLEDGEMENTS

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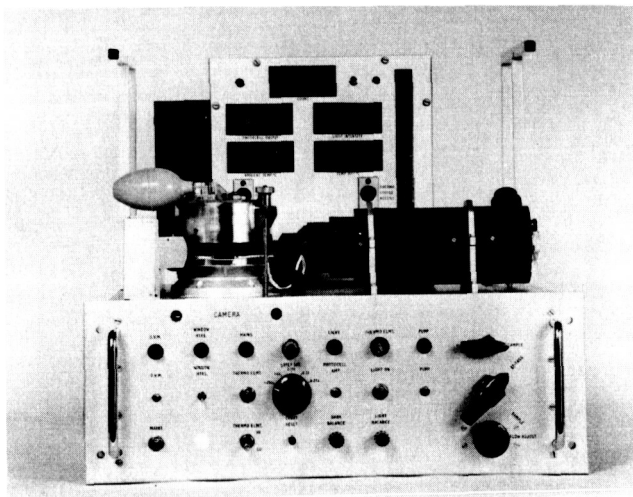
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STATIC DIFFUSION CLOUD CHAMBER

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1. CHAMBER GEOMETRY

The thermal diffusion chamber is based upon an earlier design of Twomey and Davidson, the cylindrical glass walls of which have been replaced by an annular perspex (plexiglass) spacer 12.5 mm in height, which is fitted with a very thin stainless steel liner that contacts both top and bottom plates so as to linearize the edge temperature gradient. Gagin and Terliuc (1968) used a somewhat thicker liner for this purpose.

The optical arrangement is similar to that used by Lala and Jiusto (1977). In the present design heated windows have been installed at appropriate places to allow a waisting light beam from a projector - lamp/lens combination to pass through the chamber. A window is placed at 45° with respect to the beam to view the forward scattered light with a photocell and lens. A further window at 90° to the beam allows the number of drops in the illuminated cloud to be recorded photographically.

Summary of Chamber Geometry:

Inside diameter	75 mm
Height	12.5 mm
Scattered light at 45° recorded with a peak detector	
Cloud droplets recorded photographically at 90°	
Sample volume	10 mm x 3 mm x 3 mm

2. SAMPLING SYSTEM

Air is drawn continuously through the equipment at about 1 - 1.25 dm³/min by means of a simple aerator (aquarium) pump. The flow bypasses the

chamber until a measurement is required, at which point air is admitted by turning a four-way selector valve to the "sample" position. A two-way selector valve in the air circuit enables the sample air to be passed through an absolute filter so that periodic zero checks can be made.

Sampled air enters the chamber from six peripheral holes in the underside of the top plate after passing twice around the edge in a gallery to attain the same temperature as the top plate.

The chamber and associated air lines are sealed from the ambient atmosphere so that, if required, the equipment may be operated at pressures below ambient. A prototype of this design has been successfully operated in the Division's pressurized aircraft.

2.1 Supersaturation Range

Readings are taken at five fixed points: 0.25%, 0.5%, 0.75%, 1.0% and 1.25%.

2.2 Detection System

Light source: Sylvania EJV Projector Lamp

150W - 21V operated at approximately 19V, illuminating at a distance of 45 mm a 2.4 mm square of ground glass screen. Light from this screen is passed through infrared absorbing glass and focused by means of a Bell and Howell projector lens (F1.2; focal length 51 mm) to give a beam waisting down to a 3 mm square section over a 15 mm length at the center of the diffusion chamber. The central 10 mm is used for droplet detection.

Camera: Canon AE-1 fitted with a Vivitar

55 mm Macro F2.8 lens, power-winder and data back. Film used is Tri-x (400 A.S.A.) processed for 600 A.S.A. Photographs are taken at 1/4 sec. and F4, though the effective aperture is much smaller (higher F number) since the camera views the chamber through a small window set in a short tube. Photographs of a mm graticule placed at 45° to the incident light beam and camera window confirmed both the beam cross sectional area and that depth of field was sufficient to encompass the 3 mm width of the beam.

Photocell detector: An EG and G Electro-Optics Silicon Photovoltaic Detector type PV-215 coupled to a two-stage amplifier having an output gain of 100V per micro-amp of cell current. Amplifier output is fed to a peak detector having a digital panelmeter readout. A manual reset is used. The photocell views the scattered light from virtually the same sample volume as the camera, but at an angle of 45°, and through an 8 mm focal length lens. A mask in front of the lens serves to define the viewed sample volume.

3. TEMPERATURE CONTROL

Temperature of the top and bottom plates is measured using semi-conductor transducers (National Semiconductor type LM3911 or similar). The transducers are coupled to amplifiers to give outputs of 100mV per K and have been set to within 1 or 2 hundredths of a K at ice point and at 293.15K. Digital panelmeters read the top plate temperature to 0.1K and the top-bottom temperature difference to 0.01K, though the bottom plate temperature varies by ± 0.04 K due to "hunting" of the controller. Thermocouples have been built into both plates to provide an independent check on the performance of the temperature measurement and control circuitry.

Bottom plate cooling is by means of a Komatsu thermoelectric cooling element, type KSF-2012. Simple circuitry uses the temperature transducer outputs to establish and maintain automatically the required ΔT , regardless of top plate temperature variations.

4. CALIBRATION

Calibration amounts to relating the peak value of scattered light intensity to the number of droplets in the sample volume at the time at which the peak occurred. In the present case, a chart recorder was used to register, simultaneously, variation in scattered light intensity and the time at which photographs of the scattering volume were taken (usually a series of 5-7 frames at 1 per second). It was then a simple matter to determine which frame corresponded most closely in time to the peak in the scattered light intensity. In this way, calibration curves were constructed for each supersaturation in the form of plots relating "film count (cm^{-3})" to "peak reading".

The simple analysis of a thermal gradient chamber discussed by Lala and Jiusto (1977) suggests the peak value in scattered light intensity is dependent on supersaturation and droplet concentration only and, in particular, that the number of droplets present in the sample volume at the time the peak is reached is only two-thirds of the initial nucleus concentration, independent of supersaturation. Thus our final estimate of CCN concentration is made by multiplying the "film count" by a factor of 1.5.

Calibration curves used at the workshop were derived from room air samples by blending varying amounts of filtered and unfiltered air to obtain a range of peak readings.

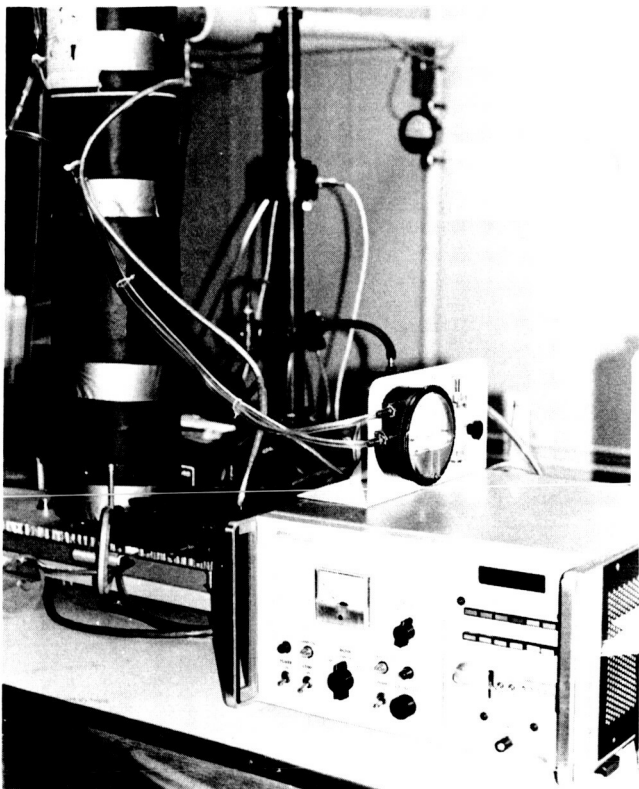
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AN AIRBORNE ISOTHERMAL HAZE CHAMBER

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1. INTRODUCTION

Thermal gradient diffusion cloud chambers (TGDC) are used to determine the concentrations of cloud condensation nuclei (CCN) with critical supersaturations (S_C) greater than or equal to about 0.2% (Hoppel and Wojciechowski, 1976). The concentrations of CCN active at $S_C \lesssim 0.2\%$ can be determined using an isothermal haze chamber (IHC) following the work of Alofs (1978), Hoppel (1979) and Hudson (1980). They have shown that CCN spectra over a supersaturation range $0.01 \leq S_C \leq 1.0\%$ can be obtained from simultaneous measurements with a TGDC and an IHC. Their instruments are designed to operate either in the laboratory or in the field.

The IHC discussed in this paper and operated at the CCN Workshop was designed for use in a light aircraft. It is based on the IHC design of Hudson (1980). The objectives of our participation in the Workshop were to (1) compare the response of the airborne IHC to the theoretical response (based on accurate sizes, numbers and compositions of aerosol particles), (2) compare the response of the IHC to the response of the larger laboratory IHC's, and (3) compare the response of the IHC to the response

of the CSU-Mee TGDC (Borys, 1980) in the region of overlapping supersaturations. The Mee instrument resembles that constructed by Lala and Jiusto (1977).

2. DESCRIPTION OF THE CHAMBER

Hudson (1980) provides a schematic of the IHC and the operating principles. Here we will provide details of the airborne version of Hudson's chamber which we have constructed. The line drawings for the chamber are given in Figure 1. The air flow and water flow through the chamber are given in Figure 2.

The supersaturation range of the instrument is determined by the size range of the solution droplets (in equilibrium at 100.0% RH) which the ROYCO sensor can detect. The equilibrium droplet size is related to its critical supersaturation as follows:

$$r_{100} = 4.1 \times 10^{-6} / S_C \quad (1)$$

where r is in centimeters and S_C is in percent, after Laktionov (1972), Hoppel and Fitzgerald (1977), Alofs (1978) and Hudson (1980). The ROYCO sensor latex-sphere calibration was revised for water droplets by Fitzgerald (1980, private communication) following Cooke and Kerker (1975); see Figure 3. The sensor was adjusted such that the critical supersaturation thresholds corresponding to the droplet sizes from (1) were 0.15%, 0.11%, 0.041%, 0.027% and 0.016%, respectively.

The ROYCO Model 225 optical particle counter which is described by Liu, *et al.* (1974) was used to size and count the solution drops produced in the IHC. The instrument was operated at a 60 s sampling interval. Consequently, it took 1 minute for a CCN spectra to be obtained.

The maximum flow F to permit droplet equilibrium in the chamber was estimated following a procedure outlined by Hudson (1980). The results of the variations in particle concentrations as a function of F were similar to the results reported by Hudson. We found the maximum value of F should be $\sim 35 \text{ cm}^3 \text{ s}^{-1}$. Values of F greater than this value would provide insufficient time for droplets to reach equilibrium size before passing through the ROYCO sensor.

The volume sampled for a CCN spectra corresponds to the flow rate f through the sensor multiplied by the sample period: $\sim 1 \text{ cm}^3 \text{ s}^{-1} \times 60 \text{ s} = 60 \text{ cm}^3$. This result is valid because the main flow F is essentially particle free.

There is no cooling or heating requirements for the chamber; the chamber is isothermal.

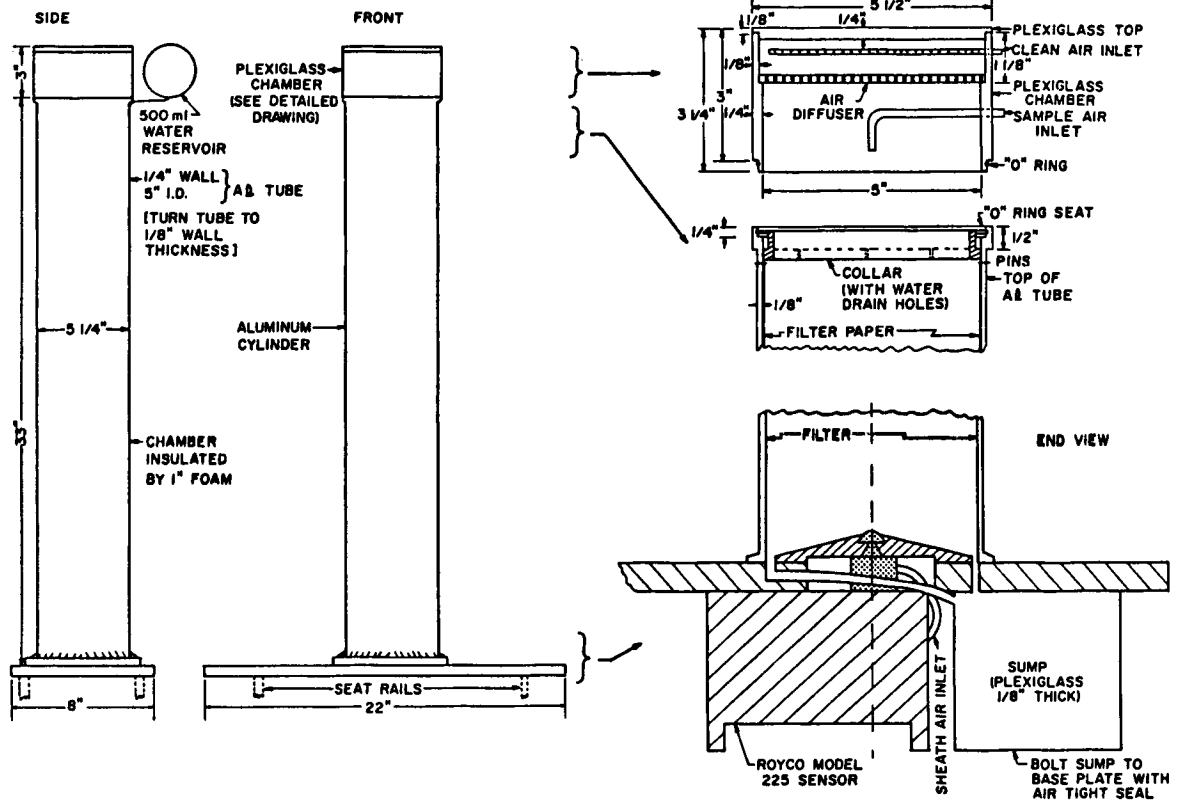


Figure 1. Isothermal haze chamber and main components: Aluminum cylinder with inside walls covered with Whatman 41 cellulose high-volume filter paper; plexiglass top into which flows particle-free air, sample air and water by gravity feed from the 500 ml reservoir; the ROYCO Model 225 sensor which sizes and counts the solution droplets which form in the chamber; the sump for receiving the water which drains from the filter paper.

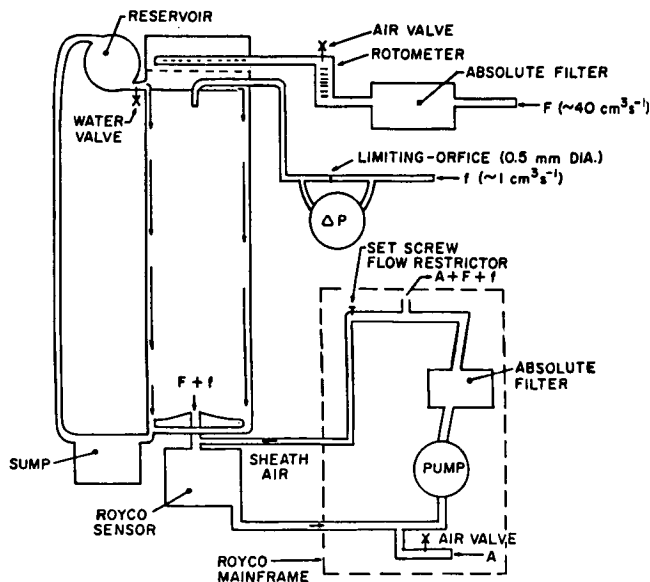


Figure 2. Airflow and water flow schematic.

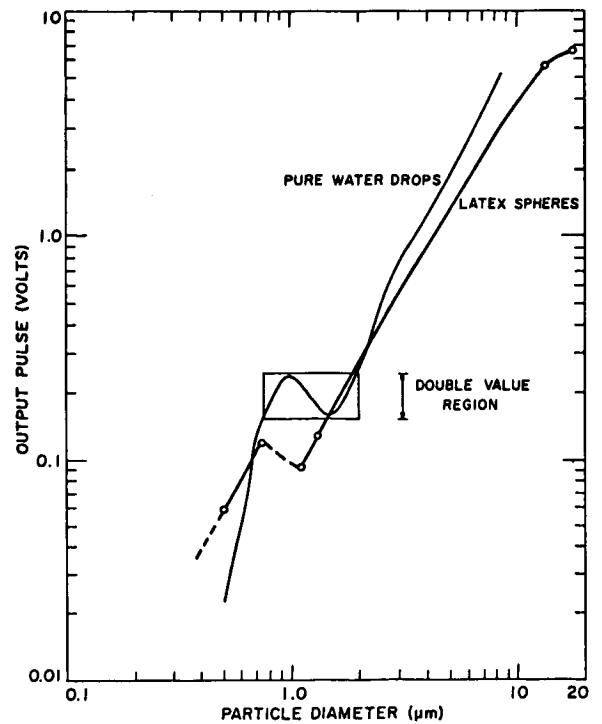


Figure 3. Calibration of ROYCO sensor for latex spheres. The calibration for pure water drops was from Fitzgerald (1980, private communication).

The minimum detectable drop size is 0.62 μm diameter (see Figure 3). This is the minimum size setting before electronic noise interferes.

3. RESULTS

During Experiments 1 through 13, the IHC was operated in the same fashion ($f = 1.3 \text{ cm}^3\text{s}^{-1}$, $F = 35 \text{ cm}^3\text{s}^{-1}$). It was clear from these early experiments that the concentrations of CCN detected by the airborne IHC were significantly below the concentrations detected by the laboratory IHC's (see the Table). For example, in the Table, the airborne IHC detected an average of 94 times fewer CCN active at $S_c \leq 0.05\%$ than did the laboratory IHC's.

One reason for the CCN concentration differences between the laboratory and airborne IHC's was thought to be a malfunctioning CSU ROYCO 225 sensor. The sensor was checked with the York University ROYCO 225 sensor and no significant difference was found between the dry particle size distributions measured by the two instruments. Further, during Experiments 14 and 15, the spare NRL ROYCO 225 sensor was connected to the CSU IHC and the CCN concentrations continued to be too low. Consequently, the CSU ROYCO 225 sensor appeared to be operating normally.

Another reason for the CCN concentration differences was thought to be a subsaturated chamber. Consequently, a humidifier (courtesy of Dr. Fred Rogers) was placed in the main-flow air line downstream of the filters during Experiment 17. The humidifier remained in the line from Experiment 18

to the end of the Workshop. The results in the Table show that the humidification reduced the differences in concentrations between the lab IHC's and the airborne IHC. For example, the factor of 94 difference mentioned earlier reduced to a factor of 7.5 following humidification. The Table presents a comparison of the results from theory and from the airborne IHC. Following humidification, the airborne IHC measured 9.0 times fewer CCN active at 0.05% and 7.9 times fewer CCN active at 0.14% than predicted to occur by theory.

Figure 4 illustrates the results obtained from the simultaneously operating CSU IHC and CSU TGDC. Also plotted are the results obtained using measured particle sizes and the theoretical relationship between the dry $(\text{NH}_4)_2\text{SO}_4$ particle size and its critical supersaturation from Fitzgerald (1973):

$$r_d = 1.53 \times 10^{-6} S_c^{-2/3} \quad (2)$$

where r_d is in centimeters and S_c is in percent. It can be seen from Figure 4 that the results from the two instruments tracked the theoretical response but with concentrations significantly less than the theoretical response. Further, the supersaturation ranges of the two instruments did not overlap because the lowest value of the TGDC range was 0.2% as established by Borys (1980) and the smallest droplet detected by the ROYCO was 0.62 μm diameter which is equivalent to a S_c of 0.13%. Nevertheless, it can be seen the slopes of the curves in this region approximate the theoretical slope.

TABLE
Selected CCN Concentration Measurements for $S_c \leq 0.05$ and $\leq 0.14\%$

Humidity?	Experiment	Theory*	NRL	UMR	DRI	CSU	Theory/CSU	Lab IHC's**/CSU	
$N(S_c \leq 0.05\%) \text{ cm}^{-3}$	No Humidity	300	120	100	80	1.5	200	67	
		600	270	250	200	5.0	120	48	
		66	30	17	8	0.09	733	203	
		-	90	26	-	1.0	-	58	
							351	94	
	Humidity	19	110	-	65	25	4.0	28	11
		20	0.4	2	2	2	0.1	4	20
		22	5	1	1	1	1.4	3.6	0.7
		23	1	0.5	1	0.5	0.15	6.7	4.4
	24	10	5	7	4	4	2.5	1.3	
							9.0	7.5	
$N(S_c \leq 0.14\%) \text{ cm}^{-3}$	No Humidity	420	290	300	240	120	3.5	2.3	
		800	600	850	350	300	2.7	2.0	
		270	240	210	100	30	9.0	6.1	
		-	350	460	-	70	-	5.8	
							4.1	4.1	
	No Humidity	19	1000	-	1000	350	250	4.0	2.7
		20	140	190	140	58	30	4.7	4.3
		22	150	100	100	30	17	8.8	4.5
		23	28	17	13	5	2	14	5.1
	24	400	170	230	70	50	8.0	3.1	
							7.9	3.9	

* The dry-particle sizes and numbers were known and their S_c could be calculated using theoretical relationships developed by Fitzgerald (1973).

** The Lab IHC value is the average of the NRL, UMR and DRI measurements.

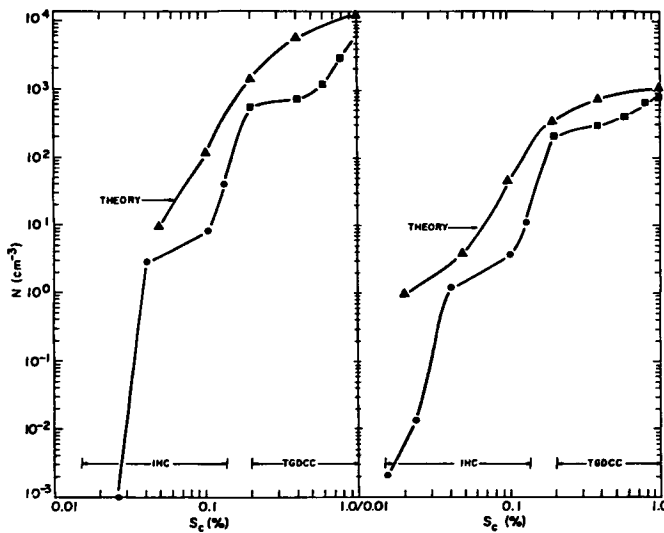


Figure 4. Results from the CSU IHC and the CSU TGDC for Experiment No. 24 (left) and Experiment No. 22 (right). The particles in both cases were polydisperse $(\text{NH}_4)_2\text{SO}_4$.

4. DISCUSSION

How subsaturated must the airborne IHC have been to account for the differences between the theoretical results and the measurements? From Figure 4 it can be seen that if the S_c values were shifted from 0.1% to about 0.05% then the theoretical results and measurements would be identical. The r_{100} corresponding to these two S_c values are 0.41 μm and 0.82 μm , respectively, using (1). Consequently, we may have been measuring the concentrations of droplets with radii $\geq 0.41 \mu\text{m}$ at an unknown RH. The unknown RH can be estimated from a graphical representation (Figure 5) of the relationship between S_c , $r_{\text{equilibrium}}$ and RH ($\leq 100.0\%$) provided by Hoppel (1980, private communication). As can be seen from Figure 5, droplets $\geq 0.82 \mu\text{m}$ at 100.0% RH become droplets $\geq 0.41 \mu\text{m}$ at 99.1% RH. So, if the chamber achieved only 99.1% RH, this subsaturation would explain all of the differences between theory and measurements.

How long a growth time is required for the largest solution drop measured ($r_{100} = 2.5 \mu\text{m}$) to closely approximate its equilibrium size? Hoppel (1980, private communication) has calculated the growth times to reach the critical radius (r_c); r_c is greater than r_{100} . The times are shown in Figure 6. Interpolating between his worst case values (condensation coef. = 0.036) the growth time to reach 2.5 μm is about 280 s. The growth time available in the chamber is about 307 s ($35 \text{ cm}^3\text{s}^{-1} / 126 \text{ cm}^2 \times 83 \text{ cm}$). Consequently, sufficient time

exists for the largest solution drops to reach equilibrium. Insufficient growth times cannot explain the differences between theory and the measurements.

Could the water droplet calibration for the ROYCO sensor account for the differences between the theoretical results and the measurements? To do this, the droplets with diameters indicated to be 0.62, 0.82 and 2.0 μm , according to the calibration in Figure 3, would have had to indicate 1.13, 1.64 and 3.56 μm , respectively. These latter values are a dramatic departure from the calibration. The calibration works adequately with the NRL ROYCO 225 which is connected to their IHC. Consequently, it is believed the sensor calibration cannot account for all of the differences between theory and measurements.

The only particle losses in the system occur across the limiting-orifice (Figure 2). The losses were measured to be at most a factor of 1.3. This loss cannot account for much of the factors of 3.9 and 7.5 reported in the Table.

What is the possibility that droplets entering the ROYCO sensor at equilibrium size (RH = 100%) shrink in size by the time they pass through the optics? It can be seen from Figure 1 that the sensor inlet protrudes into the IHC. Therefore, the time for a particle to travel from the inlet to the optics is approximately 1.1 ms ($7.9 \times 10^{-3} \times 5.1 \text{ cm} / 37 \text{ cm}^3\text{s}^{-1}$). Since the sheath air entering the sensor was slightly warmer and dryer than air entering the sensor from the IHC, conditions

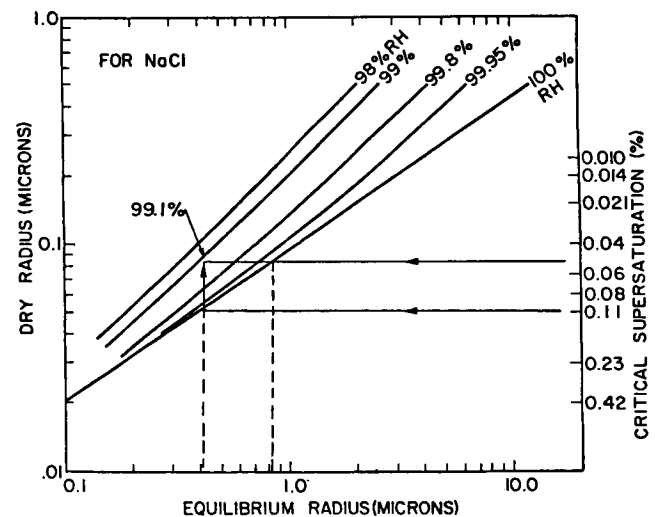


Figure 5. Relationship between r_{dry} , $r_{\text{equilibrium}}$, RH ($\leq 100.0\%$) and S_c for NaCl particles (close approximation to $(\text{NH}_4)_2\text{SO}_4$ particles). From Hoppel (1980, private communication).

for droplet evaporation were present. Alofs (1978) calculated effects of droplet shrinkage on S_C for different droplet residence times. Applying our 1.1 ms time to his worst case situation (95% RH in the optics) it was found that the effect would be confined to $S_C \geq 0.1\%$ and at most would change the inferred S_C of 0.16% to about 0.15%. Consequently, droplets will not shrink significantly due to the extremely short residence time.

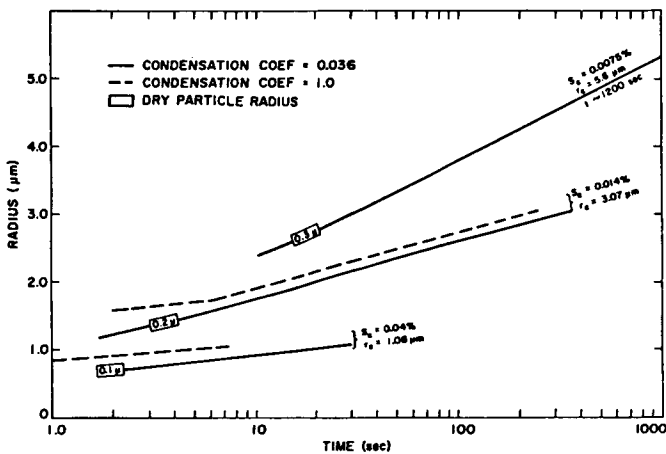


Figure 6. Growth times for dry-particles to reach their critical sizes (r_c). From Hoppe1 (1980, private communication).

Finally, errors in the theoretically derived CCN response should be negligible. Gerber, *et al.* (1977) demonstrated experimentally the soundness of the theoretical relationship between r_d and S_C given by (2). The same particle sizing and counting equipment used by Gerber, *et al.*, was used by Hoppe1 during the CCN Workshop to provide the measurements for deriving the CCN concentrations.

5. CONCLUSIONS

The following conclusions can be made based on the results obtained from the CCN Workshop. The CCN concentrations measured with the airborne IHC were lower than theoretically predicted by factors ranging between 7.9 and 9.0. The CCN concentrations measured with the airborne IHC were lower than the concentrations measured with the larger laboratory IHC's by factors ranging between 3.9 and 7.5. The bounds of the supersaturation ranges of the airborne IHC ($S_C \leq 0.16\%$) and the CSU-Mee TGDC

($S_C \geq 0.2\%$) do not overlap. Nevertheless, the slopes of the interpolated data between the bounds agree favorably with the theoretical slopes. Slight subsaturations in the IHC plus uncertainties with the sensor calibration are the most probable causes for the discrepancies between the measured and predicted CCN concentrations.

6. ACKNOWLEDGEMENTS

The Desert Research Institute is gratefully acknowledged for organizing and executing such an excellent and productive Workshop. The author's participation was sponsored by ONR Contract N00014-79-C-0793.

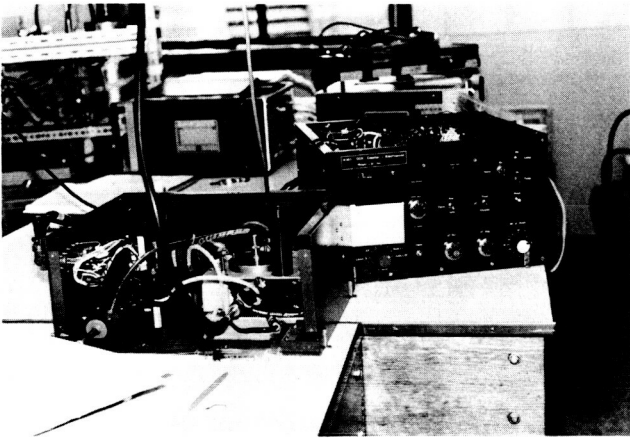
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A DESCRIPTION OF THE UK METEOROLOGICAL OFFICE
CCN COUNTER

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The Meteorological Office CCN Counter is a modified Mee Industries Model 130 CCN Counter which was purchased in 1977.

1. THE CHAMBER

The chamber is a thermal gradient diffusion type with the upper plate at room temperature and the lower plate thermoelectrically cooled. The light from a 25W projector bulb is focused in a beam of approximate dimensions 1x3 mm in the central part of the chamber. The light scattered by drops is viewed in the forward direction (40°) by a microscope and a sensitive photodetector.

Experiments have been performed to relate the peak output from the photocell with the maximum number of drops in the beam recorded photographically. The microscope is used to check the calibration at low concentrations of CCN.

2. ANCILLARIES

Modifications to the original Mee design are as follows:

(a) In order to reduce the time necessary to obtain an activation spectrum, the thermal capacity of the cooling system has been reduced by mounting the thermoelectric cooler adjacent to the bottom plate. In order to reduce the temperature difference between the plates (ΔT) and hence the supersaturation, the current is reversed through the cooler, instead of a separate heater being employed.

(b) The temperature of the top and bottom plates is monitored using precision surface mounting thermistors (YSI 400) bonded directly to the back of the sintered bronze plate. Horizontal temperature gradients across the lower plate are minimized by placing a brass disk, into which water drainage channels have been machined, directly below the plate and on top of the Peltier cooler.

(c) The new control circuits enable a sampling sequence in which an activation spectrum consisting of CCN measurements at a predetermined number of supersaturations and over any desired range to be made. ΔT is stepped up and back down through this range automatically. There is also a circuit which monitors ΔT and prevents new air samples being taken until the desired level of temperature stability in the chamber has been reached. At present, this is set to $d(\Delta T)/dt \leq 0.01^\circ\text{Cs}^{-1}$. A typical portion of this sequence is shown in Figure 2 below.

(d) The chief use of the instrument is on-board the Meteorological Research Flight C-130 aircraft. To make the instrument suitable for this use, it has been repackaged and is powered by a 115V, 400 Hz aircraft supply. For use in ground-based field projects, data logging is performed by an HP9830 calculator.

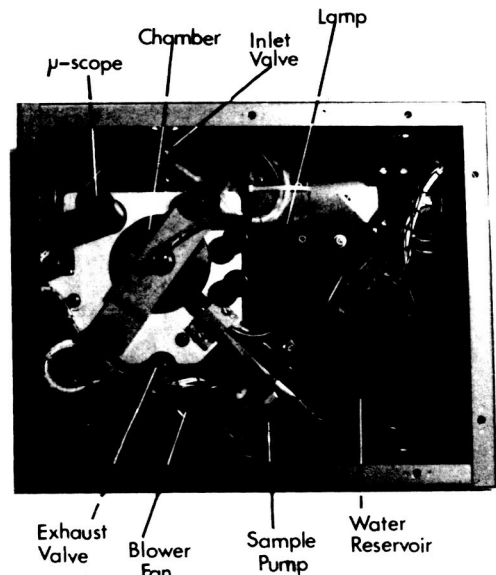


Figure 1. A view of the chamber; the control electronics, power supplies, etc. are in a separate box, not shown).

* Mee Industries Inc., Rosemead, CA 91770

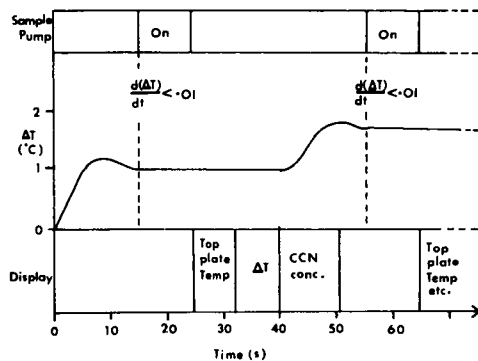


Figure 2. A typical sampling sequence. The "display" is a digital display on the front panel and the "CCN concentration" is the peak photocell output after amplification. A four-point activation spectrum in the range of supersaturation 0-1.5% requires approximately 10 minutes to complete.

3. PERFORMANCE

From experience in the operation of the modified Mee counter at the CCN Workshop and during field projects, it has become clear that there are a number of deficiencies in the design which have yet to be corrected.

(a) The low intensity light source results in a poor signal-to-noise ratio in the photodetector output.

(b) Calibration is difficult because photography of the drops growing in the chamber is hindered by the low illumination.

(c) The high fall speed of the drops viewed through the microscope is evidence of flow inside the chamber air sample. This raises doubts about whether a quasi-thermodynamic equilibrium is achieved.

(d) The sintered bronze plates are difficult to clean and may be subject to contaminants.

(e) Condensation occasionally occurs on the chamber windows, reducing the amount of scattered light focused onto the photodetector.

4. CONCLUSION

The modifications made to the Mee CCN counter have improved the temperature control and stability and also reduced the time taken to produce an activation spectrum. However, measurements of CCN concentration using this instrument have shown that its performance is still affected by design deficiencies.

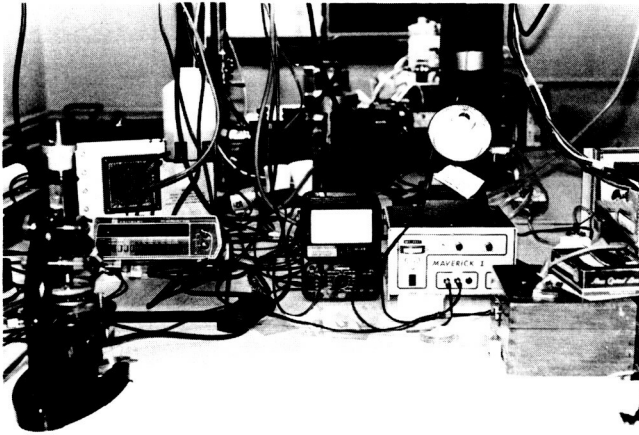
5. ACKNOWLEDGEMENT

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CLOUD CONDENSATION NUCLEUS COUNTER BY
IMPACTOR SAMPLING TECHNIQUE

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Unlike typical CCN counters, this device counts the numbers of water droplets condensed on aerosol particles sampled on a microcover glass at various different relative humidities. The relative humidities ranged from 75% to a calculated value of 110%. A schematic of the apparatus is shown in Figure 1. The individual CCN can be identified in an optical micrograph and scanning electron micrograph and may be inspected for their chemical composition later.

Sampling is made by means of an impactor and a vacuum pump of about 3 liters or less air volume onto an area of 3.35 mm by 0.25 mm of a microcover glass (0.2 mm thick). Then the cover glass is transferred to a small humidifying chamber where temperature is controlled by a thermoelectric cooler at the bottom of the chamber. The chamber has a closed glass top, and the circular side wall of the chamber has a blotting paper soaked with a saturated aqueous solution of sodium chloride, giving an equilibrium relative humidity of 75% in the temperature range between 20° and 70°C.

As the cover glass is cooled down by applying a direct current to the thermoelectric cooler, it is chilled to a lower temperature producing a relative humidity of 100%. Additional current produces relative humidities beyond water saturation. The exact saturation point is identified by the observation of dew on the cover glass which was previously coated with aluminum, using a vacuum evaporation to form a mirror. Simultaneously, the temperature difference between the cover glass and the air is observed by means of a thermojunction and a microvolt meter from the starting point of cooling, giving the "0" reading.

Humidity values are determined from interpolation and extrapolation of the microvolt reading at the point dew forms. A usual microvolt reading at the dew point is about 150 microvolts for a 25% humidity difference. In this case, every 6 microvolt change indicates a 1% relative humidity change. Consequently, a 12 microvolt reading higher than that necessary to obtain the dew point gives a nominal 2% supersaturation. Due to various cooling water temperatures, the reading at the dew point has been found to change slightly.

The water drops condensed are photographed for counting by a Polaroid micrograph camera or a normal microscope camera. Since the sizes of the water drops on a cover glass are larger than 5 microns, the resolution of an optical microscope is adequate.

The greatest advantage of this technique is to allow the inspection of sizes and chemical composition of the individual nuclei by an electron microscope combined with an X-ray energy spectrometer with a specimen cooling device to prevent possible heating on volatile nuclei. Such analysis is in progress for the aerosols sampled over the Arctic Ocean. The second advantage is that the aerosols sampled can be stored for a long time and their condensation ability can be examined later.

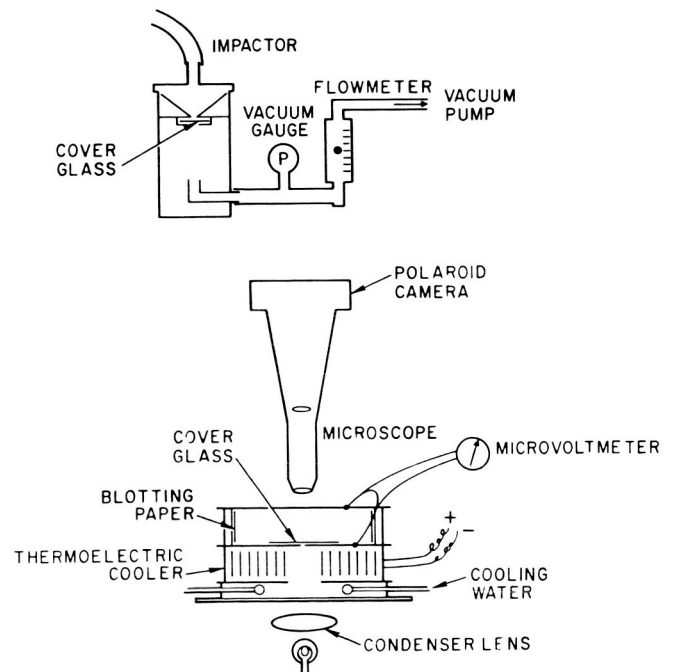


Figure 1

However, as it has probably been seen, this technique may not give an accurate absolute concentration of cloud nuclei. The poor collection efficiency on the sampler may limit the collection of aerosol particles smaller than 0.1 microns diameter, although this may be improved by use of a better sampling technique (possibly by a micro-orifice impactor). If the vacuum or air speed at the nozzle is reduced, small particles will be missing from the cover glass. Thermal precipitator will result in the particles being evaporated. Cooling thermal precipitator using liquid nitrogen will result in frosting on the substrate.

Another problem is the humidity value on a substrate where the aerosols were sampled. Water vapor molecules tend to diffuse to the particles and some condensed water drops will restrict many adjacent aerosols from condensing to water drops as pointed out by Lala and Jiusto (1972) for the case of ice nucleation. Effective relative humidity on substrate may therefore never reach water saturation. For CCN counts at room temperatures, humidity reaches saturation values confirming by observations of visible dew on a very clean mirror. However, the actual degree of supersaturation may not exceed by more than 0.1%, even though the calculated value of supersaturation may be 2%. The real supersaturation degrees for calculated humidity values remain unknown. In order to break the microscale boundary layer over hygroscopic particles, circulation of humidity-controlled air should be adequate. Preliminary experiments have confirmed this procedure to be successful.

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DIFFUSION TUBE

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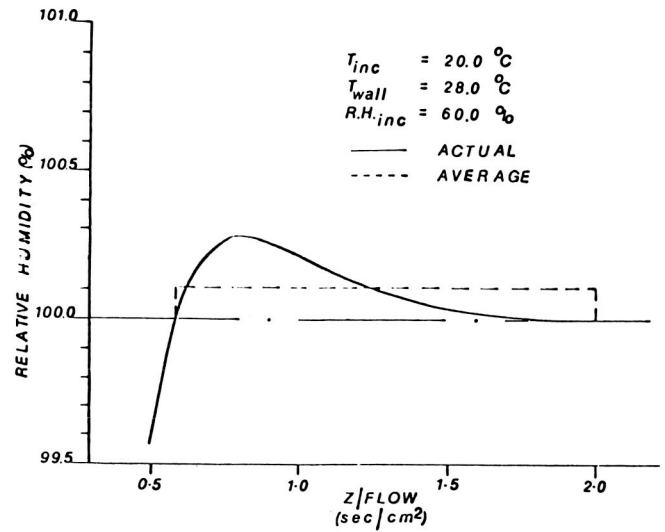
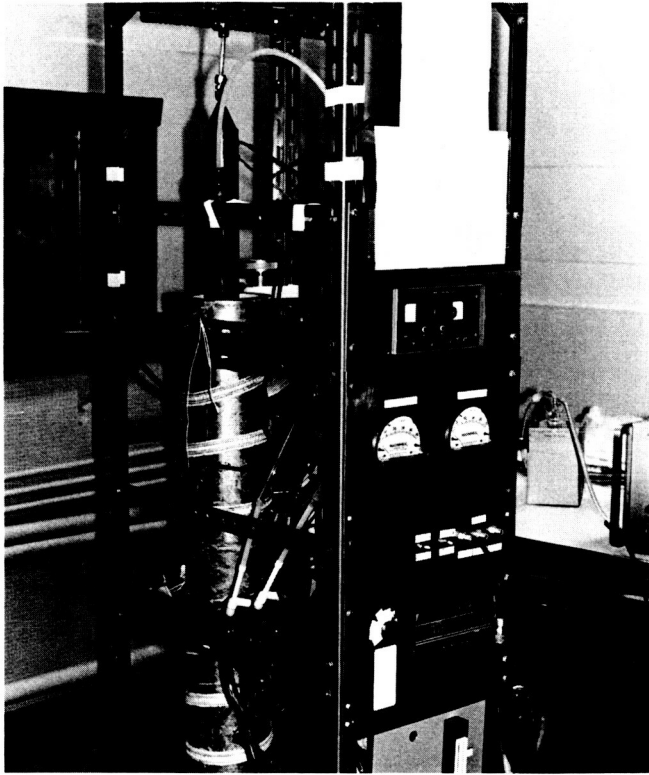


Figure 1. Relative humidity profile, average supersaturation = 0.11%. $Z \equiv$ tube length = 87 cm.

1. EQUIPMENT DESCRIPTION

The diffusion tube is designed to operate below about 0.25% of water supersaturation. It is simply a long tube lined on the inside with a damp chamois cloth, and heated isothermally to a few degrees centigrade above the incoming air.

The diffusion coefficient for water vapour is slightly larger than that for heat, making it possible to supersaturate the airflow. This is the same principle by which transient supersaturations may occur in parallel plate cloud chambers. Our elementary analysis considers only the diffusion of vapour and heat from the walls into the moving air.

The droplet sampling tube which inserts into the main diffusion tube draws the central 4 or 5% of the total volume from the tube. It is only necessary then to compute the supersaturation for the air stream in the centre of the tube. A typical supersaturation profile is shown in Figure 1 (solid line). It is dependent upon the relative humidity and temperature of the incoming air and the temperature of the tube walls. The abscissa represents the tube length divided by the total flow rate. The cutoff is chosen at 2.0 to permit reasonably isokinetic droplet sampling and because the relative humidity is reduced to nearly 100%.

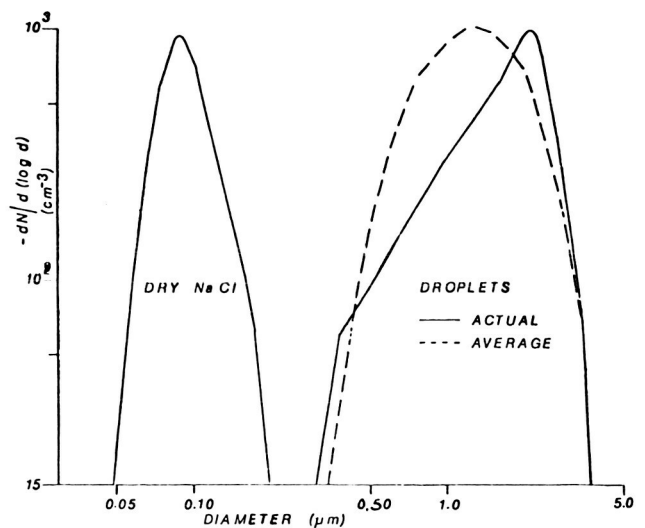


Figure 2. Numerical growth study for profile shown in Fig. 1. Dry distribution is nearly monodisperse.

TABLE 1

Royco Channel No.	Channel Threshold Dia. (μm)	NaCl D100 Value Cr.S.S.(%)	$(\text{NH}_4)_2\text{SO}_4$ D100 Value Cr.S.S.(%)	Channel Used Down to S.S.(%)
1	1.25	0.066	0.088	0.08
2	1.5	0.055	0.073	0.07
3	2.0	0.038	0.056	0.05
4	2.5	0.025	0.037	0.035
5	3.0	0.016	0.024	0.02

In order to define a single supersaturation from this transient, we average the water in excess of 100% relative humidity over the period from the point of reaching 100% to Z/flow equal to 2.0. This we define as the operating supersaturation. The dashed line in Figure 1 indicates this average for this curve. The validity of this approach has been investigated numerically. A dry salt distribution was grown along both paths in Figure 1. The resulting distributions are given in Figure 2. Studies have indicated agreement between droplet distributions is good below 0.1% supersaturation and adequate up to 0.2% at least.

The most serious difficulty occurs with the interpretation of the CCN. Since the available growth time (about 10-12 seconds) is short, for small S.S.'s, the droplets are small (1-4 μm dia.) and confusion may result with inactivated haze droplets, whose D100 values are greater than our threshold. During the Workshop, only one channel on our Royco O.P.C. was used to count the droplets. The threshold level was set at 1.5 μm dia.; as a result some haze particles were counted at supersaturations below about 0.06%. This was noted in Experiment 22 and particularly during our second run in Experiment 26. We have tried to overcome this by evaluating the CCN according to Table 1. The CCN are determined from the counts in Channel 1 down to an operating supersaturation of about 0.08%. Channel 2 is then used to 0.07%, etc.

Experiments with monodisperse aerosols generated with our own classifier have suggested the following:

1. Very good resolution is possible at low supersaturations, down to at least 0.05%.
2. Our estimation of the operating supersaturation is high by about 30% of our value.
3. Results for similar conditions are very reproducible.

The combination of 1 and 3 are encouraging, but work must be continued to assess the proper supersaturations. The high values are thought to be attributable to free convective activity and a slight temperature gradient between the wall temperature sensor and the actual wet surface. Both are being investigated. We expect that at higher supersaturations the free convection will be a greater problem because of larger temperature gradients involved. This is in agreement with a general decrease in accuracy with increasing supersaturation noted during our classifier experiments. The prob-

lem may be somewhat alleviated by beginning with higher initial humidity levels.

The computed operating supersaturations for different wall temperatures and humidity conditions, with an incoming air temperature of 20.0°C, are plotted in Figure 3. It is quite obvious the supersaturation is dependent equally upon the initial relative humidity as well as temperature differences between the wall and air. The dependence, on both parameters, lessens towards the lower supersaturations thus indicating better accuracy, in terms of temperature and humidity measurements, for the small values.

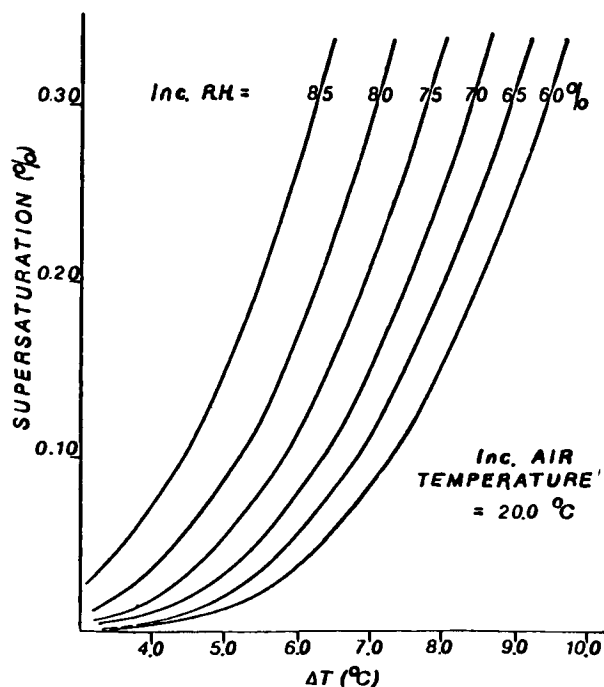


Figure 3. Operating supersaturations as a function of the temperature difference between the wall and ambient, for different incoming relative humidities.

2. OPERATION

The total flow rate is 2.55 L/m; there are two input flows. Humidified filtered air is mixed 3 or 4 to 1 with the air to be measured. A Royco 225 samples about 250 cc/m from the centre of the air stream (0.2 of the radius). The sampling tube extends 12 cm into the tube bottom (see Figure 4), with the remaining flow (about 2.3 L/m) being drawn out by two tubes at the very bottom. The residence time in the sampling tube is no more than 0.1 seconds.

The relative humidity of the incoming air is kept between 65 and 80%. The water bath is heated to about 8 or 9°C above the ambient. Measurements are taken in between adding cool water to lower the wall temperature. Initially it was hoped to vary the incoming humidity while keeping the temperature gradients constant but we encountered some difficulty with the response time of the humidity sensor to large changes.

The temperatures are recorded with YSI precision thermistors ($\pm 0.1^\circ\text{C}$). There are six located in the tube walls. The measured temperature appears stable and quite isothermal. The humidity is measured with dry and wet bulb thermistors. The wet bulb thermistor is encased in a cotton wick and thoroughly wetted.

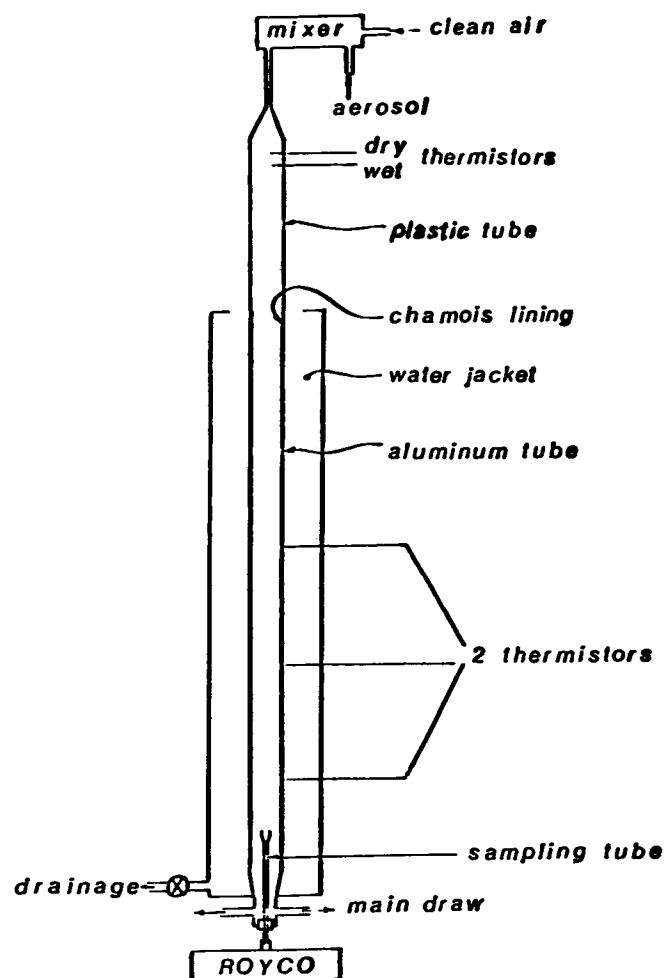
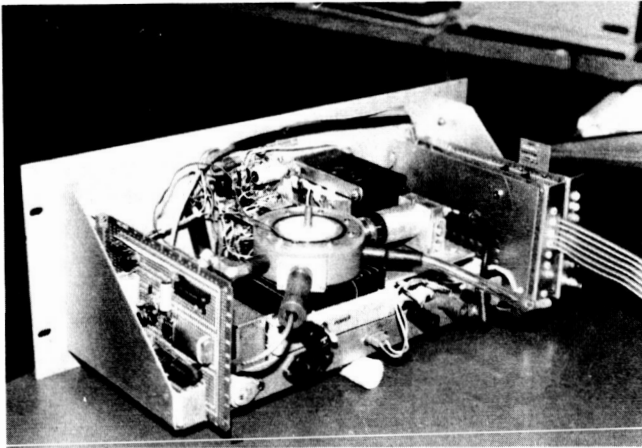


Figure 4. Schematic, in section, of diffusion tube. Aluminum tube is about 100 cm in length, plastic entrance tube is 30 cm long, and sampling tube is 16.5 cm.

AN AUTOMATIC LIGHT SCATTERING CCN COUNTER

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1. PRINCIPLE OF OPERATION

The ASRC cloud condensation nucleus counter is a static thermal diffusion chamber which has been modified to include an optical system for the determination of droplet concentration by the measurement of scattered light. The principle of operation is the same as that described by Lala and Jiusto (1977). The determination of concentration is made by measurement of the peak scattered light signal from the cloud of growing droplets which is a function of both the droplet concentration and chamber supersaturation. Because the formation of the peak is related to the rate of growth of the droplets and sedimentation, both of which are determined by supersaturation, the system calibration can be uniquely determined by comparison with an absolute counter such as a static diffusion chamber with a photographic recording system. This approach to the measurement of droplet concentration in the cloud chamber has made possible the design of a compact system with low power requirements which can be operated automatically under electronic control to provide real time measurement of cloud nucleus concentration.

2. CCN SYSTEM COMPONENTS

2.1 Diffusion Chamber

The cloud chamber is a cylindrical volume bounded by temperature controlled wet plates at the top and bottom of a cylindrical plastic wall. The chamber diameter is 7.6 cm with a plate separation of 1.0 cm with a volume of 45.6 cm³.

The temperature of the top and bottom surface of the chamber is measured by means of integrated circuit transistor thermometers (Analog Devices, Inc. AD590) which have a high sensitivity with excellent linearity and long term stability. The sensors are embedded in the center of the aluminum plates approximately 0.1 cm below the surface. The difference in the plate temperature, as sensed by the transistor thermometer, is used as the feedback signal for a closed loop temperature regulator which maintains the temperature difference at a fixed reference level by controlling the current to a thermoelectric cooler attached to the lower plate. This system has been designed to have a long term stability of $\pm 0.02^{\circ}\text{C}$, while having the capability of rapidly changing the plate temperature by heating or cooling (8 sec to stabilize to within $\pm 0.05^{\circ}\text{C}$ for a 1°C change in plate temperature).

The moisture supply for both the top and bottom surface is provided by water saturated blotter paper. Complete saturation of the upper surface is insured by a connection to an external reservoir through holes in the plate with capillary forces providing the flow of water required. A water supply is not provided for the lower surface because once wet, the normal operation of the chamber maintains saturation of the blotter by the vertical diffusion of water vapor from the upper surface.

2.2 Illumination System

The original design of the system described by Lala and Jiusto (1977) used polychromatic illumination from a tungsten halogen lamp. At that time, it was thought that broadband illumination was necessary to insure the smoothing of the Mie scattering peaks required to produce a scattered light signal which was monotonically increasing with droplet size. Subsequent calculations of the scattering of monochromatic light has shown that the angular integration over the scattering volume and collection aperture provides sufficient smoothing of the scattering peaks. The present design, taking advantage of this, uses monochromatic illumination from an infrared-emitting diode (General Electric F5D1) with a nominal emission wavelength of 880 nm. In order to provide sufficient illumination, the diode is operated in the pulsed mode at 6.25 kHz. The diverging illumination from the diode is collimated by a simple lens to form a cylindrical beam with a diameter of 0.4 cm. The principal advantages of using the infrared diode are that its emission wavelength matches the peak sensitivity of the scattered light detector, the power requirements are small in comparison to the tungsten lamp, and the long lifetime of the diode.

2.3 Scattered Light Detection System

The scattered light detection system is identical to that used in the previous design. A lens located at a 45 degree scattering angle forms an image of the cloud on a slit which serves to eliminate stray light to define the length of the scattering volume (1.5 cm). Directly behind the slit is a hybrid photodetector-amplifier (Bell and Howell 529) which converts the scattered light to a voltage signal. The low level light signal is amplified and converted to a D.C. voltage signal by means of a synchronous detector. The overall system gain produces a sensitivity of 500 $\mu\text{V}/\text{droplet}$ at a supersaturation of 0.5%.

2.4 Measurement and Control System

All data handling and timing requirements of the system are handled by a single board microcomputer (Rockwell International AIM65). Communication between the computer system and the temperature control and light scattering system is performed by analog to digital and digital to analog converters. By means of this interface, the computer is able to control all of the system functions, as well as the processing and display of the data. All user control of the system is handled through computer software, making it possible to alter the system operating parameters without having to configure new control circuits for special applications. The use of computer control allows for the optimization of the overall system performance, as well as complete flexibility in the application of the system for CCN measurements.

3. SYSTEM OPERATION

A measurement cycle can be initiated either by the computer or by an input from the operator. After receiving the start command, the computer determines the chamber top plate temperature and calculates the necessary temperature difference required to produce the desired supersaturation. By means of the digital to analog converter, the computer establishes the temperature reference and waits for the actual temperature difference to settle to within $\pm 0.05^\circ\text{C}$ for two seconds before proceeding. After temperature stabilization, the computer opens the sample valves and purges the chamber by means of a small pump. During the last second of the six second sample cycle, an average value of the background signal is determined. The sample cycle is terminated by stopping the pump and closing the valve after a one-half second delay. During the period which follows, the computer continuously samples the scattered light signal and waits for the occurrence of a peak defined by the signal falling below 80% of the highest previous reading. At this time, the peak value minus the background signal is converted to a concentration and displayed along with the chamber operating condition. The complete cycle is completed in 20 to 45 seconds, depending on the supersaturation used.

4. SYSTEM CALIBRATION

The calibration of the system is obtained by direct comparison with a second thermal gradient diffusion chamber equipped for photographic recording. The chamber and temperature control system for the photographic unit are identical to the automatic system. The illumination for the photographic system is provided by a 100 watt mercury

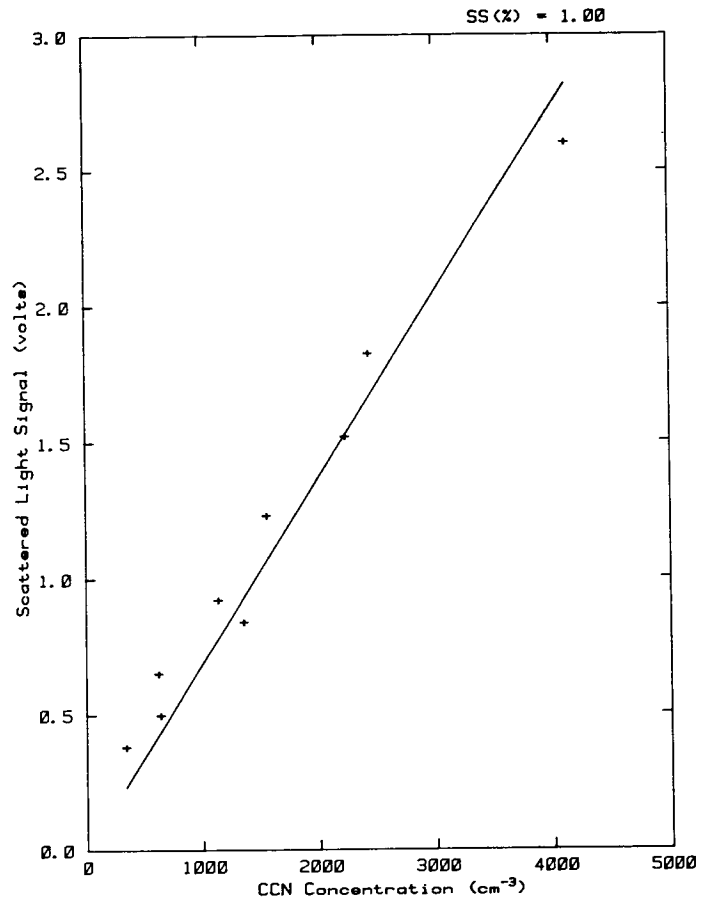


Fig. 1

arc lamp and a large aperture lens system which forms a rectangular beam with an 0.2 cm by 0.4 cm cross section. Droplets in the illuminated volume are photographed at right angles to the beam with a modified oscilloscope camera.

The calibration procedure consists of taking a 200 liter sample of ambient aerosol which is used as a source for both systems during simultaneous measurements. Concentrations are adjusted by diluting the sample with clean air and repeating the measurement after stabilization. Figure 1 shows a plot of the scattered light signal against the concentration determined by the photographic system at a supersaturation of 1%. This procedure is repeated at five supersaturations between 0.25% and 1% which is used to determine the system calibration in terms of concentration and supersaturation.

5. ACKNOWLEDGEMENTS

The assistance of Dr. J.E. Jiusto in the design of the instrument and Mr. L.E. Weed in the construction of the instrument is gratefully acknowledged.

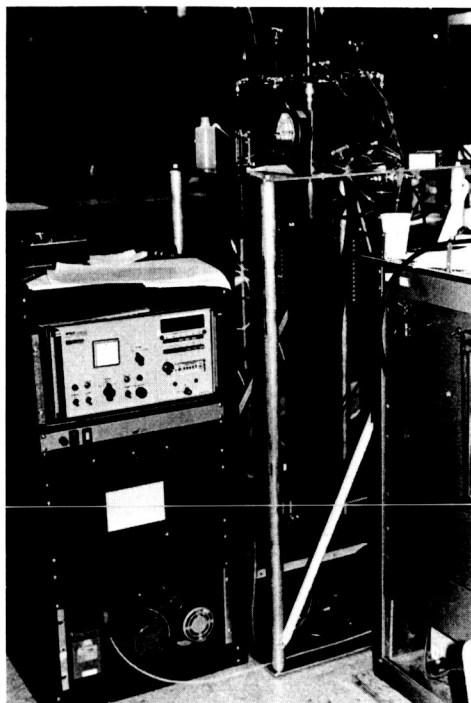
6. REFERENCES

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DESCRIPTION OF THE NRL ISOTHERMAL HAZE CHAMBER

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The background and rationale for using the isothermal haze chamber (IHC) concept to extend the range of CCN measurements to lower supersaturations than is possible with the thermal gradient diffusion chamber is given in the overview section. In the IHC the critical supersaturation is inferred from the measurement of the size of particles which have grown to their equilibrium size at exactly 100% RH. Here the Naval Research Laboratory (NRL) isothermal haze chamber will be described, and the reason for its unique design will be explained.

There are three major design constraints which had to be met:

1. Calculations of the equilibrium size as a function of relative humidity show that the humidity in the IHC must be within about 0.01% of saturation if the inferred critical supersaturation is not to be significantly in error.

2. The smaller the critical supersaturation of the particle, the larger is its equilibrium size. The larger its equilibrium size, the longer it takes to reach that size. Over 100 sec of growth time is needed for a particle with a critical supersaturation of about 0.015% to reach 95% of its equilibrium size at saturation. Much longer times are necessary for particles with smaller crit-

ical supersaturation. The second design criteria was that the chamber must provide over 100 sec of residence time at high humidity. It is this limitation which places the lower limit of about 0.015% on the critical supersaturation of our IHC.

3. The commercial optical counter (Royco) which was available has a sample flow rate of 47 cc per sec. Our decision to stick with the manufacturer's flow rate determined the third major design constraint.

The NRL IHC has cylindrical symmetry as is shown schematically in Figure 1. The design is a two-tube design. The sample first enters a small tube lined with ceramic saturated with water. The pressure drop across the small inlet orifice is less than a quarter of an inch of water and this pressure drop is used to monitor the sample flow rate. The sample flow rate which is about 3 cc per sec was recalibrated at the altitude of the Workshop prior to its start. Filtered air enters through four ports on the outer circumference of a humidifier which consists of passages between concentric cylinders lined with saturated, sponge-like material (see Figure 1). The filtered air, after leaving the humidifier, flows downward through the annular ceramic-lined passage between two tubes and forms a sheath flow around the sample flow beyond the point where the sample emerges from the small tube. At the bottom of the IHC the sample, together with sheath flow (47 cc per sec), enters the optical particle counter where the humidified particles are sized into five size channels between about 0.25 and 3 μ m radius.

The distance required for air passing through a wet-walled tube to reach saturation depends only on the volume flow rate and length of tube and not on the radius of the tube. The reason for using the small tube for initial humidification of the sample is to bring the sample to high humidity in the shortest distance possible. This can best be accomplished by keeping the sample flow, which is very small, isolated from the larger sheath flow until both are fully humidified. The small tube in our IHC is 45 cm long, has a diameter of 1.27 cm and a volume flow rate of 3 cc per sec. Solutions to the diffusion equation show that a sample entering at 50% RH is within 0.004% of saturation by the time it has passed 10 cm down the tube. The residence time in the small tube is about 20 sec.

When the sample joins the sheath air, there is an increase in the cross-sectional area of about 36 and an increase in flow of about 15, resulting in a net increase in residence time per unit length of 2.3. The total residence time in the large tube is about 90 sec, giving a total residence time for growth at high relative humidity of about 110 sec.

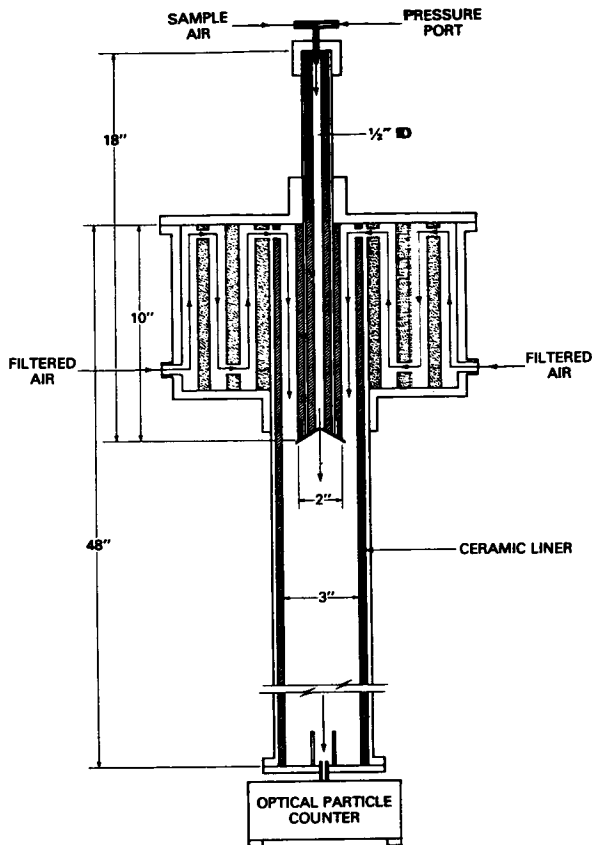


Figure 1. NRL Isothermal Haze Chamber

The value of the filtered sheath flow is multiple: (a) the filtered sheath flow helps prevent gravitational fallout by confining the sample to the central region of the chamber; (b) sheath air is necessary for dilution. The large number of small particles which swell to optically detectable sizes at high humidities will swamp the optical detection system and cause coincidence counting problems if the sample is not diluted with sheath air; and (c) the total air flow required by the optical particle counter is supplied with the minimum overall length by introducing the humidified sheath air after the sample has already reached high humidity.

During a measurement period, the optical counter was on a one-minute counting cycle controlled by its internal timer. However, resetting, starting and reading the BCD output was accomplished with the same HP 9825 computer used to control the NRL mobility analyzer described elsewhere. During each size distribution taken with the mobility analyzer, 12 individual one-minute readings of the NRL IHC were taken and printed out and, at the end of the size distribution, the average of the 12 readings for each size channel was printed out. For most experiments, it was the average of the 12 one-minute samples that was supplied to the Workshop.

The factory calibration of the optical counter was done with latex spheres which have a much different index of refraction than does the water droplets. A size calibration for water droplets

was estimated from the factory calibration and the work of Cook and Kerker (1975).

It is our opinion that the NRL IHC functioned well throughout the Workshop with only a minor problem created by the failure of one of the metering pumps which wet the ceramic. This failure required periodic, partial disassembly and manual wetting on a schedule less frequent than would have been done if the pump had been functioning.

The absolute accuracy of the measurement is difficult to assess. The biggest source of possible error in our system is thought to be the stability and accuracy of the calibration of the optical particle size spectrometer.

The largest size channel corresponds to a critical supersaturation of 0.014%. The growth time required for particles of this size to reach their equilibrium size exceeds the 110 second residence time in our IHC. Since the supersaturation spectrum is always very steep in this region, the contribution of these larger particles which have not yet attained their equilibrium size to smaller size channels is negligible. However, failure of these particles to reach their equilibrium size could result in a significant lowering of the count in the size channel corresponding to the smallest critical supersaturation.

Acknowledgements

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Reference

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MODIFIED MEE INDUSTRIES STATIC THERMAL
GRADIENT DIFFUSION CLOUD CHAMBER

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1. EQUIPMENT DESCRIPTION

The purpose of attending the CCN Workshop was to calibrate and determine the usable operating range of the cloud chamber. The instrument consisted of a Mee Model 130 Cloud Condensation Nucleus Counter (Serial #3). The original optical bench, light source, and detection components were removed and replaced with a simple laser illumination-photographic counting system. The purpose of the modification was to permit discrete droplet counting rather than the original method of scattered light detection and to extend the minimum concentration detection limit to approximately $10/\text{cm}^3$ for use in remote areas where aerosol concentrations were expected to be low. The system consisted of a 5 mw HeNe laser with an operating wavelength of 0.65 microns. The laser beam was incident to the camera field of view at a 40° angle, the same angle used in the original instrument configuration utilizing a photo diode as a scattered light detector. In both arrangements, the detector, in this case the camera, sensed the forward scattered light at the given angle from the center of the parallel plate thermal gradient diffusion chamber. The film used was Kodak 2475 recording film developed at 3000 ASA. Photographs were made at 1/2 second exposure through a microscope attachment to the chamber producing a magnification of 3.47 x on the film negative. The observed volume was 0.011 cm^3 . Temperature and ΔT were controlled by thermoelectric cooling of the bottom plate with the upper plate at ambient temperature.

The operation procedure consists of flushing the chamber for 10 seconds at a sample flow rate of 4 liters per minute. An observation is then made through the eyepiece to determine the time required for maximum droplet number concentration to occur. This time is then used for subsequent measurements

at the same selected chamber conditions. One to 30 exposures are made on each frame to obtain 50-100 droplet images per frame, depending upon the CCN concentration. This produces uncertainties of 10-15% in the CCN concentration determination. At the upper limit of 30 exposures used, concentrations of 10 cm^{-3} can be determined with uncertainties of 50%. The time to determine a four-point CCN spectrum of low concentrations ($<500 \text{ cm}^{-3}$ at 0.5% SS) is 90 minutes. At high concentrations the time reduces to about 30 minutes. Since the droplet counting is from images of droplets on film, the minimum detectable concentration and uncertainty can be selected by the operator with the subsequent loss or gain of time resolution. The instrument in its present form can only be operated manually.

2. RESULTS OF THE WORKSHOP

The experiments performed at the Workshop were undertaken to confirm the calculated sample volume, determine the usable range of supersaturation, and minimum detectable size.

Problems with chamber leaks were discovered for the experiments 0-10. Hence the data from these experiments cannot be used for comparison. Comparison of absolute CCN concentrations with other state-of-the-art continuous flow diffusion chambers at supersaturations near 1% indicated that the volume used produced CCN concentrations well within the range of CCN concentrations determined at the Workshop. This agreement is interpreted to mean the sample volume was correct. Direct measurements of the beam geometry done in the laboratory indicated a factor of 4-5 error. This error is apparently due to the larger apparent visible beam diameter versus the actual usable beam diameter given the droplet illumination, chamber optical geometry, microscope optics and film characteristics. It is therefore suggested that the only way in which the sample volume should be determined is by experimentation with known droplet sizes and numbers via known aerosol generation and sampling.

By comparison of the results from experiments 19 and 15, it was determined that the minimum usable supersaturation for this chamber was between 0.09% and 0.22%, respectively. By using the dry aerosol data from experiments 15, 18, 19 and 20, and the number of droplets detected at indicated supersaturation of 0.1%, the minimum supersaturation the instrument was able to achieve was 0.11%, 0.17%, 0.09%, and 0.19%, respectively. Therefore, 0.20% was determined as the lowest usable supersaturation.

A minimum detectable size of about one micron radius was determined, based upon the smallest observed image on the film, the known magnification

of the image by the microscope attachment, and the artificial enlarging of the image by diffraction of the relatively long wavelength monochromatic light through the small diameter lenses of the microscope. This agrees roughly with a droplet size of 0.7 microns radius at which the maximum in light scattering efficiency occurs for the wavelength light used.

The chamber response to known aerosol at different supersaturations compared favorably to theory, both in the predicted slope for polydisperse aerosol and the determination of the critical supersaturation for monodisperse aerosol. However, there appears to be some tendency to undercount at mid-range supersaturations giving an anomalous concave upward appearance to the curve for some experiments, but not all. The cause of this effect is unknown. Further testing will be required to look more closely at this effect.

TABLE 1. INSTRUMENT PARAMETERS

Detection System:

Laser light source	5 milliwatt 0.65 micron wavelength
Photographic counting	2475 recording film ASA 3000
Minimum detection size	~1 μm radius

Sampling:

Flow rate	4 lpm for 10 sec
Sample volume	0.011 cm^3
Time for spectra (4 pts)	30-90 min

Chamber Specifications:

Aspect ratio	4.6:1
Temperature control	Thermoelectric cooling of lower plate
Usable supersaturation range	0.2 - 2.0%

SIMULTANEOUS OPERATION OF THREE CCN COUNTERS
AND AN ISOTHERMAL HAZE CHAMBER AT THE
1980 INTERNATIONAL CCN WORKSHOP

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1. INTRODUCTION

The Desert Research Institute (DRI) operated four devices for the detection and characterization of cloud condensation nuclei (CCN) and fog condensation nuclei (FCN) during the International CCN Workshop, Reno, Nevada, October 6-17, 1980. In chronological order of development, the CCN devices are the conventional continuous-flow diffusion chamber (CFD), the rapid-cycle CCN spectrometer, and the instantaneous CCN spectrometer. The FCN detection device is an isothermal haze chamber (IHC). These four instruments will be discussed in sequence, and general comments on their performance at the Workshop will be given.

2. CONVENTIONAL CONTINUOUS-FLOW DIFFUSION CHAMBER

2.1 Instrument Description

One of the DRI Continuous Flow Diffusion (CFD) Chambers (Hudson and Squires, 1973, 1976) was used in this Workshop. Since this instrument is largely unchanged since 1976, only the more important features will be described here, along with the values of some of the operating parameters used during the Workshop.

The CFD was operated in the vertical mode; that is, the parallel plates were vertical although the sample traveled in a horizontal direction. The plate separation was 1.3 cm while the width of the plates (this is actually a vertical distance when the chamber is operated in the vertical mode) is 30 cm. The sample enters the working volume of the chamber through a slit in a transverse tube which is at one end of the chamber midway between the plates. It is confined to a 1 mm thick zone about the central plane by the two particle-free air flows which go around each plate. The sample slit is about one-third of the chamber width so that in the other dimension, the sample comes no closer than 10 cm from the "side" walls of the chamber. This allows a minimum aspect ratio of 8. The length of the plates is 40 cm which is the total distance the sample travels through the instrument while the length of the wet paper on the cold plate is 38 cm which allows a 2 cm dry space for the sample after its introduction. The wet paper on the warm plate does not begin for another 10 cm as it is 28 cm long.

2.2 Operation

The main flow of particle-free air enters the chamber at two ports on the backsides of the diffusion plates. These flows then pass over the plates and then turn 180° at their ends where these two flows merge with each other and with the sample flow. By this time, the air should have attained a

reasonably complete temperature equilibrium with each plate. The total flow of air then moves between the plates to the end of the chamber where all of the air is pulled through an optical particle counter (OPC). It is actually the pump in the OPC which produces the air flow through the chamber and the slight underpressure (~ 8 " of H₂O) within the chamber. A valve on the flowmeter which measures the main flow, F, can be adjusted to change F which, during the Workshop, varied from 20 to 58 cm³/sec. Thus the velocity in the central plane of the chamber was $v = 3/2 F/A$, where $A = 39$ cm² is the cross-sectional area. Thus, v ranged from 0.77 to 2.23 cm³/sec so that the time which the sample spent in the supersaturated volume of the chamber varied between approximately 12 and 36 seconds. The lower flow rates (longer times) were generally used for the lower supersaturations where longer growth times are necessary while faster flows or shorter times were used for the higher supersaturations where fallout could be a problem.

The sample flow rate is controlled by a glass capillary tube which restricts the flow to specific values depending on the pressure difference across the tube. The rate of flow through a given capillary was calibrated for the pressure differences to be used while it is in operation. For most of the Workshop experiments, the sample flow rate was about 0.6 cm³sec⁻¹ although it varied from 0.1 cm³sec⁻¹ to 1 cm³sec⁻¹ during the course of the Workshop.

Supersaturations in this chamber during the Workshop ranged from 0.02% to 1.5%. Although supersaturations below 0.1% are normally beyond the useful range of the instrument, certain monodisperse aerosols may yield useful results at such low supersaturations. The time required to complete a spectrum depends on how many supersaturations are used. Although a smaller number of steps generally requires less time, if more steps are used, the time required to change plate temperatures is reduced since the steps are smaller. Typically, it takes at least five minutes to change plate temperatures and at least three minutes to acquire useful data at a given supersaturation.

The drops exit the chamber along with the total air flow. All of this passes through the active volume of the Royco 225 optical particle counter (OPC). In this instrument, the drops (or particles) are counted individually as they scatter light when they pass through an incandescent light beam. The scattered light is gathered by a PM tube which produces a signal which is roughly proportional to the amount of light scattered which is somewhat proportional to the particle size. Since the light beam is larger than the cross-sectional area of the particle stream, all particles which enter

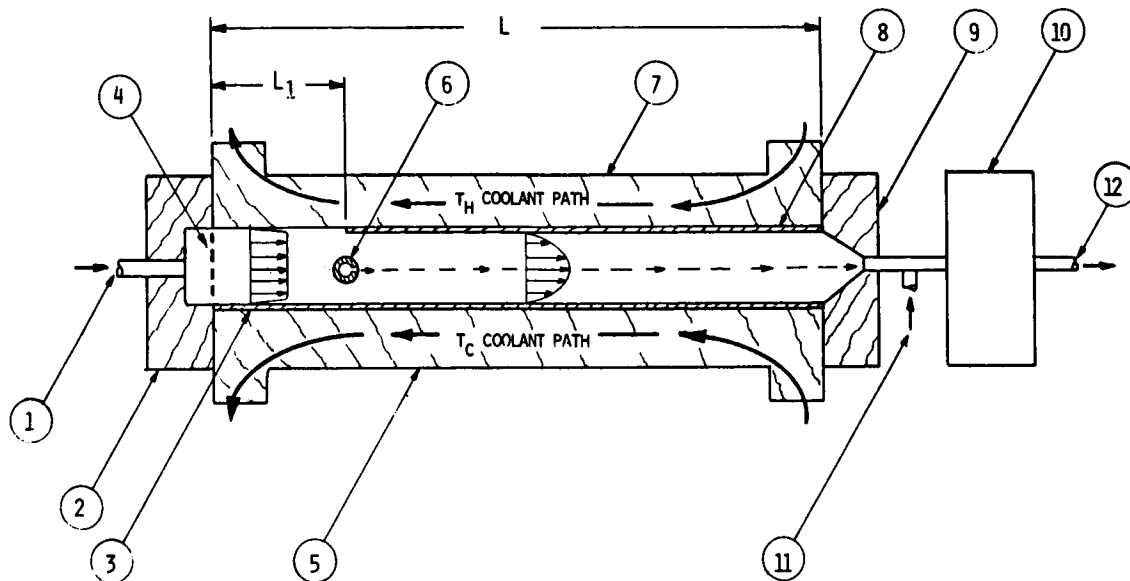


Figure 1. Physical Schematic of the Rapid Cycle CCN Spectrometer.

Legend

- | | |
|-------------------------------|-------------------------------|
| 1. Carrier flow entrance | 7. Warm thermal plate |
| 2. Entrance manifold | 8. Warm plate wicking surface |
| 3. Cold plate wicking surface | 9. Exhaust manifold |
| 4. Diffuser screen | 10. Optical particle counter |
| 5. Cold thermal plate | 11. Sheath flow entrance |
| 6. Sample injection tube | 12. OPC exhaust |

Dimensions: $L = 44.45$ cm
 $L_1 = 10$ cm
 Width = 29 cm
 Plate spacing = 1.6 cm

the OPC scatter light. The chances of more than one particle being within the beam simultaneously are insignificant unless the OPC count is more than 100,000/min so the drops are usually counted and sized individually. When count rates were higher than this, lower sample flows were used.

The plate temperatures are controlled by circulating fluid from constant temperature baths. Plate temperatures are monitored by stainless steel thermistors which are imbedded in the plates just below the wet filter paper on its surface.

This instrument participated in and yielded valid data for all of the Workshop experiments.

3. RAPID CYCLE CCN SPECTROMETER

This instrument was built as a prototype for the NASA low gravity Atmospheric Cloud Physics Laboratory (ACPL). Rapid changes in supersaturation are accomplished by injecting a surge of fluid into the temperature-controlled plates. This displacing fluid is at a temperature different from the original plate temperatures. Thus the change in temperature of the plates is accomplished by proper mixing of an appropriate amount of fluid from a reservoir at an extreme temperature. A hot and a cold reservoir are on hand for this purpose. Microprocessor control is used to inject the fluid into the plates so that a smooth cycle of supersaturations can be obtained. Thus four or five supersaturations can

be examined within a period of 5 to 10 minutes, depending upon the desired accuracy and the particle concentration. This same period of time is required just to make one change in supersaturation with the conventional CFD's.

Figure 1 is a schematic diagram of the rapid cycle CCN spectrometer. It is quite similar even in dimensions to the earlier versions of the CFD's. A notable exception to this is the elimination of the flow around the back sides of each plate. In this version, the carrier flow enters the chamber at the opposite end of the chamber from the optical bench (Royco 225) unlike the previous CFD's, where the carrier flow entered at the optical bench end of the chamber behind both plates only to flow toward the other end of the chamber around the plates and back between the plates. The laminar flow so accomplished is obtained in the spectrometer by the use of a diffuser screen. A stainless steel wicking surface is used instead of filter paper as the wet surface on the inside of the plates.

A hypothetical experiment is shown in Figure 2. Each supersaturation can be maintained as long as desired; each subsequent supersaturation can be established and stable within about 30 seconds.

Each thermal plate is accurately controlled in temperature such that the temperature difference between the two plates is stable and known to

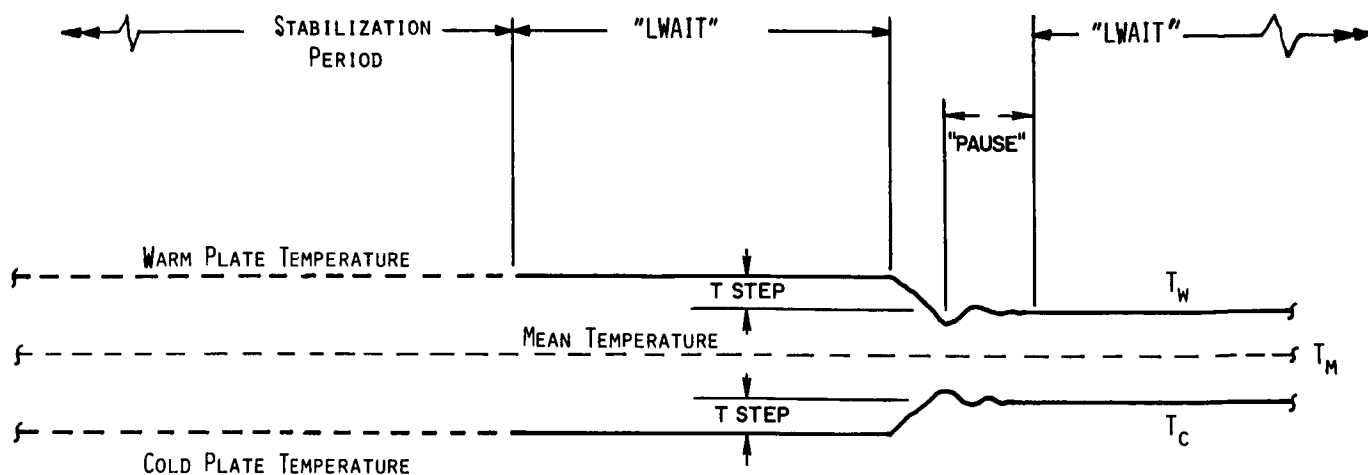


Figure 2. Temperature control of the thermal plates of the rapid cycle CCN spectrometer is programmed to follow the sequence shown below.

Event Sequence:

1. *Stabilization:* ~15-20 min is required to allow the CFD to become equilibrated at the initial temperature settings.
2. *"LWAIT" Period:* The experiment is initiated by a manual command at the start of "LWAIT". This period is long enough to allow sufficient time for counting particles in the OPC.
3. *"TSTEP":* Counting stops at the end of the "LWAIT" period. TSTEP is a temp. parameter

required for reaching the next supersaturation level in the experiment protocol. The computer control servos around the value T_w defined by this variable. Time required for this step is ~2-5 sec.

4. *"PAUSE":* Allows time for system to stabilize at new temp. settings. This has been a period of ~30 sec in preliminary experiment. The end of "PAUSE" initiates another "LWAIT" period at a new supersaturation. The instrument continues to sequence until ΔT reaches a predetermined minimum.

within about $\pm 0.01^\circ\text{C}$ for each supersaturation. Figure 3 is a schematic of the hydraulic circuit for the warm thermal plate. A centrifugal pump circulates water through the thermal plate and a housing containing four electrically resistive immersion heaters in a coolant flow of about $60 \text{ cm}^3/\text{sec}$. The coolant is also circulated over a small glass bead thermistor just downstream of the pump; this method of temperature measurement provides an accurate estimate of the average temperature because of the thorough mixing present at the pump exit. The thermal plate consists of a covered channeled metal surface with large distribution manifolds at both ends. The large manifolds assure that flow through each of the 18 water channels is uniform. Placing the exit and entrance ports on the same side of the thermal plate assures kinematic mixing of the water in the primary circuit. The time required for a parcel of water to flow completely around the primary circuit is one to two seconds.

Temperature control is provided by a secondary hydraulic circuit consisting of a thermoelectric module (TEM) powered "cold source" heat exchanger and a small gear pump. Since this is a closed hydraulic system, operation of the gear pump (trickle pump) displaces cold water into the primary circuit and returns warmer water to the cold source heat exchanger. By correctly metering the proper water flow, any equilibrium temperature of the thermal plate above the cold source temperature can be maintained. Very fine temperature control can be achieved by metering slightly more cold water than is required (slightly overcooling the primary circuit) and adding electrical energy with the use of

the immersion heater. Both immersion heater output and flow of water through the trickle pump are under servo control of the control computer.

A second pump is included in the secondary hydraulic circuit in order to provide rapid changes of thermal plate temperature during the "TSTEP" period shown in Figure 2. The surge pump is operated full on for a period ranging from about 1 to 7 seconds, depending upon the magnitude of the desired temperature decrease in the warm plate. A relatively large parcel of cold water is displaced into the primary circuit during this period, causing a rapid drop in temperature. A period of about 10-30 seconds is required for temperature equilibrium. The surge pump is not used again until another change in plate temperature is desired. The hydraulic circuit for the cold plate is similar to that described above except that the surge pump for the cold plate is attached to a separate TEM-powered heat exchanger (Hot Source), controlling the temperature of the water above the mean temperature of the experiment. Operation of the surge pump for the cold circuit raises the temperature of the cold plate, such that the temperature difference between the plates is reduced in steps as the experiment progresses.

The ΔT measurement is accomplished with the use of a ten-element thermopile mounted on the back surface of each thermal plate. The thermocouple junctions are chromel-constantan and are potted in good thermal contact with an aluminum-base with thermally conductive epoxy. The thermopile is assembled in the form of a yoke; the thin thermo-

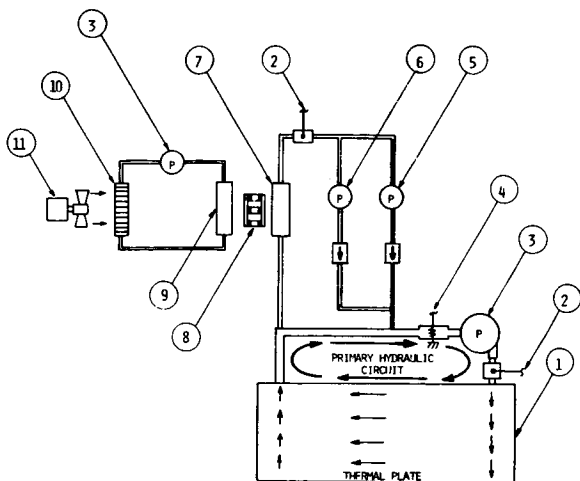


Figure 3. Schematic representation of hydraulic circuit of the rapid cycle CCN spectrometer.

Legend

1. Thermal plate
2. Thermistor
3. Centrifugal pump
4. Servo heater
5. Trickle pump
6. Surge pump
7. Cold source heat exchanger
8. Thermoelectric module
9. Water jacket
10. Waste heat radiator
11. Fan

couple leads are sandwiched between two layers of grounded copper foil tape for support and electrical shielding. The entire bundle is mounted on a thin (0.04 cm x 2.5 cm) strip of fiberglass that forms a yoke around the two thermal plates. The base of the yoke mounts on the thermopile amplifier, located at the bottom edge of the CFD chamber, reducing the lead length of the thermocouples to a minimum. The resolution of measurement is 0.0024°C.

Carrier flow and sample flow rates are similar to those used in the earlier model CFD's. The high degree of precision and accuracy achieved with the earlier models is preserved in the spectrometer. In fact, the increased temperature uniformity and measurement sensitivity allows for increased accuracy. Moreover, this accuracy holds true during the supersaturation cycles of the spectrometer. Thus, the entire spectrum can be monitored with precision and accuracy limited primarily by statistics.

Figure 4 displays an example of an F-plateau in the rapid cycle CCN spectrometer. This shows that the nucleus concentration is independent of the time the sample is exposed to the supersaturation. Thus all particles are allowed to activate but not fall out. This is more fully explained in Hudson and Squires (1976). Figure 4 also displays a comparison in particle concentration between the rapid cycle spectrometer and the earlier model CFD's. Agreement was within 3% which is about the same as the experimental uncertainty.

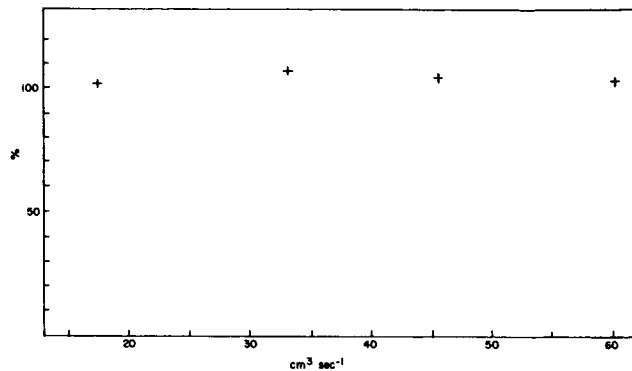


Figure 4. Relative concentration of CCN detected by the rapid cycle spectrometer vs. carrier (main) flow through the chamber. Here the count is normalized to the deduced concentration in a conventional CFD operating side-by-side. The supersaturation in both instruments was fixed at 0.80%.

The rapid cycle CCN spectrometer operated very well during the International Workshop; during many of the experiments, it was actually operating under computer control with no operator present. There were three problems, two of which were instrument-related, and a third which may be classified as operator error. First, the chamber had been newly cleaned and reassembled at the time of the Workshop, and was operating with a background count of order 10-20 cm⁻³ (at 1% supersaturation). This count is about an order of magnitude higher than the normal background for this device; after the Workshop it was found to be entirely due to a slightly loose fitting on the carrier flow inlet. Second, the losses of CCN in the sample inlet system were higher than designed for, due in large part to the presence of charged aerosols in many of the experiments. This problem is discussed in greater detail by Hudson and Alofs in a companion paper discussing CFD design and performance. Finally, due both to occasional, necessary operator absences and operator unfamiliarity with CFD operation at low supersaturations, the main, carrier air flow was not always adjusted properly as the microprocessor stepped the chamber through its range of supersaturations. The result generally manifested itself in a tendency to undercount.

4. INSTANTANEOUS SPECTROMETER

4.1 Equipment Description

This instrument was built along the same lines as the DRI continuous flow diffusion (CFD) chamber (Hudson and Squires, 1976). The most important feature of this instrument is that it uses the sizes of the drops detected by the optical counter to deduce the critical supersaturations of the nuclei. Since several size thresholds can be used, this allows the possibility of simultaneously determining the number, N, vs. critical supersaturation S_c, for several S_c's. This is difficult in a conventional CFD where the drops usually achieve a nearly monodisperse size distribution regardless of the range of S_c's in the sample aerosol (Hudson, 1976).

4.2 Theory of Operation

The instantaneous spectrometer, however, contains three supersaturation steps which disperse the drop spectrum over a wider size range. This range is further widened since the sample is exposed to these supersaturations in ascending order. The device, which is shown in Figure 5, is a series of three CFD's inside one chamber. It contains a sequence of three pairs of temperature controlled plates so that a sample aerosol can be exposed to three separate supersaturations (S_1, S_2, S_3). This means that in the first zone only the largest nuclei become activated drops. That is, only those nuclei with S_C 's below S_1 grow into droplets while the remaining nuclei remain as unactivated haze drops. After being exposed to this constant supersaturation, these drops approach a monodisperse distribution.

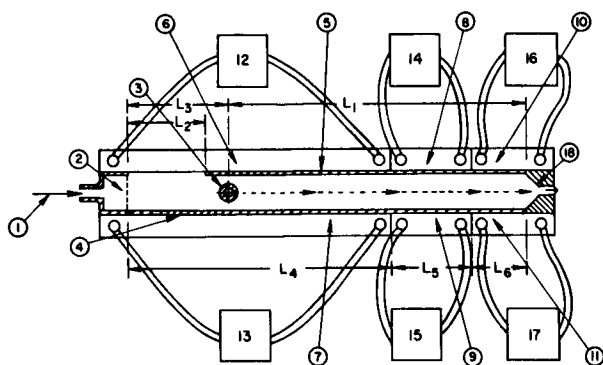


Figure 5. Schematic of the instantaneous CCN spectrometer.

Legend: $L_1 = 48$ cm; $L_2 = 8.4$ cm; $L_3 = 10.4$ cm; $L_4 = 38.4$ cm (1st supersaturation zone - S_1, T_3, T_4); $L_5 = 12$ cm (2nd supersaturation zone - S_2, T_2, T_5); $L_6 = 8$ cm (3rd supersaturation zone - S_3, T_1, T_6).

Supersaturation: $S_1 < S_2 < S_3$

Plate Temp: $T_1 < T_2 < T_3 < T_4 < T_5 < T_6$

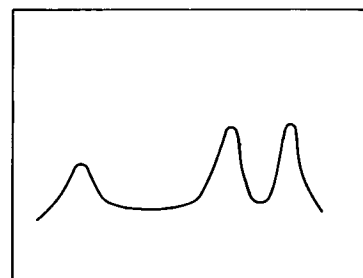
(1) Carrier flow entrance; (2) Diffuser screen; (3) Sample injection tube; (4) Cold plate wicking surface; (5) Warm plate wicking surface; (6) First warm section, T_4 ; (7) First cold section, T_3 ; (8) Second warm section, T_5 ; (9) Second cold section, T_2 ; (10) Third warm section, T_6 ; (11) Third cold section, T_1 ; (12) Temperature bath at T_4 ; (13) Temperature bath at T_3 ; (14) Temperature bath at T_5 ; (15) Temperature bath at T_2 ; (16) Temperature bath at T_6 ; (17) Temperature bath at T_1 ; (18) Exhaust to OPC.

In the next zone nuclei with $S_1 < S_C < S_2$ become activated and grow into cloud droplets with similar sizes. In the meantime the drops which were already activated in the first zone grow even larger in the second zone. In fact, their growth rate is speeded up due to the higher driving supersaturation in zone two. Thus, the nuclei with $S_C < S_1$ grow even larger and somewhat more monodisperse and at the end of the second zone a bimodal drop distribution should result. Finally, the third zone acti-

vates the smallest nuclei (largest S_C) with $S_2 < S_C < S_3$ and a trimodal distribution should result.

The most significant result is not the trimodal distribution but the fact that the drop size spectrum has a wider spread than it has in a CFD. In the spectrometer the drop concentration is less sensitive to drop size and it is easier to discriminate nucleus S_C 's based on drop sizes. Therefore, a small change in the drop size thresholds results in a smaller change in apparent concentration than would be the case with the monodisperse distribution in a CFD. Thus it is much more feasible to relate drop sizes to S_C and to establish size thresholds which correspond to certain S_C 's. If there were no other factors than S_C affecting drop size, then a trimodal drop distribution with clear separations between modes would always result. In that case, size discrimination could be made between the modes and a definite N vs. S_C spectrum could be made which would correspond to the three supersaturations used in the chamber. In such a case, a cumulative distribution would have three plateaus where the number concentration would be constant over a range of sizes. In the CFD there is one drop size plateau which ensures that all nuclei are activated but that none fall to the floor so that a direct determination of N vs. S_C can be made with S_C being the applied supersaturation in the chamber. There are some situations when the instantaneous spectrometer has three drop size plateaus which then allow direct determinations of N vs. S_C for the three S_C 's. However, in most situations the modes are not completely separated (Fig. 6) and instead of plateaus in the cumulative distribution, we find decreases in the slope of N vs. r (Fig. 7). Although this is a much better situation than in the CFD, the lack of a plateau limits the accuracy of direct measurements of N vs. S_C . Accuracy can be increased by setting the voltage thresholds so that the number concentration in the spectrometer matches that in a CFD monitoring the same sample at a specific supersaturation.

INSTANTANEOUS SPECTROMETER
NUMBER VS SIZE



0.93% 0.60% 0.23%

Figure 6. Relative number of drops vs. relative sizes (voltages) for the instantaneous spectrometer. Note that this is a differential and not a cumulative plot.

Although this can also be done with two CFD's (Hudson, 1976) (where one of the CFD's takes the role of the spectrometer and the other one is used to calibrate the first CFD), the process works much

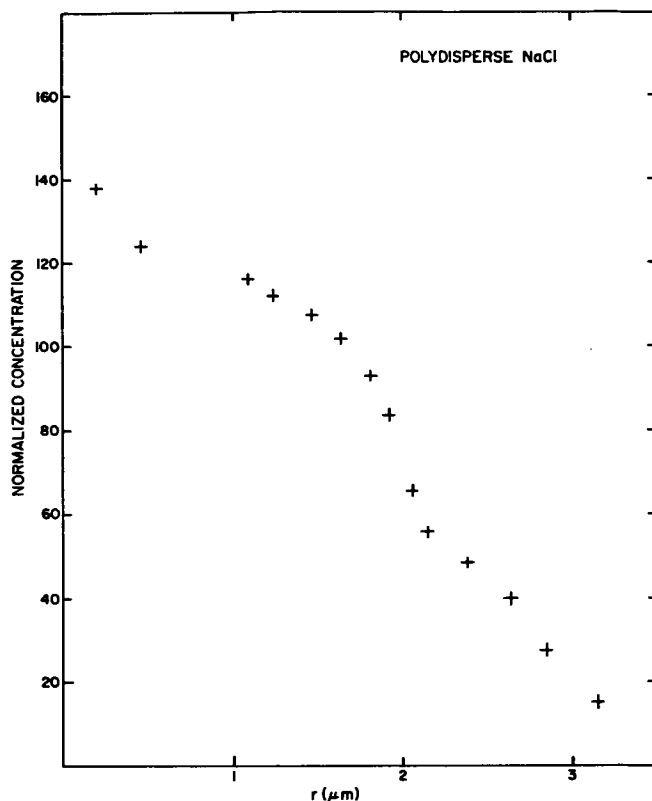


Figure 7. Relative cumulative number of drops vs. size in the instantaneous spectrometer. Applied supersaturations were 1%, 0.4% and 0.15%.

better with the instantaneous spectrometer where there is nearly a constant concentration over some parts of the size range. This considerably reduces the requirement for stability of the various operating parameters. With the instantaneous spectrometer it has been possible to keep the operating parameters constant enough that voltage settings can be used for many days or weeks with continued good accuracy.

4.3 Description

Figure 5 shows most of the dimensions of the instantaneous spectrometer. The plates are separated by 1.6 cm while the plate width is 29 cm. This chamber was also operated with the plates vertical and sample moving horizontally. The main flow through the chamber was $50 \text{ cm}^3 \text{ sec}^{-1}$ throughout the Workshop. This resulted in a particle velocity of $v = 1.62 \text{ cm sec}^{-1}$ so that the sample spent about 31 sec in the chamber; 18.6 sec at S_1 , 7.4 sec at S_2 , and 5 sec at S_3 .

As with the DRI CFD, a Royco 225 optical particle counter is used as the detecting device for the instantaneous spectrometer. In addition a 512 channel analyzer (MCA) (Northern Scientific, Inc.) is also interfaced to the Royco to increase particle size resolution so that greater detail in the concentration vs. size spectrum can be displayed.

The plate temperatures were roughly the same for the entire Workshop so that the supersaturations were nearly constant at $S_1 = 0.30\%$, $S_2 = 0.55\%$ and $S_3 = 0.90\%$. The droplet size thresholds were

set by matching the number concentrations in the instantaneous spectrometer with the concentrations measured with the CFD set at the three different plate temperatures in the instantaneous device. The largest drops corresponded with the lowest supersaturations, etc. All of the drops which could be detected down to the smallest sizes ($\sim 0.2 \mu\text{m}$ radius water drops) corresponded to the number of CCN active at the highest supersaturation in the spectrometer.

Channel 2 was set for about $1.42 \mu\text{m}$ radius water drops while Channel 3 was set for $1.75 \mu\text{m}$ radius water drops. A slight number vs. drop size plateau was observed here and the concentration of CCN in the CFD at 0.55% supersaturation (which was S_2 in the spectrometer) was found to be always less than the number of drops in Channel 2 but more than that found in Channel 3 of the instantaneous spectrometer. This meant that nuclei with S_C of 0.55% produced drops within the size range of $1.42 \mu\text{m}$ and $1.75 \mu\text{m}$ radius in that particular configuration of the instantaneous spectrometer. Thus, the average of Channels 2 and 3 were used to deduce the number of CCN active at 0.55% in the spectrometer. Channel 4 was set at $2.77 \mu\text{m}$ radius water drops and Channel 5 was set for $3.0 \mu\text{m}$ water drops. In a similar fashion, these corresponded to 0.30% S_C . It was found necessary to make a small adjustment in the size thresholds only once during the Workshop.

The sample flow rate was usually the same as the CFD, $0.60 \text{ cm}^3 \text{ sec}^{-1}$, although it was at times as low as $0.1 \text{ cm}^3 \text{ sec}^{-1}$. The plates were also controlled by the same regulator baths and the same types of thermistors were embedded in the plates. This chamber differed from the CFD in three other respects: (1) There were no flows of particle-free air around the backside of the plates. Instead, a diffuser screen was used to eliminate any turbulence; (2) A metal mesh screen was used instead of filter paper for the moist plate surfaces; and (3) Instead of dripping water onto the plates as in the CFD's, water was fed to the metal screens by capillary action from a reservoir of distilled water.

4.4 Operation

Several tests can be performed to check the performance of the instantaneous spectrometer. When the upstream lowest supersaturation, S_1 , is increased, the larger sized droplet peak increases and becomes larger as it should. When the higher downstream supersaturation, S_3 , is increased, the magnitude of the smaller sized peak is increased and there is an increase in its size. Under these conditions, the larger sized peak is only shifted to a slightly larger size. These observations are all in keeping with the operating principles. Thus, sizes which allow separations between the peaks can be chosen. Moreover, the Royco voltage thresholds can be set so that certain size channels can be used to monitor the concentration at specific supersaturations. The size channels can be adjusted so that an individual drop size plateau can be obtained for each supersaturation (see Hudson and Squires, 1976). This assures that all drops which should have been activated at a certain supersaturation were activated and counted. Changes in the downstream supersaturation, S_3 , do not affect the detected concentration active for instance at S_1 or S_2 .

The spectrum of three supersaturations was available simultaneously as soon as the OPC counted and printed out the numbers. Agreement with the DRI CFD was very good and consistent throughout the Workshop as shown by the results.

5. ISOTHERMAL HAZE CHAMBER

The measurement of CCN can be extended to lower values of supersaturation by using the isothermal haze chamber (IHC) first described by Laktionov (1972). The basic operating principle of the IHC relates to the fact that the equilibrium size of a haze droplet, r_{100} , at 100% RH (supersaturation = 0) is uniquely related to the critical supersaturation S_c of the nucleus. Following Laktionov (1972) and Alofs (1978), at $T = 20^\circ\text{C}$ the relationship is

$$r_{100} = \frac{4.1 \times 10^{-6}}{S_c} \quad (1)$$

where r is in centimeters and S_c is in percent. According to Hoppel and Fitzgerald (1977), this relationship is unique if the particle is at least 1% soluble. Eq. (1) can be applied if drops are grown to their equilibrium sizes in a saturated environment and then counted as a function of size.

The Desert Research Institute IHC is a device which subjects sample aerosol to 100% relative humidity for 100-200 s. In most cases, this is enough time to allow the drops to attain their equilibrium sizes at which time they are counted and sized by an optical particle counter (OPC Royco 225). The size is then related to the critical supersaturation S_c so that an N vs. S_c curve can be drawn.

This instrument was built along the same lines as the continuous flow diffusion (CFD) chamber (Hudson and Squires, 1976), in that the sample occupies only a small volume of the cloud chamber which is made up mostly of particle-free air. Figure 8 is a schematic diagram of the DRI IHC. The geometry, of course, is different from the CFD in that the IHC is a right circular cylinder, while the CFD is a rectangular parallelepiped. In the IHC, the sample travels vertically downward (the sample always travels horizontally in the DRI CFD) so that fallout of the large drops still carries them into the optical counter instead of onto the walls where they would be lost.

As with the CFD the total flow F of air can be changed without altering the sample flow f . This allows the capability of changing the time which the sample spends in the saturated volume of the chamber. Therefore, like the CFD, it is possible to give the sample nuclei various growth times before counting and sizing by the OPC which is attached at the outflow end of the cylinder which is the bottom of the chamber.

Using this device, it is possible to experimentally determine whether the drops have had sufficient time to reach their equilibrium sizes. If a plateau exists in the relationship between F and the number of drops larger than specific threshold sizes (Fig. 9), then it is reasonable to assume that these drops have reached their equilibrium sizes r_{100} . According to calculations made by Laktionov (1972), 100 s are required to grow nuclei with $S_c = 0.02\%$ to within 95% of their r_{100} which

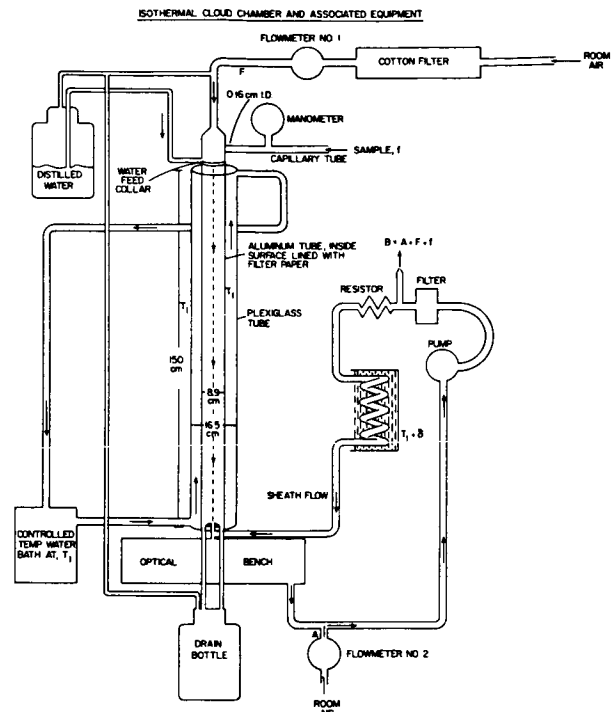


Figure 8. Schematic diagram of the DRI isothermal haze chamber (IHC) and associated apparatus.

is $2 \mu\text{m}$. The available time in the DRI IHC usually approaches 200 s. The existence of an F plateau ensures that drops have enough time to grow to their equilibrium size and that they do not evaporate in the OPC.

Various flows of air through the IHC imply various velocities through the optical counter. In theory, this should not affect the size calibration of the instrument which should only depend on pulse height and not pulse width. Although Alofs (1978) has found that this dependence is not always true at extremely low flows, the flows used in the DRI IHC were never that low. In fact, the size calibration was found to be the same within experimental error for the various flows which were used.

The response of the Royco OPC depends on the index of refraction of the particles. The instrument is calibrated with latex spheres which have an index of refraction of 1.59. Calculations for various OPC's including a Royco have been made by Cooke and Kerker (1975). Although this is probably the best treatment of the subject, it has many shortcomings in this application. The limits of the scattering angles are slightly different for the Royco 225's used in this study and the values for the Royco 245 referred to by Cooke and Kerker. The largest discrepancy is the angle β (the upper limit of the scattering cone) which Cooke and Kerker refer to as 25° for the Model 245; the manufacturer quotes a value of 28° for the Model 225 while the author measured 27.4° on one of the 225 instruments. This discrepancy may or may not be enough to change the results of the Mie scattering calculations. Moreover, Cooke and Kerker did not perform a calculation for index of refraction 1.59 which is the index of refraction of the latex spheres used

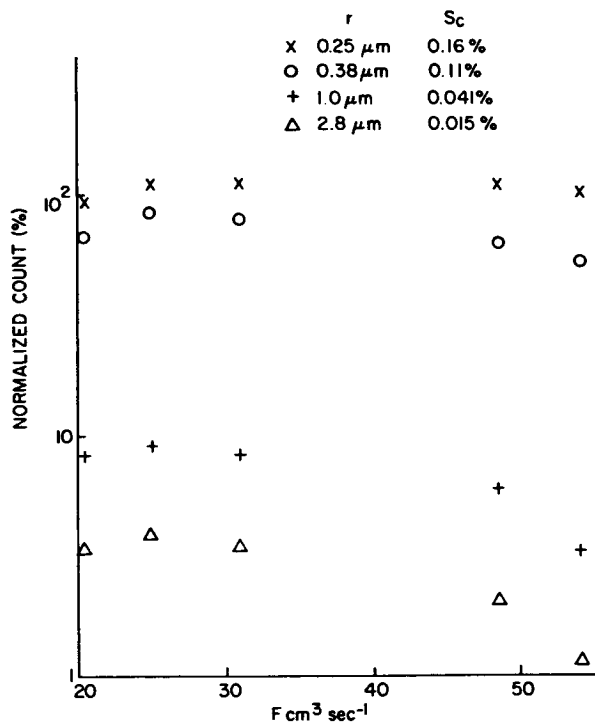


Figure 9. Number of drops exceeding various size thresholds as a function of the total flow in the IHC. The numbers were normalized to the concentration in a simultaneously operating CFD at 0.15% supersaturation. This figure illustrates a plateau in number concentration for various growth times for various critical supersaturations.

to calibrate the device. The best one can do is to interpolate between the 1.54 and 1.70 curves and compare this with the response curve for index 1.33. A correction curve can then be developed so that the instrument calibration performed with latex spheres can be applied to water drops. This index of refraction correction was then applied to the IHC.

This device operated fairly well during the Workshop, but there are indications that the smallest droplets, those grown on FCN of highest critical supersaturation, may have suffered evaporation in the Royco OPC. The consequence of evaporation of this sort is that the IHC tends to undercount in the region of operation overlapping CFD chamber operation, or about 0.1% to 0.2% supersaturation.

6. ACKNOWLEDGEMENTS

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STEADY GENERATION OF AEROSOLS WITH AN IMPROVED CONSTANT OUTPUT ATOMIZER

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1. INTRODUCTION

It is common practice to generate laboratory aerosols of soluble materials with pneumatic atomizers. In a typical device, a solution of the substance to be aerosolized is injected into a jet of air and the liquid is broken up into very small droplets. After forced evaporation, a dry aerosol of the solute is produced. In most atomizers, a solid surface is placed into the droplet-laden jet such that the larger droplets are impacted and a narrower size distribution is produced.

In preparation for the Workshop, a number of commercially available devices were tested that operated on the above principle. Devices included the DeVilbiss No. 644*, the Nano-Mist** and other inhalation nebulizers, and the TSI*** Model 3076 Constant Output Atomizer (COA), specifically intended for laboratory use. The schematic example (Figure 1) shows a vertical cross-section of the TSI-COA which amounts to a slightly modified version of the Collison Atomizer (May, 1973). Its distinctive feature is the ducted portion of the jet within which the liquid is injected, in contrast to the open jet mostly used in various other inhalation nebulizers (as shown schematically in Figure 3).

Despite differences in design, all the atomizers tested suffered from short- and/or long-term fluctuations of their output particle number concentrations. Some of the devices displayed a constant unsteady behavior while others, such as the TSI-COA, occasionally performed well but then flipped into a very unstable mode for prolonged periods. For many types of laboratory investigations, especially where several instruments are involved as in the case of the CCN Workshop, it is essential that temporal stability of test aerosols be maintained within a few percent. The aim of the present study was to elucidate the mechanisms responsible for atomizer instabilities and to find methods for alleviating these problems.

2. STUDY OF OUTPUT INSTABILITIES

In most atomizers, it is not possible to observe the critical nozzle area during operation nor can the nozzle geometry be modified to effect changes in the performance. In order to overcome these difficulties a variable geometry atomizer (VGA) with a transparent housing was designed. Using this device, the relative position and the size of the air jet, liquid feed tube and impaction surface could be varied. Atomizer performance was

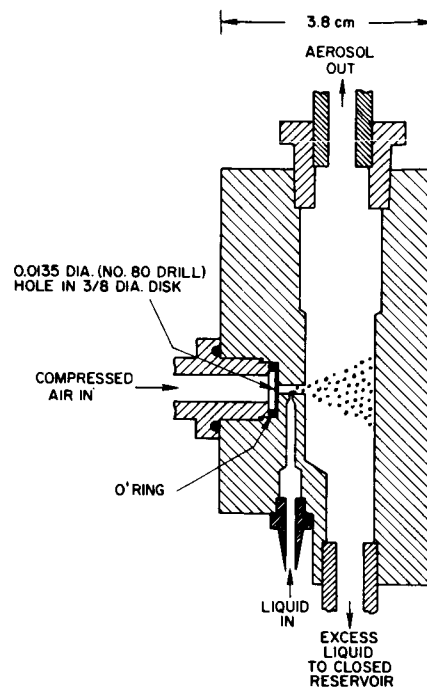


Figure 1. Cross-section of TSI constant output atomizer (from TSI Manual).

assessed by analyzing its output in an electrostatic classifier (EC, TSI Model 3071) and an electrical aerosol detector (EAD, TSI Model 3068) the signal of which was recorded on a stripchart.

Initially, it was thought that build-up of electric charges on non-conductive components of the aerosol flow system were responsible for the sometimes very abrupt changes in aerosol output. However, this interpretation had to be ruled out on the basis of experiments with systems that were made totally conductive.

A series of tests in conjunction with other aerosol measuring devices and with various atomizers confirmed that the observed output fluctuations were not artifacts of the aerosol instruments but, indeed, originated in the atomizer system. The same tests also indicated that the output variations occurred over the entire particle size range, though not necessarily to exactly the same extent. In the case of the TSI-COA, the average output in the unstable operating mode contains a considerably higher proportion of larger particles than in the steady mode as Figure 2 attests.

* DeVilbiss Company, Somerset, PA.

** Eastfield Corporation, Noroton, CN.

*** TSI Incorporated, St. Paul, MN.

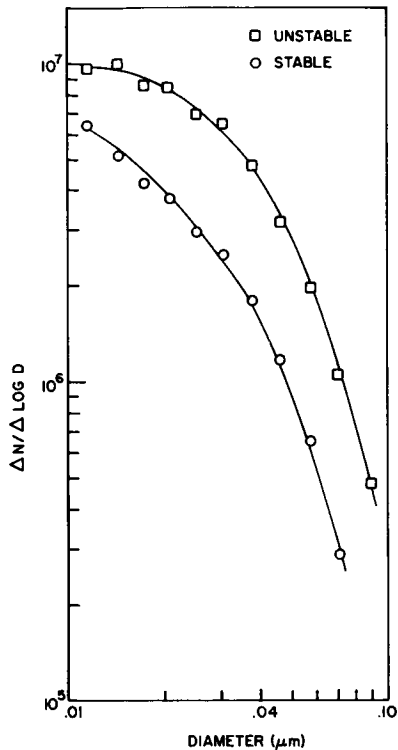


Figure 2. Size distributions of NaCl aerosol generated by TSI-COA at 30 psi with a 0.01% solution in stable and unstable operating modes.

Visual observation of the VGA in operation uncovered several phenomena causing unsteady aerosol outputs. By monitoring the impact area of the jet on a transparent plate, it was found that the size and position of that area was fluctuating in correlation with the particle output fluctuations. Measurements, on the other hand, showed a strong dependence of the output from the mutual position of jet and liquid feed tube. Closer scrutiny (by stereo microscope) indicated:

(a) that in case of wide (~1 mm diameter) liquid feed tubes, the jet and associated turbulence caused the liquid surface to oscillate and, as a consequence, to release liquid at irregular intervals into different portions of the jet, thus generating output irregularities as observed, e.g., in the DeVilbiss nebulizer. This is schematically depicted in Figure 3a, while a narrow tube as in Figure 3b alleviates the problem;

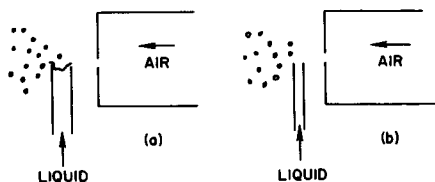


Figure 3. Schematic of open-jet atomizer, (a) with wide liquid feed tube causing irregular output; and (b) with narrow liquid feed providing improved output stability.

(b) that salt deposits at the rim of liquid feed tubes were forming after a period of operation - sooner when working with concentrated solutions, later for dilute solutions. Both the liquid discharge into the jet as well as the jet itself are influenced by the presence of the salt accumulation resulting in erratic output behavior. Removal of the minute salt obstacle immediately restored a steady output; and

(c) that gradually accumulated solution from spray droplets being whirled around between nozzle and impaction surface or solution directly from the liquid feed tube is periodically entrained, or drips into, the jet resulting in temporary output maxima. Concentrated solutions aggravate the problem due to their tendency for foaming, as frequently observed in the TSI-COA where the narrow vertical cavity promotes this undesirable effect.

3. DESIGN AND PERFORMANCE OF THE IMPROVED CONSTANT OUTPUT ATOMIZER (ICOA)

With temporal stability being the main criterion, a new atomizer design was considered essential in order to eliminate or minimize the above mentioned causes of fluctuations found in other units. Thus in a succession of test models, a geometrical arrangement of air jet, liquid feed tube and impaction surface evolved that showed none of the previously described problems.

The top of Figure 4 depicts a cross-section and face-view of the air nozzle and liquid feed combination that evolved. By allowing the liquid to travel in a short, open channel from the tube end to the air nozzle (aided by capillarity and Venturi effect), the formation of salt deposits as well as entrainment of spilled-over liquid was eliminated. A sufficiently large housing (typically ~1 x volume) and the impaction surface positioned

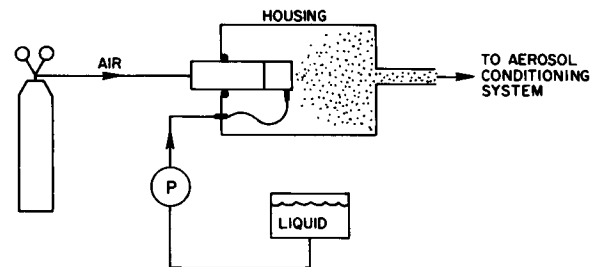
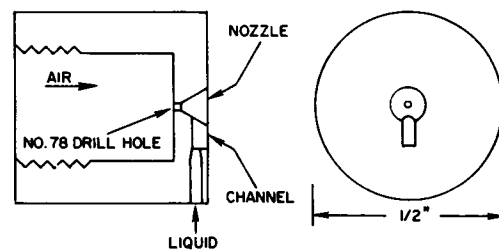


Figure 4. Top: Schematic of ICOA nozzle. Bottom: Schematic of ICOA operational assembly.

several (~ 4) centimeters from the nozzle provided enough space for the spray to dissipate without forming deposits that would drip into the jet. The exit cone of the 0.41 mm nozzle allowed the liquid to enter the jet from all sides and thus further prevented accumulation of liquid near the nozzle.

By operating the unit with a liquid metering pump (such as FMI-RHOCKC*) as shown in Figure 4, bottom, at a rate of 0.05 to 2.0 ml min⁻¹, size spectra as shown in Figure 5 could be obtained at less than $\pm 1\%$ deviation from the average particle number concentration for periods of one hour or more. An illustration of this can be seen in Figure 6 which shows concentration data for the

beginning and end of a six hour run. Occasional minor long term drifts of the output (typically under 2%) have tentatively been traced to the liquid metering pump. The atomizer has been operated successfully at air pressures between 10 and 100 psi; below 10 psi, the output is slightly less stable.

4. ACKNOWLEDGEMENTS

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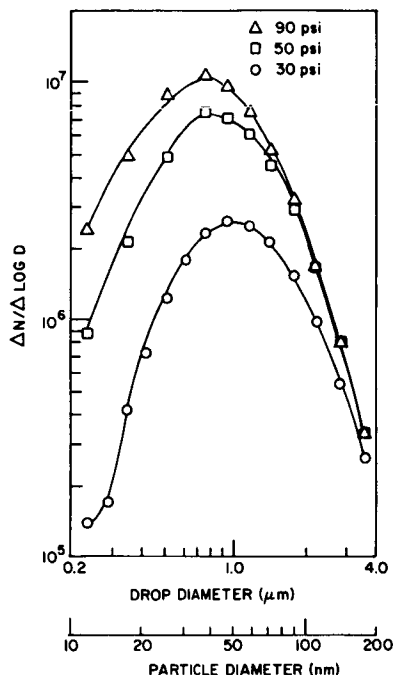


Figure 5. Aerosol size distributions obtained with the DRJ-ICOA at the three indicated air pressures with 0.025% NaCl solution. Upper scale refers to original droplet diameters as inferred from particle sizes and solution concentration.

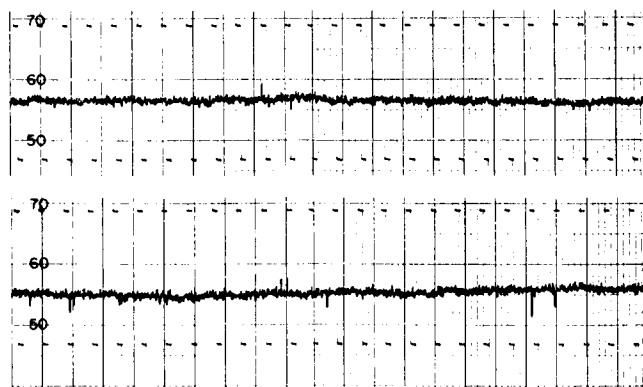


Figure 6. Strip chart record of ICOA output as measured by EC-EAD for a six hour run; shown are first and last 20 minutes, demonstrating good constancy over long periods.

* Fluid Metering, Inc., Oyster Bay, NY.

SECTION VI. SUMMARY - REVIEW PAPERS

CCN COMPARISONS OF STATIC DIFFUSION CHAMBERS

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ABSTRACT

Nine CCN counters of the static diffusion (SDC) type were compared with one another and with continuous flow diffusion (CFD) chambers. The nine SDCs showed a considerable amount of variation, largely attributable to newness and/or lack of prior calibration of some units. The five more consistent instruments agreed quite well, to within at least 20% of the NRL mobility analyzer and to within 10% at 1% supersaturation. There was satisfactory agreement between the more reliable SDC and CFD chambers.

1. INTRODUCTION

At the first International Workshop on Cloud Nuclei in Lannemezan, France (1967), it was not necessary to distinguish between types of instruments for measuring CCN concentrations. Only a few were in existence and present, and all were of the Twomey-type (1963), based on a method of Wieland (1956). In these thermal gradient diffusion chambers, nucleated droplets were detected individually on photographic film. Even at the second Workshop in Fort Collins, Colorado (1970), five of the six units present again were of the thermal diffusion (static) chamber type.

As is evident from the instrument descriptions, a whole host of CCN counters were represented at this Workshop in Reno. They may be classified by type as static diffusion chambers (SDC), continuous flow diffusion chambers (CFD), and low supersaturation "haze" chambers. In this preliminary analysis, we will concentrate on the performance of the static diffusion chambers, although some brief reference to the other instrument types may be appropriate.

At the 1970 Fort Collins Workshop, the five thermal diffusion chambers (excluding one commercial unit) agreed quite well, to within $\pm 30\%$ of the mean (Ruskin and Kocmond, 1971). Chamber operation and expansions were essentially at three supersaturations S : 0.3, 0.75 and 1.0%. By contrast we operated at Reno over a broader supersaturation range, included more S points, and did not prescribe standard S values. In retrospect, the latter was unfortunate and made the data analyses from computer printout data quite laborious and time consuming. The inclusion of many S levels did enhance the data base and allow for, among other things, consideration of CCN supersaturation spectra shapes (e.g., Jiusto and Lala article).

Some of the basic questions that such an analysis can address are as follows:

- How well did the SDC units intercompare?
- Where discrepancies occurred, were there logical explanations?
- Was there general agreement between SDCs and the more sophisticated CFD units?
- Has the science of CCN concentration measurements advanced during the past decade?

Clearly in light of the wealth of data produced and time limitations involved, we will only touch on some of these complex questions. Hopefully, more definitive answers will emerge after further analyses of the data and future experiments stimulated by this International Workshop.

2. CCN INSTRUMENTS

The static diffusion chambers operated at the Workshop are listed in Table 1. Of the nine instruments, only one - the NRL counter - was essentially the same as that used at Fort Collins. This instrument has been used extensively over the years in field programs and in comparisons with other counters. Thus it is considered a useful reference for intercomparisons. The Hebrew University counter was present at all three workshops, though at Reno it employed a different detection method.

As is evident, there has been a trend in recent years away from individual droplet counting in static diffusion chambers to measuring the total scattered light from a sample volume of droplets and transformation to CCN concentrations. Some of the advantages and disadvantages of each approach are discussed in Section 3.4.

Five of the nine SDCs employed a scattered light principle. A related approach apparently was first used by Radke and Hobbs (1969) but with "side" scatter detection over a broad angle, necessary recording of the Mie peak, and a sizable chamber geometry. The commercial unit of Mee, Inc. introduced near-forward light scattering which allows for considerable counter size reduction and portability. Lala and Jiusto (1977) established the theoretical relationship for forward scatter chambers linking supersaturation (drop growth), light scatter intensity and drop concentration; full automation and data reduction via microprocessor control were achieved.

TABLE 1
CCN Counters - SDC Type

<u>Instrument No.</u>	<u>Figure Symbol</u>	<u>Organization</u>	<u>Operator</u>	<u>Droplet (Nuclei) Detection</u>	<u>Light Source</u>
2	O	U. of Wyoming	Rogers/Politovitch	Photographic	He-Ne Laser (5 mW)
5	B	British Met. Office	Kitchen	Light Scatter	25 W Tungsten
8	E	Mee, Inc.	Mee	Light Scatter	25 W Tungsten
9	H	Hebrew U.	Gagin/Nuzitsa	Light Scatter	100 W Halogen
10	S	SUNY	Lala/Jiusto	Light Scatter	I.R. Emit. Diode (25 mW)
13	I	CSIRO	Ayers	Light Scatter	150 W Tungsten
17	N	NRI.	Wojciechowski	Video Camera	200 W Mercury Arc
24	F	France	Serpolay	Video Camera	He-Ne Laser (5 mW)
25	C	CSU	Borys	Photographic	He-Ne Laser (5 mW)

In essence the CCN instruments of SUNY, Hebrew University and CSIRO are reasonably similar in principle and all incorporate a back-up standard photographic method for calibration. The British Met Office and CSU units are modified Mee, Inc. commercial counters. The instrument from France employs a laser light beam and a video camera detector, while the Wyoming counter utilizes laser illumination and records droplet images with a 35 mm camera.

Several of the SDCs had undergone recent re-design or modification, partly in preparation for the Workshop. Others were newly assembled and admittedly brought to the Workshop to test their performance and obtain first-time cross calibrations. This was counter to the original plan of comparing "proven" instruments, but the more open approach undoubtedly added to the overall learning process. Under such circumstances, one might expect a fair spread in the data obtained, and indeed it occurred.

The NRL mobility analyzer (electrical classifier) provided accurate aerosol size information from which supersaturation spectra could be computed with aerosols of known chemical composition (see Hoppel article). These derived spectra also served as a useful reference for instrument comparison.

3. RESULTS

3.1 Overall SDC Performance

One method of evaluating the data was to examine the CCN concentration measurements made with all SDC instruments on all experiments at designated supersaturations. The supersaturations considered were 0.2%, 0.5% and 1.0%. As noted previously, not all counters operated at these specific S values, so a good deal of judicious extrapolation was necessary. The 0.2% supersaturation is at the lower bound of reliability for static diffusion chambers because of problems associated with haze discrimination, non-uniform drop sizes (light

scatter method), and temperature control. As one approaches 0.5 to 1.0%, higher reliability should be expected.

Table 2 lists by experiment the coefficient of variation, and the average CCN concentration \bar{N} of all SDCs (5 to 9) participating in a given experiment compared with the NRL SDC, the NRL mobility analyzer (MA), and the UMR continuous flow chamber. The coefficient of variation V is a dimensionless function of the standard deviation and the arithmetic mean, i.e.,

$$V = \sigma/\bar{N} \text{ or } V(\%) = 100 \sigma/\bar{N}.$$

V is a relative measure of the dispersion of the data and, in a very loose sense for these data, can be thought of as the percentage to each side of the mean that will encompass most of the CCN concentration values.

Table 2 reveals a number of noteworthy items:

1. The three classes of experiments - involving polydisperse salts, monodisperse salts, and ambient air - were not too dissimilar in terms of the presented variables. Thus for simplicity one can concentrate on the overall averages of the bottom row.

2. σ/\bar{N} values of about 0.5 at 0.2% and 0.5% S were comparable and not overly satisfying. At 1% S the degree of dispersion had reduced to 0.32.

3. The concentration ratios (\bar{N} divided by the count of each of the three indicated instruments) show that, on average, the SDCs were under-estimating CCN concentrations. This was graphically evident at the Workshop where certain instruments were obviously registering low counts. Inadequate light intensity or an inefficient optics system are usually the sources of such difficulties. Possible calibration error, depending on the standard used, can also introduce discrepancies.

TABLE 2

Average Performance of All SDC Instruments
(Coefficient of Variation and Concentration Ratios vs. S)

	0.2% S				0.5% S				1.0% S			
	σ/\bar{N}	\bar{N}/NRL	\bar{N}/MA	\bar{N}/UMR	σ/\bar{N}	\bar{N}/NRL	\bar{N}/MA	\bar{N}/UMR	σ/\bar{N}	\bar{N}/NRL	\bar{N}/MA	\bar{N}/UMR
a. <u>Polydisperse Salts</u>												
Exps. 1, 2, 10, 13, 14, 22, 23, 24*	0.51	0.80	0.84	1.01	0.50	0.70	0.58	0.72	0.33	0.86	0.73	0.79
b. <u>Monodisperse Salts</u>												
Exps. 4, 5, 8, 9, 15, 18, 19, 20*	0.46	0.75	0.89	0.87	0.54	0.70	0.80	1.02	0.31	0.87	0.91	0.95
c. <u>Ambient Air</u>												
Exps. 3, 6, 11, 12 16, 17, 26	0.54	0.67	-	0.91	0.57	0.76	-	0.71	0.33	0.85	-	0.84
Overall Average	0.50	0.74	0.87	0.93	0.54	0.72	0.69	0.82	0.32	0.86	0.82	0.86

*0.25% S rather than 0.20% S

TABLE 3

Average Performance of Five SDC Instruments:
NRL, CSIRO, SUNY, Hebrew University, and France
(Coefficient of Variation and Concentration Ratios vs. S)

	0.2% S				0.5% S				1.0% S			
	σ/\bar{N}	\bar{N}/NRL	\bar{N}/MA	\bar{N}/UMR	σ/\bar{N}	\bar{N}/NRL	\bar{N}/MA	\bar{N}/UMR	σ/\bar{N}	\bar{N}/NRL	\bar{N}/MA	\bar{N}/UMR
a. <u>Polydisperse Salts</u>												
Exps. 10, 13, 14, 22, 23, 24*	0.42	0.79	0.95	1.09	0.29	0.77	0.68	0.84	0.19	0.85	0.83	0.89
b. <u>Monodisperse Salts</u>												
Exps. 5, 8, 9, 15, 18, 19, 20*	0.36	0.84	0.98	0.97	0.30	0.84	0.90	1.08	0.31	0.93	0.99	0.97
c. <u>Ambient Air</u>												
Exps. 6, 11, 12, 17, 26	0.48	0.74	-	1.03	0.35	0.91	-	0.89	0.22	0.94	-	0.99
Overall Average	0.42	0.79	0.97	1.03	0.31	0.84	0.79	0.94	0.24	0.91	0.91	0.95

*0.25% S rather than 0.2% S

Because of problems with some instruments, a degree of smoothing was in order. Table 3 presents identical information for just five of the SDC counters whose performance was considered more reliable at the Workshop; these were the units from NRL, CSIRO, SUNY, Hebrew University and France. Even these units on certain experiments registered values that were out of the mainstream. For inclusion of an experiment in this analysis, at least four of the five instruments had to be participating.

The coefficient of variation (Table 3) improved considerably, decreasing to 0.42, 0.31 and 0.24 as supersaturation increased from 0.2 to 0.5 to 1.0%. σ/\bar{N} was somewhat better (smaller) for the laboratory aerosol tests than for ambient air, due largely to greater ambient aerosol fluctuations. Such temporal fluctuations (even with lab aerosols at times) can introduce misleading scatter in the data because the CCN counters possessed different sample processing times and also could not readily be synchronized in time to common S levels.

The overall concentration ratios of Table 3 present a similar but more consistent picture than Table 2 data. At 0.2% the SDC's appeared to underestimate (0.79) with respect to the NRL counter but were within 3% of both the NRL mobility analyzer (0.97) and UMR CFD (1.03). At 0.5% the SDC's apparently underestimated concentrations by about 6 to 20%. At the higher 1% supersaturation, the concentration ratios were all quite respectable, reaching 0.91 to 0.95 values. In general, the SDC's were in somewhat better agreement with the UMR counter than with the other two references. Note that similar agreement would have been obtained with the DRI CFD (instrument #15) which tracked very closely with the UMR counter.

3.2 Individual Experiments and Comparisons

Figures 1-5 depict some representative SDC experiments (not necessarily the best nor the worst cases, but ones that were reasonably well controlled with many participants). The CCN concentrations vs. S data were taken from the printout sheets provided each participant after the Workshop; then averaged where appropriate if several runs were made; and plotted by computer in the fashion shown. The solid lines represent CCN-S spectra computed by NRL from particle size data obtained with the NRL mobility analyzer. Refer to Table 1 for instrument symbol designation.

In the monodisperse aerosol experiments shown in Figures 1 and 2, the critical supersaturations were $\leq 0.2\%$ so that CCN concentrations should be uniform over the indicated S range. Indeed, most data points were within 20% of the plateau. In three other monodisperse aerosol experiments, S_c fell within the SDC operating range of 0.2-1% S. A glance at the Workshop data indicates that most instruments were capable of detecting S_c to within $\pm 0.1\%$ S.

It should be noted that in Figure 1 the British Met Office instrument data were not plotted and in Figure 2 the CSU data were ignored. Both instruments were experiencing difficulties and undercounting. While the Mee, Inc. instrument is represented in Fig 1 (E symbols), it was periodically malfunctioning during the limited days it was

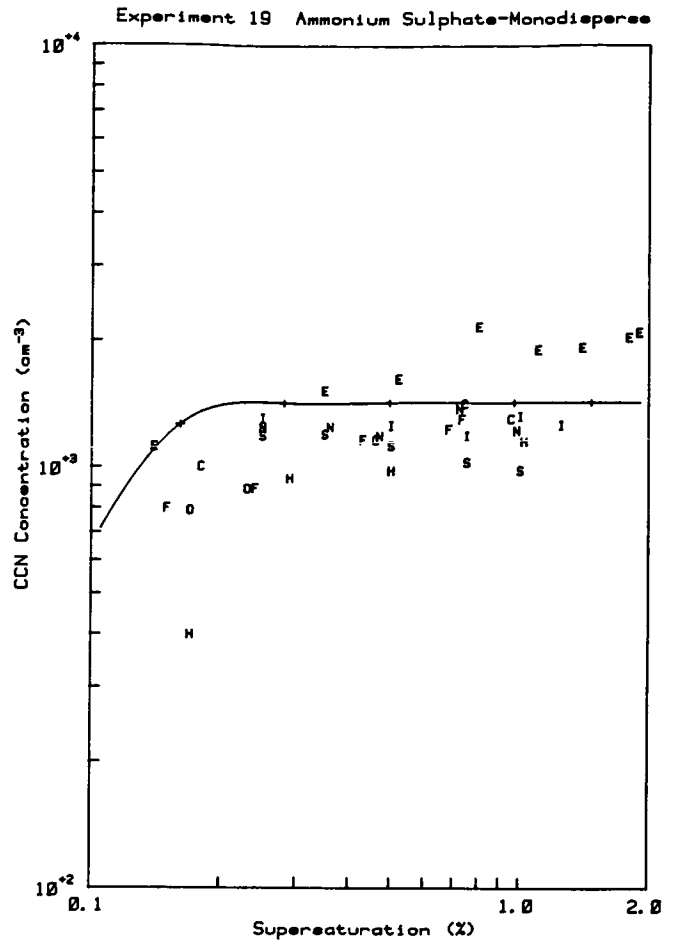


Figure 1. Static Diffusion Chamber Values and Mobility Analyzer Curve (see Table 1 for instrument symbol legend).

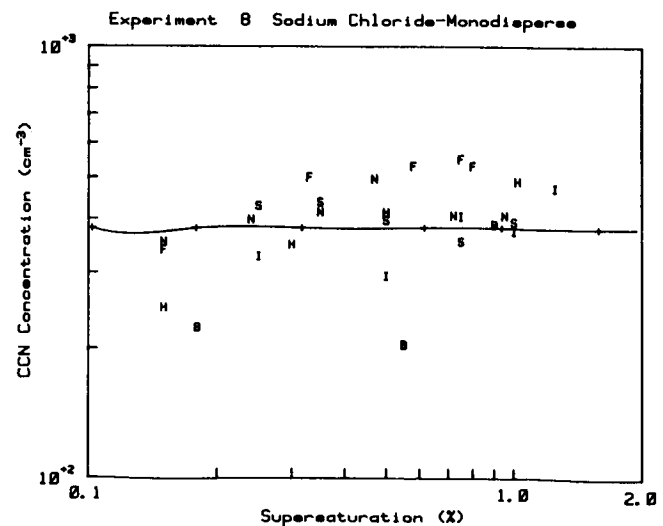


Figure 2. Static Diffusion Chamber Values and Mobility Analyzer Curve.

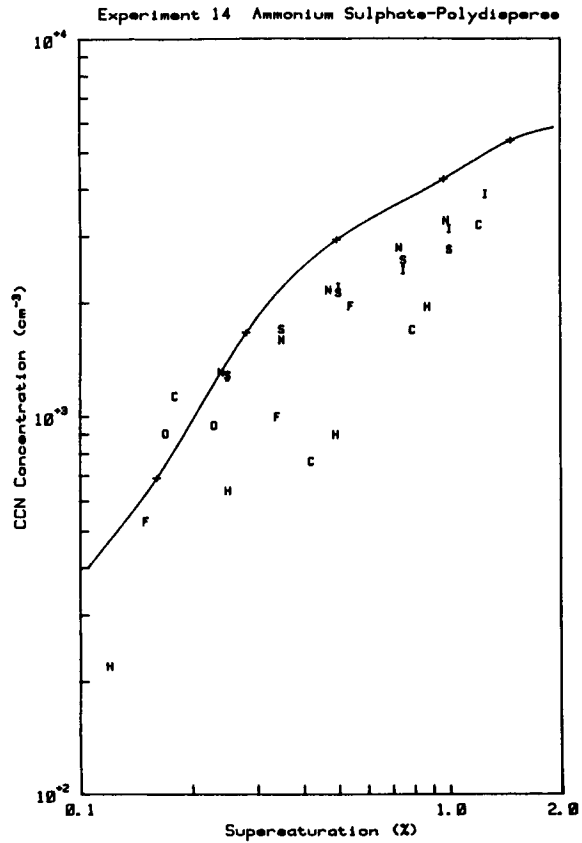


Figure 3. Static Diffusion Chamber Values and Mobility Analyzer Curve.

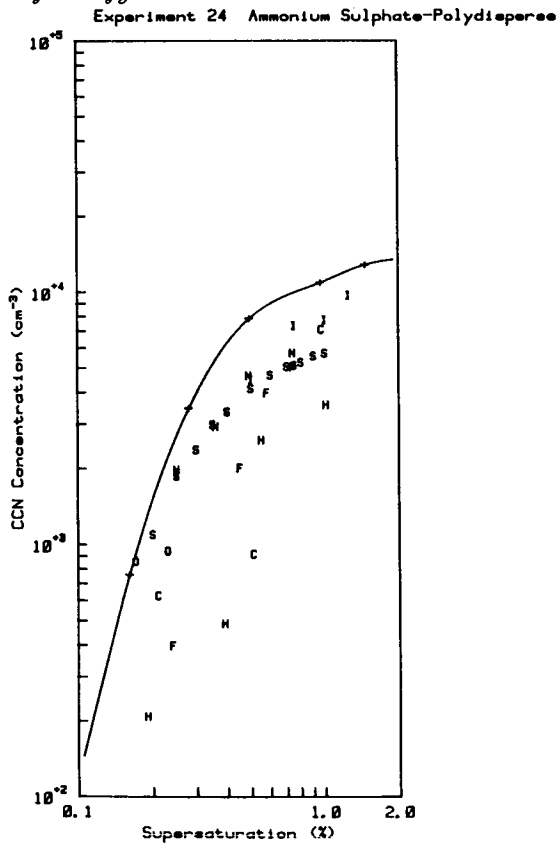


Figure 4. Static Diffusion Chamber Values and Mobility Analyzer Curve.

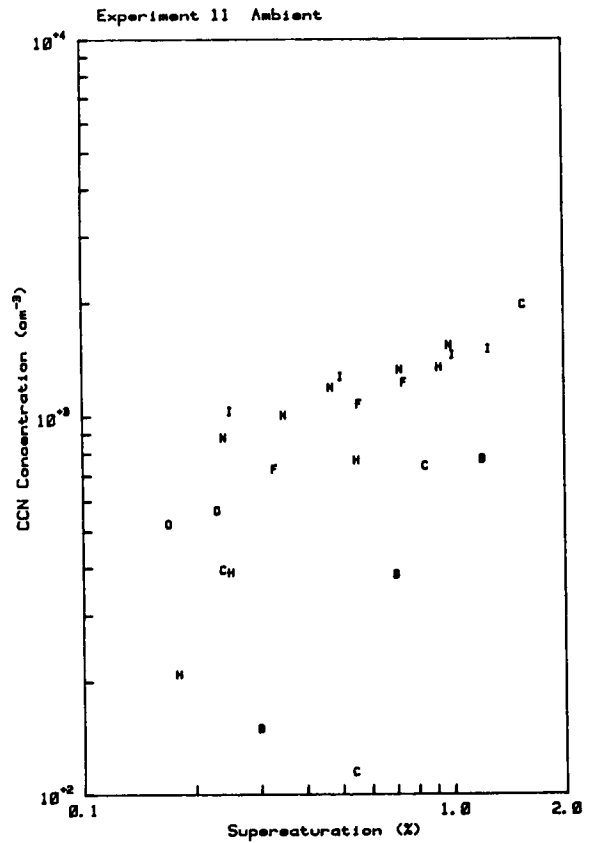


Figure 5. Static Diffusion Chamber Values in Ambient Air.

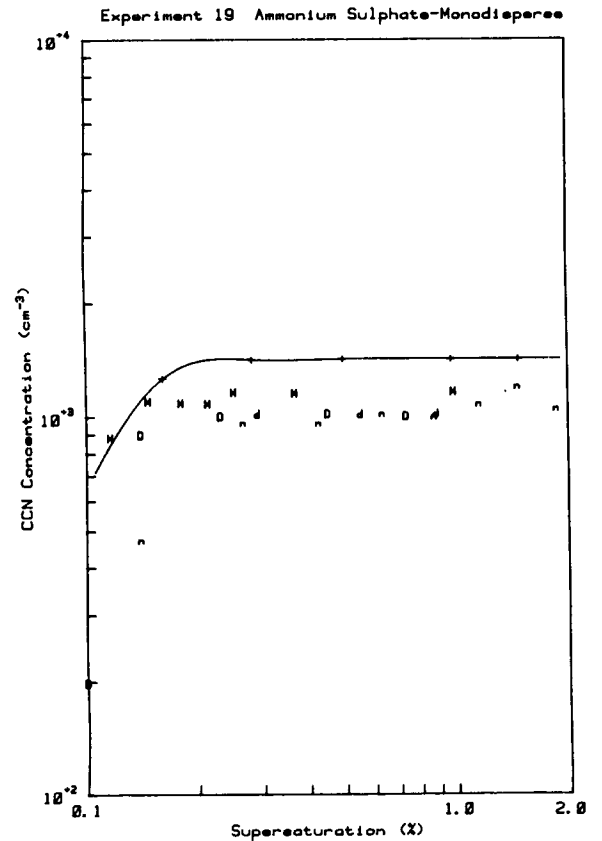


Figure 6. CFD Instrument Values and Mobility Analyzer Curve (see instrument symbols in Sec.3.3 of text).

at the Workshop. It was a new factory unit and reportedly rushed to the Workshop without the customary time for checkout.

Figures 3 and 4 illustrate polydisperse aerosol experiments. Most points in these two experiments lie below the mobility analyzer curve, with considerably more data spread in Experiment 24 (Figure 4). In these and several other cases, the NRL, CSIRO and SUNY instruments tracked reasonably close to one another, with the Hebrew University unit also close at the higher supersaturations. Overall the consistent NRL chamber data matched most closely the NRL mobility analyzer data.

An ambient air case is shown in Figure 5 with considerable data spread. Both the "B" and "C" point instrument are presumably in an early stage of development. The remaining data points cluster reasonably well.

3.3 SDC and CFD Chambers

Figures 6-8 represent CFD chamber data for three of the same experiments just described. Instrument symbols are as follows:

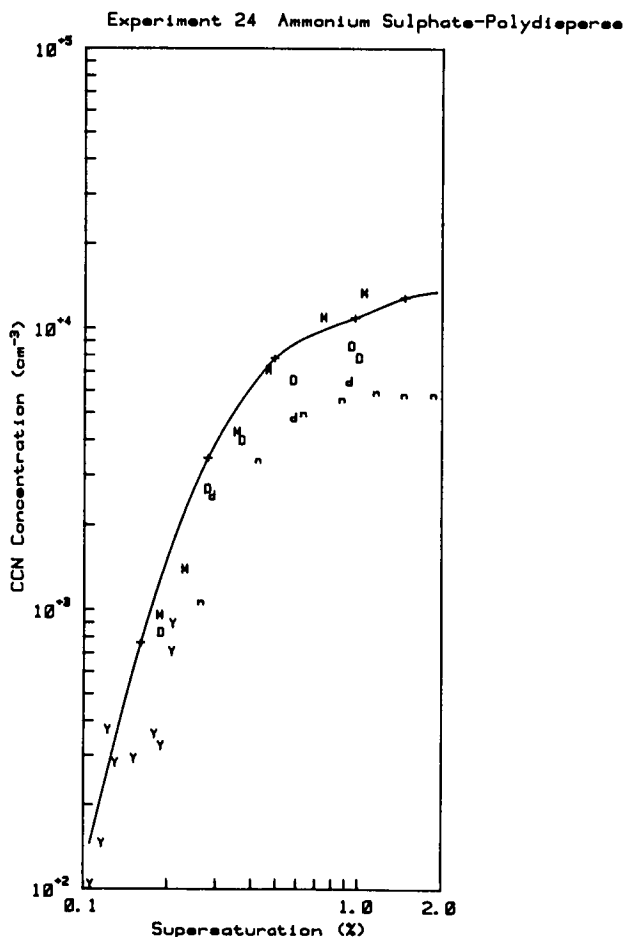


Figure 7. CFD Instrument Values and Mobility Analyzer Curve.

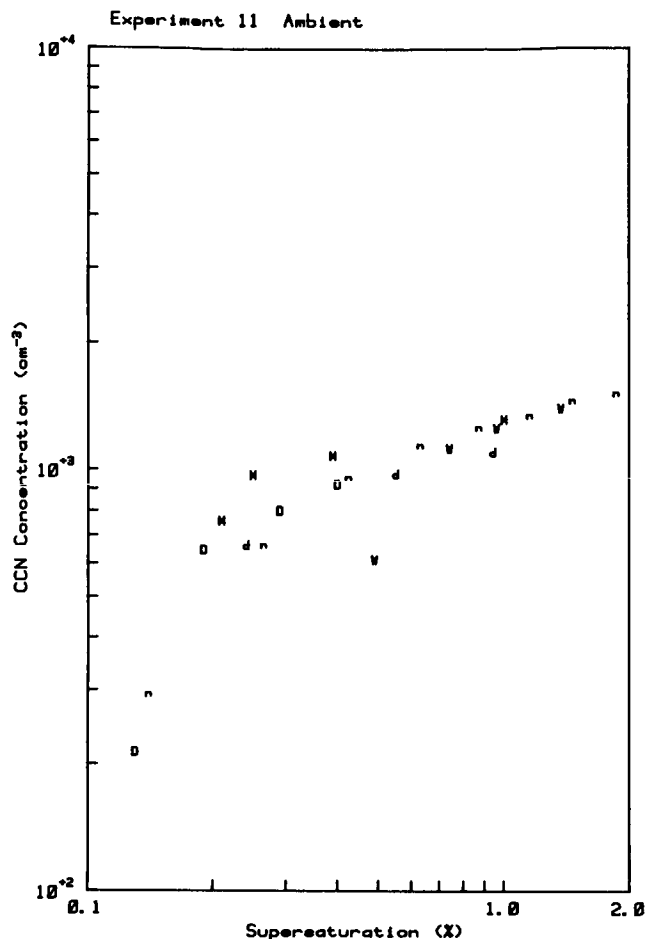


Figure 8. CFD Instrument Values in Ambient Air.

Symbol	Instrument No.	Group
n	15	DRI (NASA unit)
d	16	DRI (Spectrometer)
D	18	DRI
M	21	UMR
W	20	Univ. of Washington
Y	7	York University (different concept/see Leatch and Megaw)

One cannot help but be impressed by the relatively small degree of data spread, particularly in the ambient air case (Figure 8). The fact that only five CFD instruments are involved, that three of these were developed at one institution, and that all have benefitted from extensive engineering and intercomparisons certainly are constructive factors. The results perhaps contain an instructive message for those wishing to develop CCN units.

Comparing Figures 6 and 7 with Figures 1 and 4, it is seen that most CFD points were also just below the NRL classifier curve. By carefully examining these figures (and the ambient case), it is evident that the CFD data and more reliable cluster of higher SDC data points overlapped quite well. In short, we suggest that a well-designed SDC and a well-designed CFD will both perform well. That is evident from the concentration ratios of Table 3

(five SDC's vs. UMR CFD) and from Alofs' determination (described elsewhere in this report) that the UMR CFD and NRL SDC systems agreed on average to 6% at 0.3 and 1.0% S.

3.4 SDC Types

As mentioned previously, SDC instruments now take two principal forms: individual droplet imaging and counting via TV monitors and regular cameras or total light scattering from an ensemble of droplets. The results from this Workshop are not definitive in terms of whether either class is inherently superior. At least one or two of each type appeared to give above average performance while others did not.

An objection to the light scattering method is that at the lower supersaturations, drop sizes are less uniform and less apt to yield accurate concentration data. Similarly with individual droplet detection methods at low S, it is very difficult to discriminate haze from activated droplets. Also with low nucleus concentrations, some subjective judgement can enter as to what frame or portion of a sample volume to count. With either type system, very small sample volumes (light beams) can produce erratic results in low concentration aerosols.

Allowing that Twomey-type chambers with an experienced operator may possess higher inherent accuracy, there is much to be said for the light scattering technique. The latter approach lends itself to more miniaturization, portability, and automatic operation and recording of data. This facilitates field studies of CCN concentrations (and spectral slopes k) on fine time and spatial scales that otherwise would be formidable. A flexible compromise configuration, in evidence at the Workshop, is an SDC that incorporates both the optical and light scattering concepts with the camera operation used at least for periodic calibration.

4. CONCLUDING REMARKS

In summary, the International CCN Workshop provided a wealth of data and information on different classes of instruments and their performance. In view of the massive quantity of data available, much of which could not be thoroughly covered, this report contribution should be considered preliminary.

In general, the more reliable SDCs performed acceptably most of the time. Some are clearly in need of modification and further development. Even those with above-average performance were occasionally out of the mainstream and improvements were undoubtedly suggested to their instrument developers.

The combined average CCN concentrations from five SDC's agreed to within at least 20% with the NRL mobility analyzer, and to within 10% at 1% supersaturation. Agreement between the NRL SDC and the mobility analyzer was even closer. The coefficient of variation (σ/\bar{N}) of the five instruments was 0.42 at 0.2% S, 0.31 at 0.5% S and 0.24 at 1.0% S.

The spread of SDC data was considerable and in excess of that at the prior Fort Collins Workshop. We largely attribute this to the greater number of instruments present, many of which were relatively

new and untested previously. By contrast, four of five CFD units present showed closer agreement with one another, in part for just the opposite reasons cited above as well as for indispensable long-term engineering. The above-average SDC's agreed most satisfactorily (within about 10-15%) with the UMR and DRI CFD's over their common S range.

We began by posing the general question as to whether the science of CCN concentration measurements has advanced over the past decade. We believe it has! Certainly the capability of measuring at supersaturations lower by about an order of magnitude with haze chambers and certain CFD's is an all-important achievement; now virtually all cloud and fog condensation nuclei can be discriminated. CFD instruments have ushered in a high degree of engineering sophistication and accuracy for many applications. The Twomey-type counter, as represented by the NRL SDC, continues to maintain an enviable level of performance. It will undoubtedly remain a useful standard for SDC units. The introduction of light scattering techniques for SDC performance, while not necessarily adding to accuracy, provides for much simpler operation and data processing. The net result may be CCN information with a time and spatial resolution heretofore unavailable.

5. ACKNOWLEDGEMENTS

We acknowledge with deep thanks the DRI Workshop hosts, NASA and NSF sponsors, and all participants and their spirit of cooperation. We also appreciate the efforts of Dr. G. Lala in preparing the figures.

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CCN-SUPERSATURATION SPECTRA SLOPES (k)

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ABSTRACT

Theoretically the slope k of a CCN-supersaturation spectrum ($N = cS^k$) should equal two-thirds of the slope of the total (soluble) aerosol size-distribution. Workshop results tended to verify this relation. As has been noted before, the k values are markedly different depending on whether one is measuring ambient CCN concentrations at supersaturations S above or below ~ 0.1 - 0.2% . The larger k values for $S \lesssim 0.1\%$ is consistent with the greater decrease in large particle concentration with increasing size. Over the S range of 0.02% to 2% , two power fits (and k values) may sometimes suffice for a reasonable approximation of the CCN distribution. At other times, and with laboratory generated aerosols, such an approach is inadequate and requires refinement.

1. INTRODUCTION

Since the inception of cloud condensation nucleus measurements, it has been observed experimentally that nucleus concentration N versus supersaturation S can be expressed reasonably well by the power function

$$N = cS^k \text{ (Twomey, 1959).} \quad (1)$$

With thermal gradient diffusion chambers operating above $\sim 0.2\%$, slope values k typically range from about 0.4 to 1.0. Early indications were that lower values are associated with maritime aerosols and the higher values with continental aerosols (Twomey, 1959; Kocmond, 1965; Jiusto, 1967). Twomey and Wojciechowski (1969) in an extensive aircraft measurement program over various parts of the world reported average k values aloft of 0.5 and 0.7 (0.2 to several % S) for continental and maritime air, respectively.

From Köhler theory for droplet growth, it is evident that the critical supersaturation and size of soluble nuclei are related by $S_c = r^{-3/2}$. Similarly a Junge power-function relation exists between particle size and cumulative particle concentration: $N_{cum} \propto r^{-\beta}$. Combining these relations with expression (1) above lead to:

$$k = 2/3 \beta \quad (2)$$

where β is the slope of the total soluble aerosol size distribution. Junge and McLaren (1971) demonstrated that the shape or slope k of a CCN supersaturation spectrum is more dependent on aerosol size distribution than on aerosol composition, provided the particles are at least 10% soluble. Added confirmation was provided by an analysis of data from the 1970 Fort Collins Nucleus Workshop (Fitzgerald, 1973).

Because Junge total aerosol size distributions often have slopes $\beta \approx 3$ for particle radii $> 0.1 \mu\text{m}$,

CCN k slopes of 2 are sometimes mentioned. This misconception is occasionally encountered in the literature. At chamber supersaturations of 0.2 - 1% , nuclei much smaller than $0.1 \mu\text{m}$ radius are activated as well. Because of their much greater abundance, the resultant k values should be weighted to the flatter portion of the Junge curve and hence yield values less than 2 and generally less than 1.

Conversely, at low supersaturations < 0.1 - 0.2% , where larger condensation nuclei are activated, one should expect higher k values. Such evidence was first reported via direct measurements by Laktionov (1973). Subsequently, the development of "haze" chambers operating at low supersaturations confirmed the trend (e.g., Hoppel, 1978; Hudson, 1980).

Clearly, the topic of CCN spectra shape over a broad range of supersaturation deserves further resolution. This International CCN Workshop, with its variety of chambers, offered an opportunity to explore such questions as:

- Is the relation $k = 2/3 \beta$ generally valid?
- Over what S range can one safely apply the empirical function $N = cS^k$ with a single k value?
- For ambient air, will two k values suffice for the respective supersaturation ranges less than or greater than about 0.2% ?

2. APPROACH

Detailed particle size distributions were obtained at the Workshop by NRL with their highly advanced mobility analyzer. (See section by Hoppel and 23 size spectra so produced). From the NRL cumulative size distributions, one can readily calculate slopes (β_i) between two size increments and also average β over any desired size range. Figure 1 illustrates the equivalent k ($2/3 \beta$) values versus size for the polydisperse NaCl aerosol generated during Experiment 2. The critical supersaturation S_c scale corresponding to NaCl dry particle size is also shown. For the two salts employed at the Workshop, the approximate function relations between critical supersaturation and dry particle size are:

$$\begin{aligned} S_c &= 1.31 \times 10^{-11} r_d^{-3/2} \text{ (NaCl)} \\ S_c &= 2.01 \times 10^{-11} r_d^{-3/2} \text{ (NH}_4)_2\text{SO}_4 \end{aligned} \quad (3)$$

For experiments with ambient air, it was tentatively assumed that aerosol composition could be approximated by ammonium sulfate particles.

CCN spectra values of k were calculated for selected instruments and experiments, based on final Workshop data printouts provided each participant. Slopes k were determined for rather narrow supersaturation increments (three successive S values) as dictated by each instrument's operating

levels. Also k was calculated for broad S ranges over which each instrument normally functions, i.e., ~ 0.2 - 1.5% for static diffusion chambers (SDC) and continuous flow diffusion chambers (CFD) and ~ 0.01 - 0.2% for haze chambers.

Listed in Table 1 are the CCN instruments considered in this preliminary analysis.

TABLE 1
CCN Instrument Numbers, Types,
and Organizations

Instrument No.	Group	Operator	Chamber Type
10	SUNY	Lala	SDC
11	NRL	Hoppel	Haze
13	CSIRO	Ayers	SDC
14	DRI	Hudson	Haze
15	DRI	Rogers	CFD
17	NRL	Wojciechowski	SDC
21	UMR	Alofs	Dual CFD and Haze

3. RESULTS

3.1 CCN Slopes Above and Below $0.2\% S$

Table 2 presents the k values for broad S ranges above and below $\sim 0.2\%$. Theoretical values computed from NRL particle size data ($k = 2/3 \beta$) are indicated followed by actual measured values ($N = cS^k$) for given CCN chambers.

Overall, the comparisons between CCN instruments and between calculated and measured k values are quite respectable. This is particularly true when one considers that the instruments were not operated over identical S ranges or time intervals. The former diluting effect can be appreciated from Figure 1, where it is evident that a modest shift in the S range covered would lead to differing k averages, particularly at the large particle (low S) end of the spectrum. Also, time variations in ambient aerosol concentrations were considerable and would influence the comparative results from different instruments requiring anywhere from 10 minutes to 30 minutes, respectively, to complete a CCN spectrum. The NRL mobility aerosol (size) analyzer data represent the smoothed average of typically 2 to 4 size spectra (~ 25 min each) over the duration of an experiment.

While the number of cases is limited, one may note that for ambient aerosols, the slope values for $S > 0.2\%$ were typically less than 1, and for $S < 0.2\%$ were much steeper ($k \sim 2$ to 3). This is consistent with prior field measurements and with Junge type aerosol distributions. Laboratory-generated aerosols also showed a steeper slope k at low S , reflecting an analogous greater decrease in large particle concentration with increasing size.

If k values for ambient aerosol monotonically increased with size over, say, the 0.1 - 1.0% S range as shown for a lab aerosol (Figure 1 and others to follow), then a single k value would hardly suffice.

3.2 CCN Slopes over Narrow Size (S) Increments

Finer increments of the size and supersaturation spectra were examined in terms of their asso-

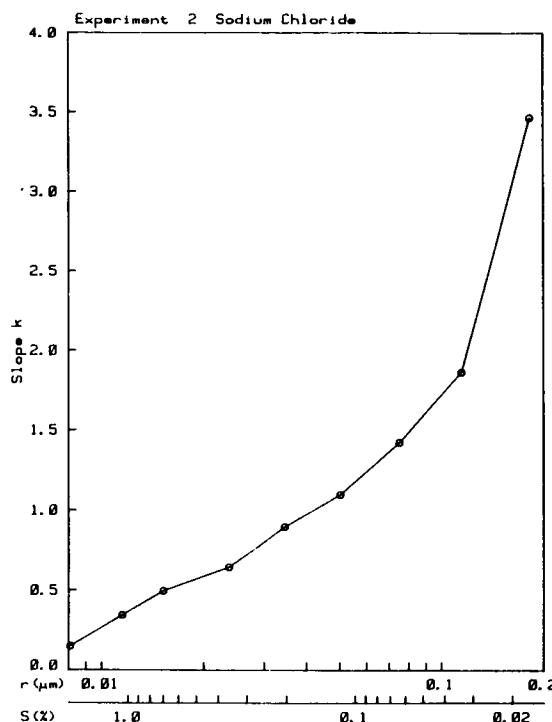


Figure 1. Slope k derived from mobility analyzer data as a function of dry particle radius (critical supersaturation computed). In this and subsequent figures, each plotted point typically represents the average over the interval bounded by immediately adjacent points.

ciated k values. The results are shown in Figures 2-6 for laboratory and ambient aerosols. Again the solid line represents k values constructed from the NRL MA particle size distributions. Superimposed are measured k values as obtained with the four indicated CCN chambers. Some observations are as follows:

With the possible exception of Experiment 17 (Figure 6), the agreement between theoretical and measured k values is considered good for $S > 0.2\%$ for both ambient and laboratory aerosols. Agreement is also reasonably good for $S < 0.2\%$ for laboratory aerosols (Figures 2 and 3), but not for ambient air (Figures 4 and 5). In ambient air, the typically higher measured k values for $S < 0.2\%$ may be due to sample time and concentration variations or to inherent accuracy limitations of sparse data. However, it may be that particles larger than $\sim 0.1 \mu\text{m}$ with suitable fractions of soluble material to serve as effective CCN have steeper aerosol distribution slopes than the total (Junge-type) aerosol distribution. Evidence is accumulating (e.g., Meszaros, 1968; Winkler, 1970) to the effect that large particles $> 0.1 \mu\text{m}$ have considerably lower solubility ratios than smaller particles.

In the 0.2 - 1.0% S range, the ambient aerosols tested do seem to possess a reasonably flat k plateau. This generally was not the case in the 0.02 to 0.2% S range.

Experiment 17 (Figure 6) was a case in which the ambient aerosol concentration fluctuated greatly with time. A good deal of the scatter in the

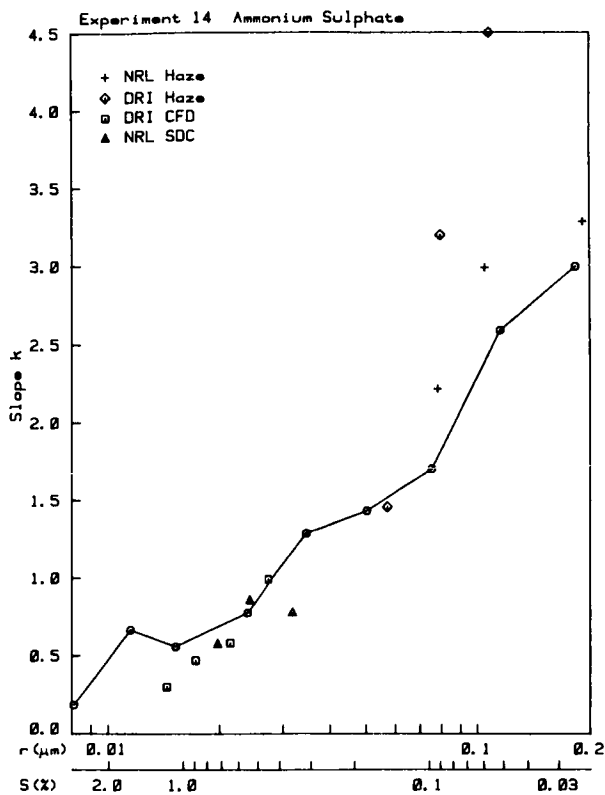


Figure 2. Comparison of k values from CCN spectra (four instruments) and from mobility analyzer derived slopes (solid line) -- laboratory aerosol.

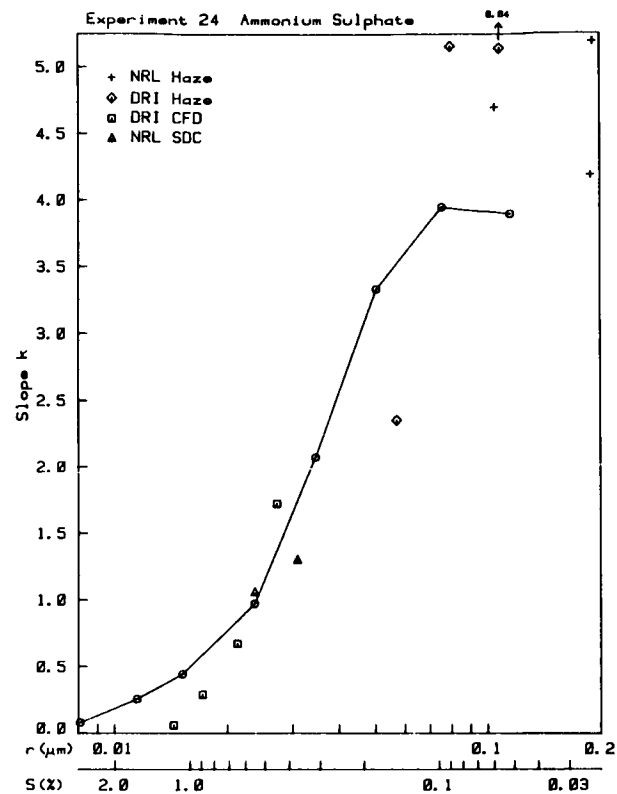


Figure 3. Comparison of k values from CCN spectra (four instruments) and from mobility analyzer derived slopes (solid line) -- laboratory aerosol.

TABLE 2
CCN Supersaturation Slopes k^*
Instrument Number and Type

Experiment No.	Aerosol	NRL MA+	SDC			CFD		Haze			
			10	13	17	15	21	NRL MA+	11	14	21
2	NaCl	0.69	--	0.50	0.49	--	0.79	2.48	1.78	2.57	1.63
14	(NH ₄) ₂ SO ₄	0.89	0.63	0.71	0.67	0.67	1.0	2.47	2.81	3.31	2.66
23	(NH ₄) ₂ SO ₄	0.96	0.66	1.49	1.07	0.93	1.33	2.62	4.80	4.61	4.16
23	(NH ₄) ₂ SO ₄	1.21	0.90	1.01	1.01	0.93	1.60	3.91	4.82	5.22	3.85
6	Ambient	0.83	0.75	0.74	0.85	0.91	--	2.28	3.08	3.76	1.83
11	Ambient	0.62	--	0.37	0.39	0.44	0.51	2.02	2.60	2.29	3.80
17	Ambient	0.93	1.08	0.83	0.63	1.27	2.32	1.95	2.98	2.14	2.71

* For SDC and CFD Chambers - S range from ~0.2 to 1.5%
For Haze Chambers - S range from ~0.01 to 0.2%

+NRL Mobility Analyzer (MA), sometimes referred to in other Workshop data as the NRL Electrical Classifier (EC).

data can undoubtedly be attributed to this factor. Under such circumstances, CCN instruments with short time constants are distinctly advantageous. Otherwise a storage vessel for "grab samples" should be employed. If the NRL mobility analyzer curve is truly representative (and not also influenced by temporal aerosol variations), it is evident that a single k value in the 0.1-1% S range would be questionable.

4. CONCLUSIONS

Much more is known about CCN concentration variations than about the shape (k) of the CCN supersaturation spectrum. While experiments at the Workshop were not specifically designed to focus on k values, some relevant information was obtained as a byproduct. Some preliminary insights gained were as follows:

a. From Köhler theory and ambient aerosol distributions, it is predicted that $k = 2\beta/3$, where β is the slope of the (soluble) total aerosol size distribution. Workshop results tended to verify this relation.

b. Over the typical S range (0.2 to 1%) of many CCN counters (SDC's and CFD's), the CCN spectrum slope in the ambient air cases was generally uniform; such is necessary for applicability of a single-valued power function of the form $N = cS^k$. A uniform k does not hold for the laboratory generated aerosols, and is not always an accurate approximation in ambient air.

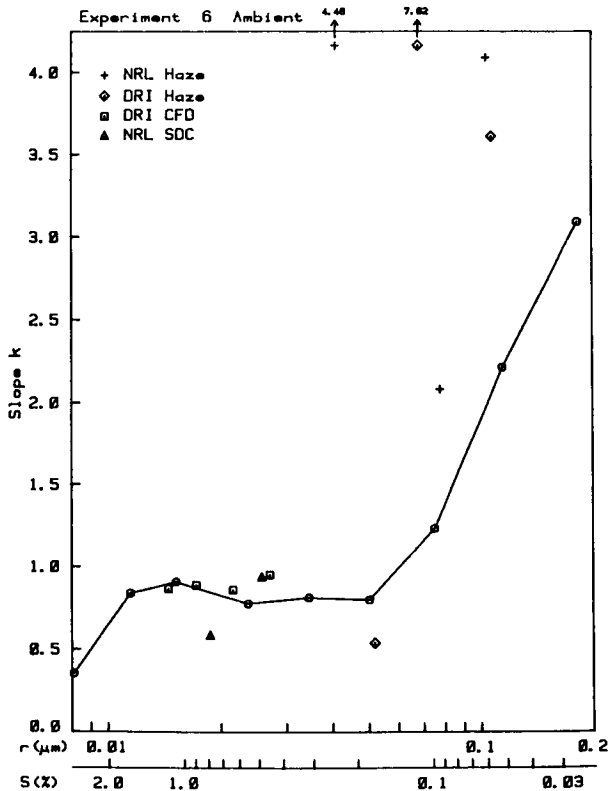


Figure 4. Comparison of k values from CCN spectra (four instruments) and from mobility analyzer derived slopes (solid line) -- ambient aerosol.

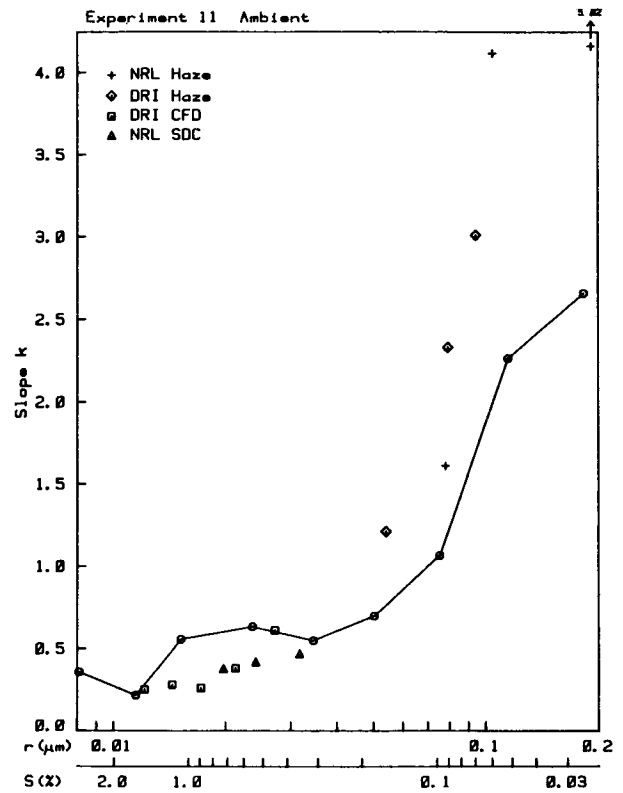


Figure 5. Comparison of k values from CCN spectra (four instruments) and from mobility analyzer derived slopes (solid line) -- ambient aerosol.

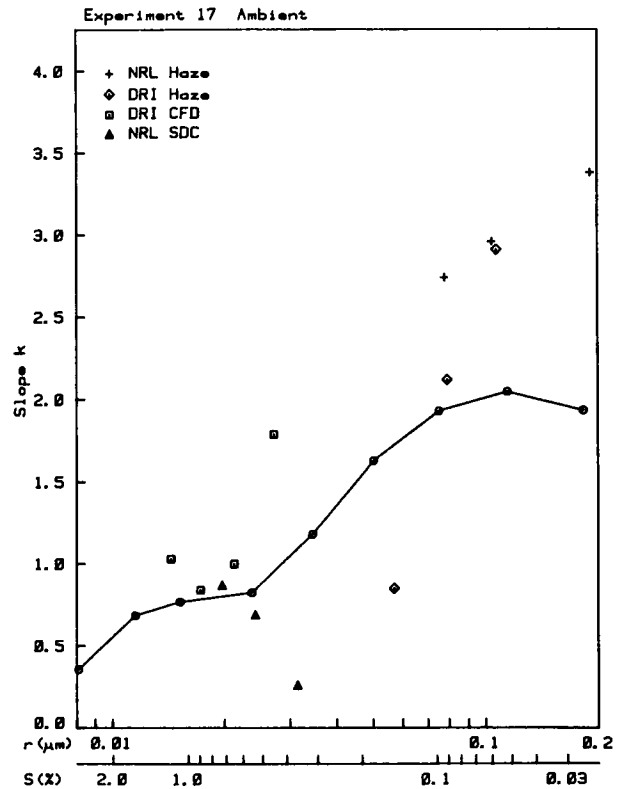


Figure 6. Comparison of k values from CCN spectra (four instruments) and from mobility analyzer derived slopes (solid line) -- ambient aerosol.

d. At supersaturations from ~0.01-0.2%, k values were much larger and seemingly less constant over the S range of interest. Future measurements will help determine whether a single power fit is adequate in this range.

e. The measured k values for ambient air in this low S range were typically larger than those calculated from an assumed soluble aerosol size distribution. This may merely reflect insufficient data and lack of suitable experimental control. However, another hypothesis to examine is whether these larger CCN particles, with suitable effective solubility ratios, decrease in number more rapidly with size than the total aerosol distribution.

f. CCN concentrations can change quite rapidly with time in ambient air. Unless fast-response CCN counters are used or large point-sample containers employed, erroneous k values and power functions can result.

5. ACKNOWLEDGEMENTS

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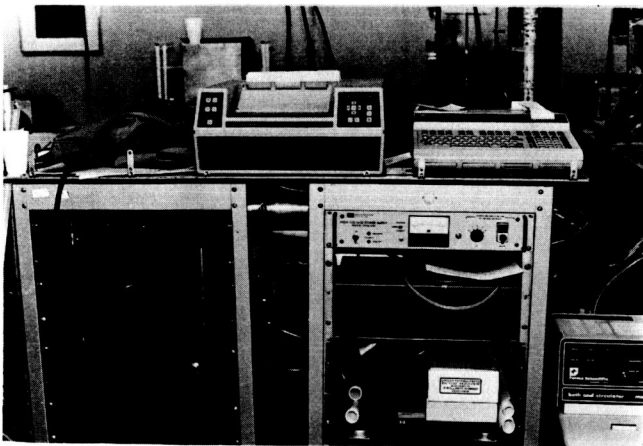
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MEASUREMENT OF THE AEROSOL SIZE DISTRIBUTION
WITH NRL'S MOBILITY ANALYZER

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ABSTRACT

The size distribution of the aerosol sample generated at the International CCN Workshop was measured with the NRL mobility analyzer/size spectrometer in the size range between 0.0057 and 0.57 μm radius. A description of the instrumentation and data analysis is given, together with the measured size distributions calculated for each of 23 experiments carried out at the Workshop.

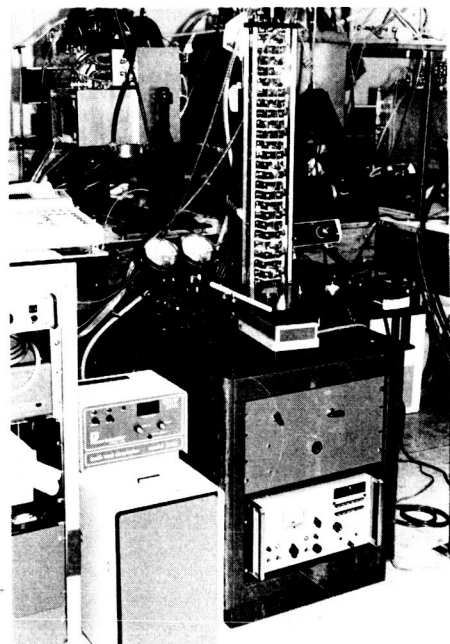


The size distribution of aerosol particles between 0.0057 μm and 0.57 μm radius was measured with NRL's mobility analyzer. The basic instrument is shown in Figure 1. The aerosol sample is first brought to charge equilibrium by passing the sample through a region of bipolar ionization. The sample air enters the mobility analyzer through a slit in the outer cylindrical wall where it is confined to a thin laminar layer along the wall by filtered sheath air. A small amount of filtered air is extracted through a slit in the inner electrode. When a voltage is applied to the inner electrode, those particles which are charged and lie in a narrow mobility range will be withdrawn with the extracted air. By measuring the particle concentration in the extracted air as a function of voltage, the mobility distribution of the charged fraction can be obtained. From the mobility distribution of the charged fraction (of one polarity) and the equilibrium (Boltzmann) charge distribution, the size distribution can be calculated. The theory of the analyzer and the analytical procedure for obtaining the size distribution has been given by Hoppel (1978).

The concentration of particles in the extracted air is much lower than in the sample air. The first data obtained with the mobility analyzer was taken by measuring the concentration in the extracted air with a Pollak counter. Although the long-tube Pollak counter is probably the most sensitive

of the light scattering CN counters, its use imposed rather severe limitations on the sensitivity of the system. Useful size distributions could be obtained in atmospheres which had high concentrations of particles but when the total count dropped below about 1000 particles per cm^3 the system could not be used. Even at higher concentrations the sensitivity was not adequate to detect particles in all size channels over the specified range.

In an attempt to increase the sensitivity of the system, a single particle condensation nucleus counter was developed at NRL, which now allows much greater sensitivity and has made it possible to automate the system. While this new continuous particle counter is currently used as part of the mobility analyzer system, it is novel and will possibly have other applications in regard to CCN and CN. Figure 2 shows the design of the segmented thermal gradient CN counter for single particle counting. The sample air enters along the axis of the chamber and is surrounded by filtered, humidified sheath air. The walls are water saturated with alternating segments maintained at alternating temperatures. The basic principle is similar to that employed in the TGDC where a supersaturation exists between horizontally oriented hot and cold plates with saturated walls. The TGDC is limited to measuring particles activated in a narrow range of supersaturations between about 0.2% and 1.0% due to well-documented reasons related to fall-out during the time required to establish the equilibrium supersaturation in the TGDC. The advantage of the



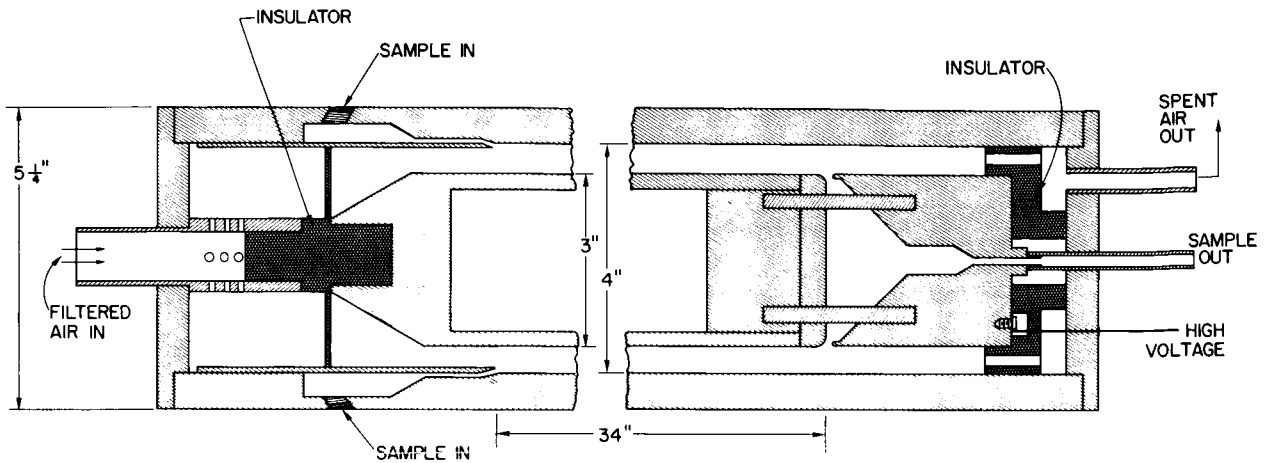


Figure 1. Diagram of NRL Mobility Analyzer.

segmented chamber is that particles falling vertically are not lost and the segmented geometry suppresses large scale convective motions which would occur between vertically oriented hot and cold walls. The criteria in designing the segmented chamber is that the air flow rate must be such that the residence time in a single segment is short compared to the time required for moisture or heat to diffuse in from the walls to the center; whereas the residence time of the sample in the entire chamber must be longer than the diffusion time. When this criterion is met, the air along the axis will reach a temperature and vapor pressure which is approximately midway between the temperature and vapor pressure of each segment. Along the axis the asymptotic supersaturation will be the same as exists in the center of a TGDC working at the same temperature difference. Figure 3 shows the development of the supersaturation within the chamber. Details of the design and analysis of the chamber are given by Hoppel, Twomey, and Wojciechowski (1979, 1980). The particles are nucleated and grown to optically detectable sizes in the segmented chamber and then transmitted directly into an optical counter for single particle counting. When the segmented chamber is used with the mobility analyzer, a small portion of the output sample from the mobility analyzer is passed from the analyzer to the segmented chamber for counting. At the Workshop, the chamber was operated at a temperature difference of about 20°C which results in a supersaturation of about 20 percent.

The system (mobility analyzer and segmented chamber) has been operated using either 21 or 11 size channels to cover the specified range. After considerable experience with atmospheric measurements, it was decided to standardize to 11 channels when the system was automated. The reason for using fewer channels is that it takes about 18 minutes to cover 11 channels and about twice that long for 21 channels. In the atmosphere the increased resolution offered by 21 channels is less important than making the measurement as quickly as possible because of temporal changes in concentrations. On those occasions at the CCN Workshop when very stable nearly monodisperse particles were generated, it would have been helpful to have had the added resolution of 21 channels.

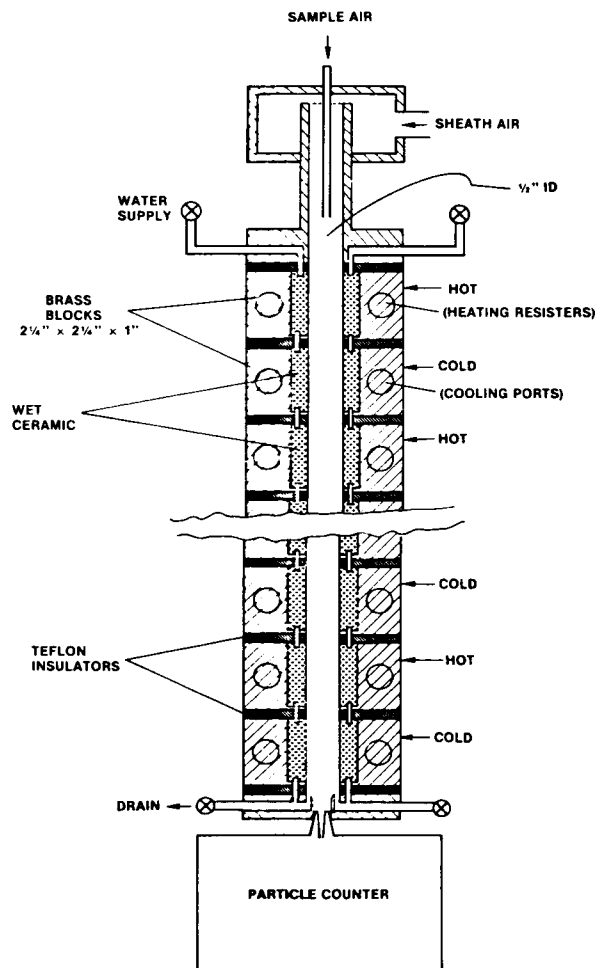


Figure 2. Design of NRL segmented CN counter for single particle counting.

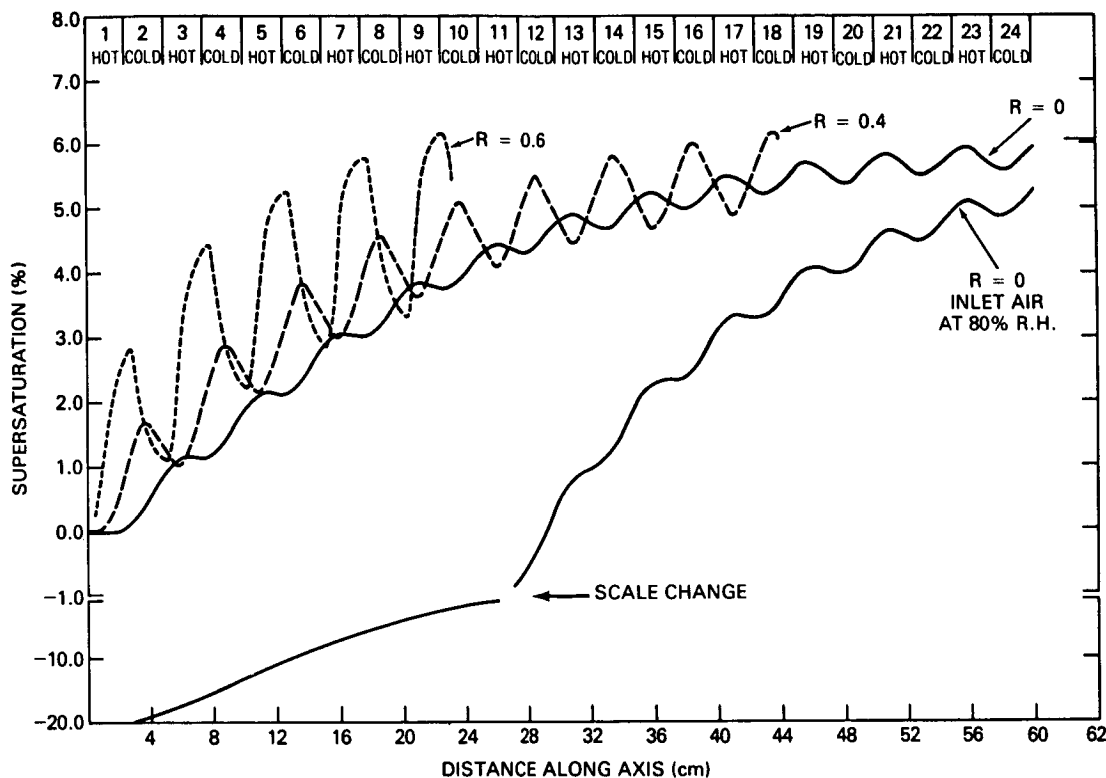


Figure 3. Supersaturation within segmented CN counter.

The system is automated with the aid of Hewlett Packard 9825 desktop computer with a real time clock. The voltage is changed every 84 seconds by a 16 bit word written to a 16 bit parallel port which controls the output of the high voltage power supply. Sixty seconds are allowed for flushing and stabilization at which time the single particle counter (Royco) is reset and two 12 second counts are recorded, averaged, and stored. The Royco is controlled through a binary port and read with a BCD interface. At the end of the cycle (12 voltage steps including zero volt background count) the computer calculates the mobility distribution and size distribution by an iterative procedure (given by Hoppel, 1978) which corrects for the effect of multiply charged particles according to the Boltzmann charge distribution. A computer controlled plotter then produces real time plots of the differential and cumulative size distribution. These size distributions were available immediately after each experiment. Usually several (two to four) individual size distributions were taken during each experiment.

After the Workshop, the raw data from the individual runs were examined and averaged. An average size distribution was then calculated for each experiment for which data was available. It is believed that the averaged size distribution is preferred to the individual distribution in all experiments for the following reasons: (1) temporal variations in the concentration may occur between voltage steps and this will distort the size distribution. The more data there is to be averaged, the less will be the distortion due to temporal variations; (2) even if the temporal variations are small, variations due to counting statistics of the samples are reduced by averaging; (3) there are occasions when the high supersaturation in the segmented chamber results in the formation of a drop-

let at the mouth of the Royco. When this droplet is drawn through the Royco a large spurious count is encountered. This spurious count is easily recognized by examining the count in the different Royco channels recorded as raw data. The spurious reading caused by the droplet results in a large disparity between channels which is easily detected. When the system is working correctly, the count is nearly the same in all Royco channels. In the averaged data, the spurious points are removed before averaging.

The size distributions calculated from the averaged data for each experiment are shown in the 23 figures given at the end of this contribution.

Experiments 27 and 28 were for silver iodide aerosol. The accuracy of the size distribution for these experiments cannot be trusted but the distributions are included for sake of completeness. Silver iodide is quite insoluble and therefore many of the particles may not have been nucleated at the 20% supersaturation in the segmented chamber. Some compensation was made by increasing the supersaturation during these runs by an unrecorded amount. The raw data from the optical counter did not behave normally in that the number of particles was not the same in all channels, i.e., the particles did not appear to grow in the manner expected for nucleated droplets.

It should be pointed out that the largest channel is uncorrected for multiple charging. In order to correct a channel for the presence of larger particles with multiple charges, some knowledge of the concentration of larger particles must be available. It is therefore impossible to make this correction in the largest channel on the basis of analyzer data. However, if the size distribution is decreasing very rapidly at the largest size

(as is nearly always the case), this correction will be small. Since the largest channel is uncorrected it must be viewed with suspicion.

Also, when the cumulative count drops below a few particles per cc, the count cannot be trusted because this corresponds to a raw count which is close to background levels. (This is not an inherent lower limit since this lower limit can be easily decreased further by increasing the dilution ratio in the segmented chamber. The dilution ratio is presently about 1 to 15.)

The accuracy with which the boundaries of the size channels are determined is more easily evaluated than the accuracy with which the total number of particles in that channel can be evaluated. The sizing accuracy is related to the accuracy with which the mobilities are determined. The midpoint mobility of an interval is given by

$$k_1 = \frac{\phi_1}{4\pi CV} \text{ (esu)} = \frac{\epsilon\phi_1}{CV} \text{ (mks)}$$

Where ϕ is the volume flow rate of the sheath air, C the electrical capacity, and V the voltage. The accuracy of C and V are not in question. The airflows were remeasured by the "bubble" method at the altitude of the Workshop and are calibrated to within 3%. The accuracy to which the airflows are set and maintained throughout a given run is about 5%. A 5% error in flow results in a 5% error in mobility. A 5% error in mobility translates into approximately a 5% error in radius at the larger sizes and a 10% error in radius at the smaller sizes. The Stokes-Cunningham-Millikin relationship is used to calculate the radius from mobility. If this relationship introduces no further errors, then the maximum error in the size boundary is certainly less than 15%. It should be pointed out that the radius in the size measurement is the equivalent drag radius which may be slightly different than the "mass" radius required in the Kohler theory.

The accuracy with which the number concentration of particles in a given size channel is known depends on several factors: (1) the detection system, (2) the shape of the size distribution and, (3) the validity of the Boltzmann charge distribution. The nucleated droplets are detected by the Royco particle counter which, in this application, is used as a single particle counter and not as a sizing device. It is generally acknowledged that the absolute counting accuracy of optical counters is much better than their sizing ability. The errors due to counting are believed to be small compared to factors (2) and (3). The method of extracting the mobility distribution assumes that the distribution is nearly linear across any given channel. This assumption is not justified in the case when the distribution is strongly peaked in a single channel as was the case for many of the monodisperse experiments at the Workshop. Since the transmission function is the greatest at the center of the interval and drops to zero at each extreme, it would appear that if the peak in the monodisperse size distribution occurs at the center of one of the (preset) channels, then the total number of particles would be overestimated, whereas if the peak occurs near the boundary of the channel, the total number would be underestimated. Below a radius of about 0.02 μm the validity of the

Boltzmann distribution is in doubt. There are experimental results which indicate that the Boltzmann law hold down to 0.01 μm (Lui and Pui, 1974 and Servaas and Krider, 1977), but on the other hand, there is theoretical justification to indicate that the real distribution departs from Boltzmann below about 0.03 μm and that the ratio of charged to uncharged particles is larger than predicted by the Boltzmann law.

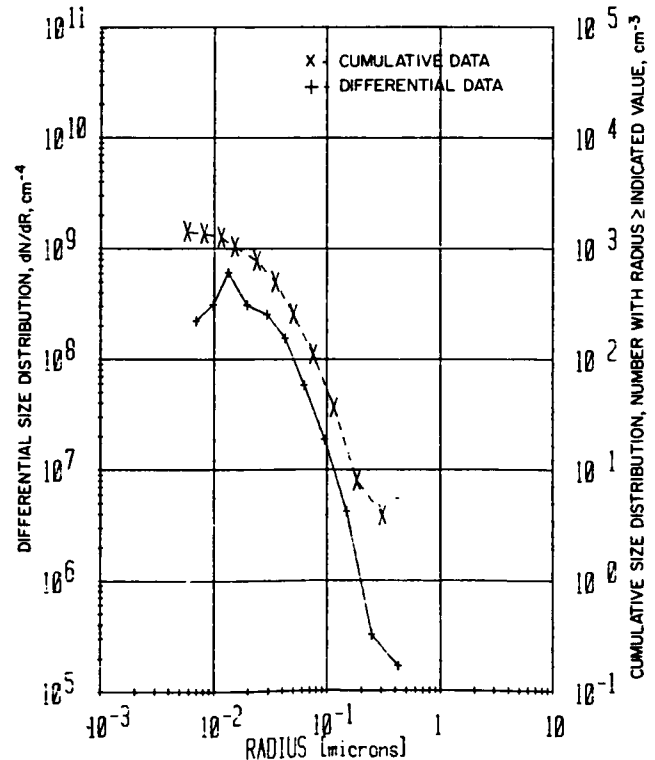
As a result of these three factors, it is difficult to quantify the accuracy of the total numbers of particles measured with the mobility analyzer. However, we believe that the second order mobility analyzer is the most accurate method of measuring the size distribution presently available.

Acknowledgements

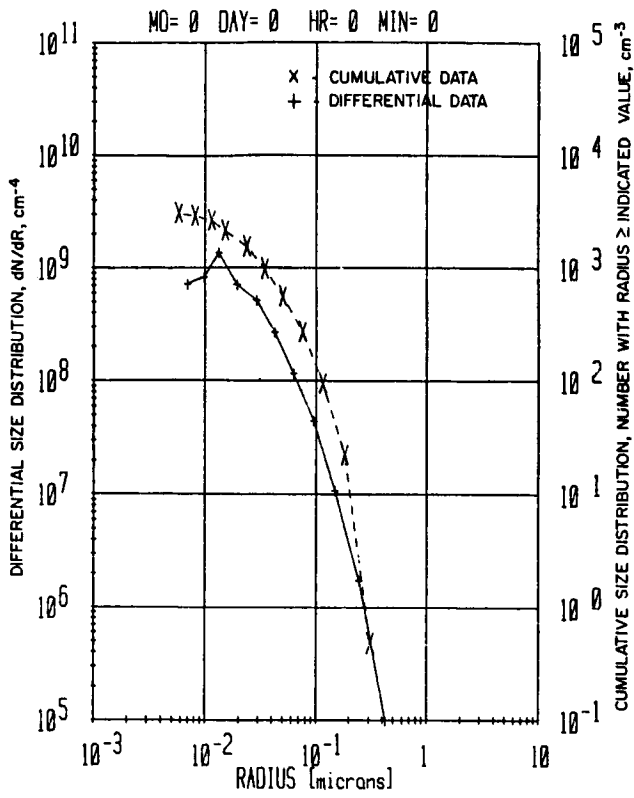
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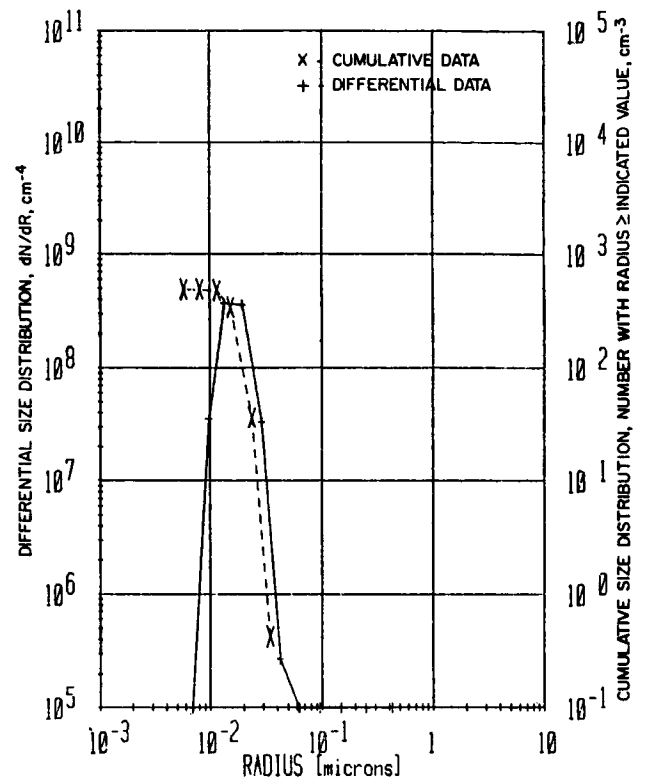
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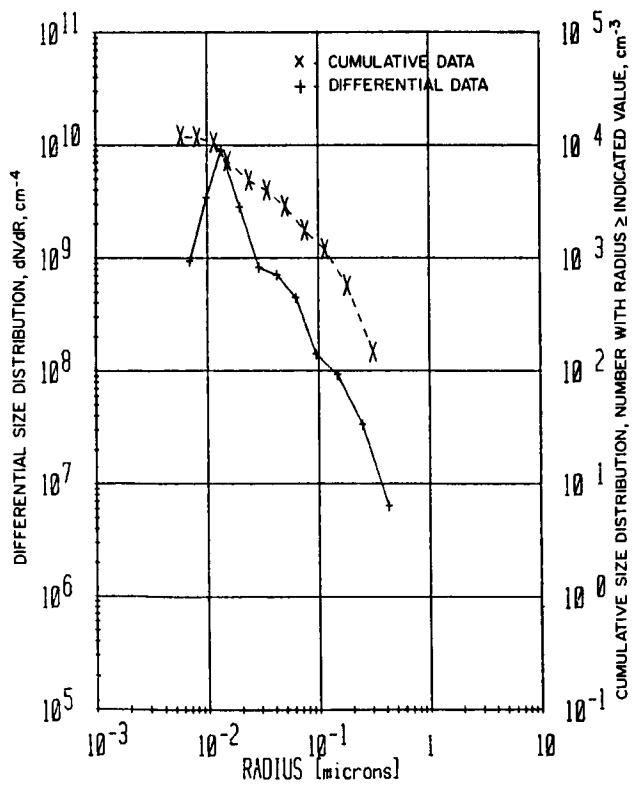
Experiment #1 - Average of 3 Runs



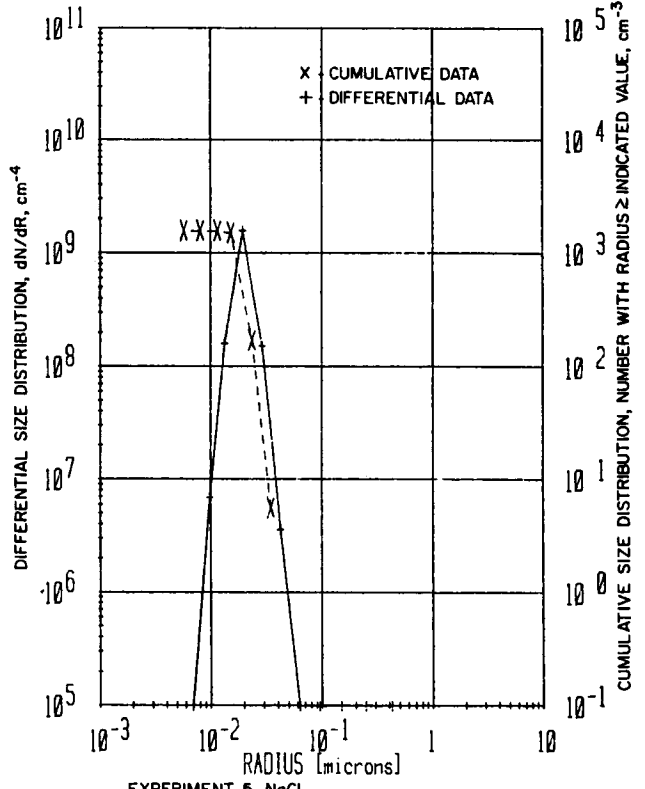
Experiment #2 - Average of 2 Runs



Experiment #4 - Average of 4 Runs

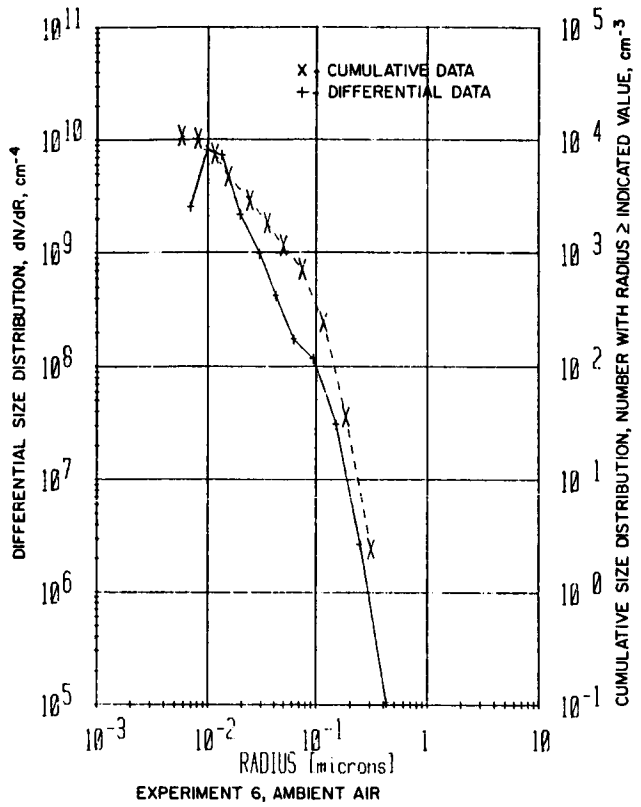


Experiment #3 - One Run Only

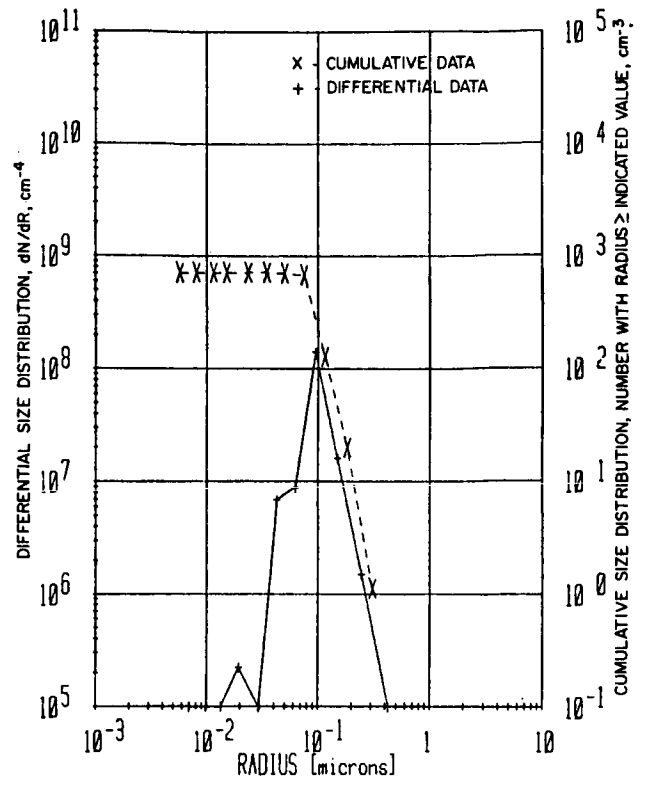


EXPERIMENT 5, NaCl

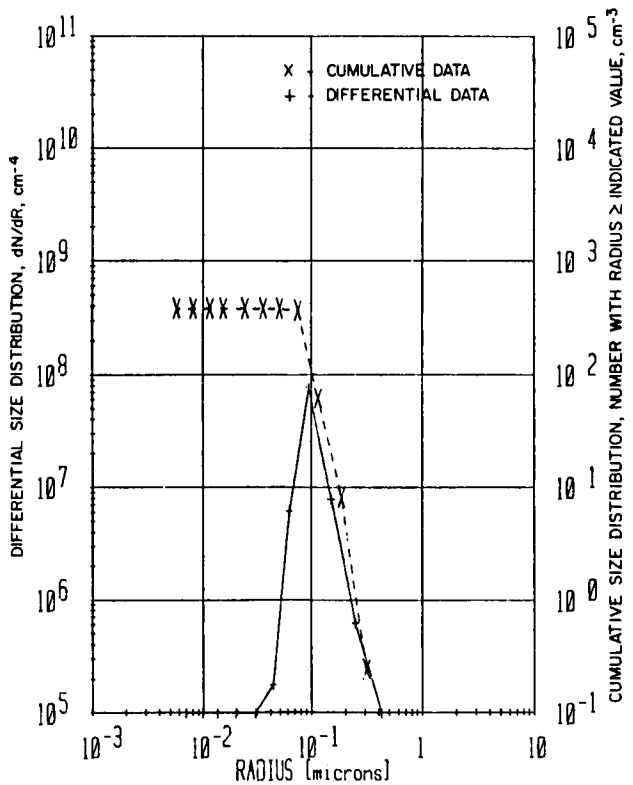
Experiment #5 - Average of 3 Runs



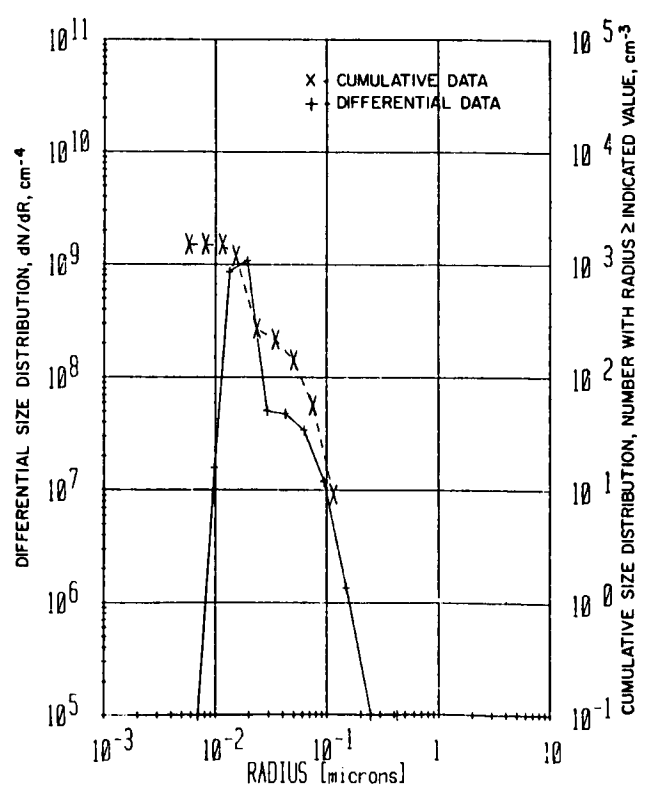
Experiment #6 - Average of 3 Runs



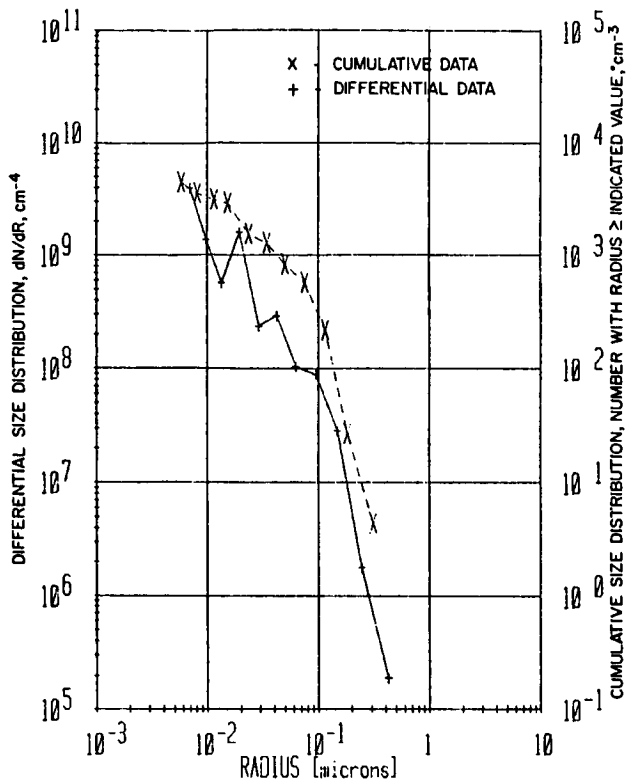
Experiment #9 - Average of 3 Runs



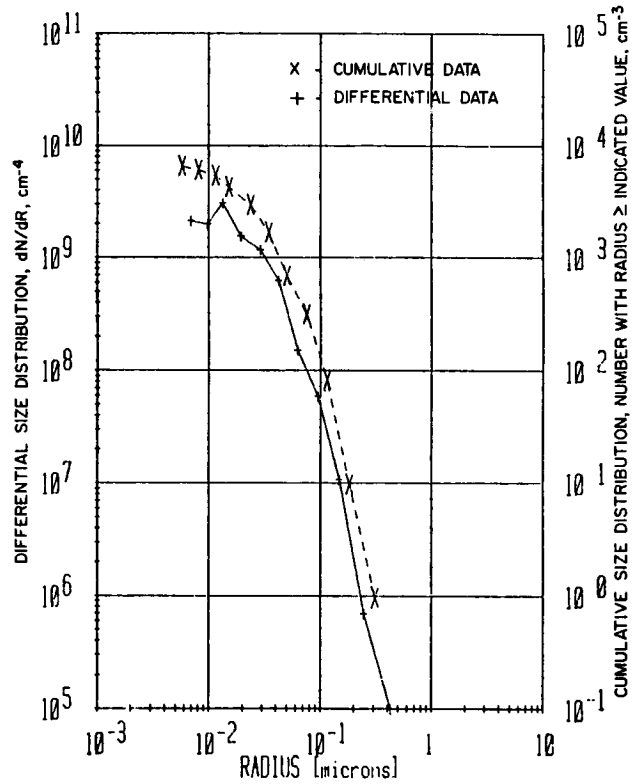
Experiment #8 - Average of 2 Runs



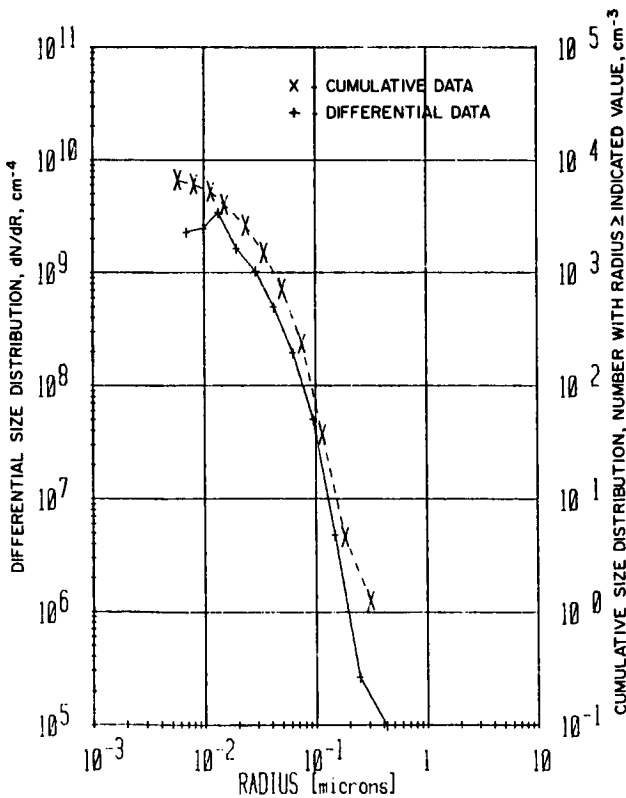
EXPERIMENT 10, BIMODAL NaCl
Experiment #10 - Average of 2 Runs



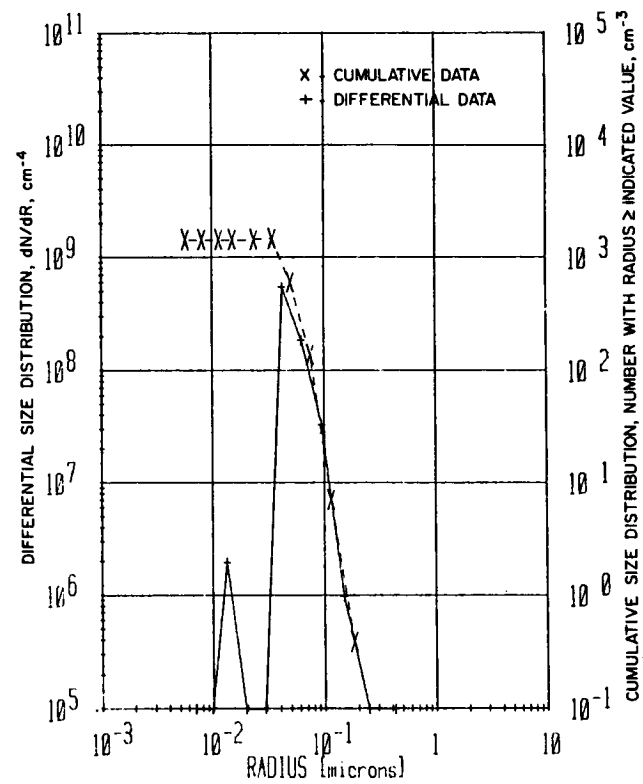
Experiment #11 - Average of 3 Runs



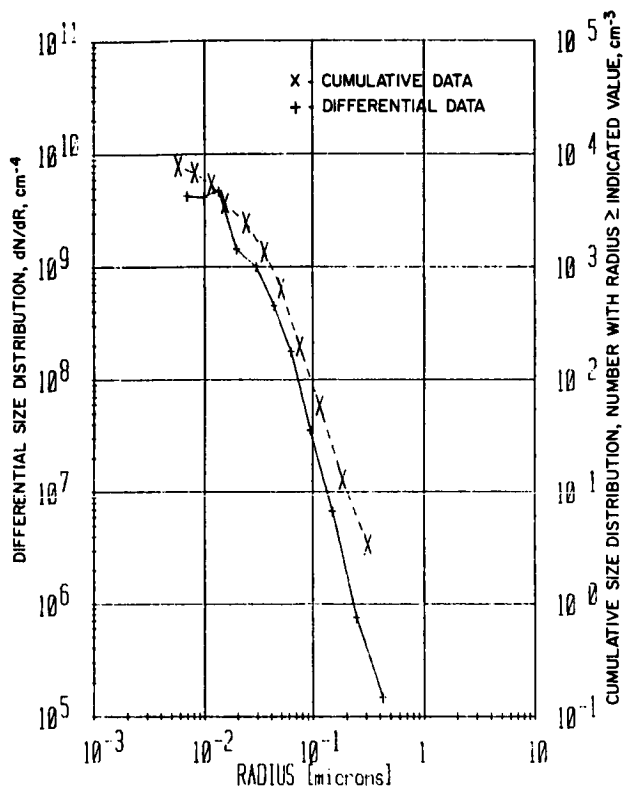
Experiment #14 - Average of 3 Runs



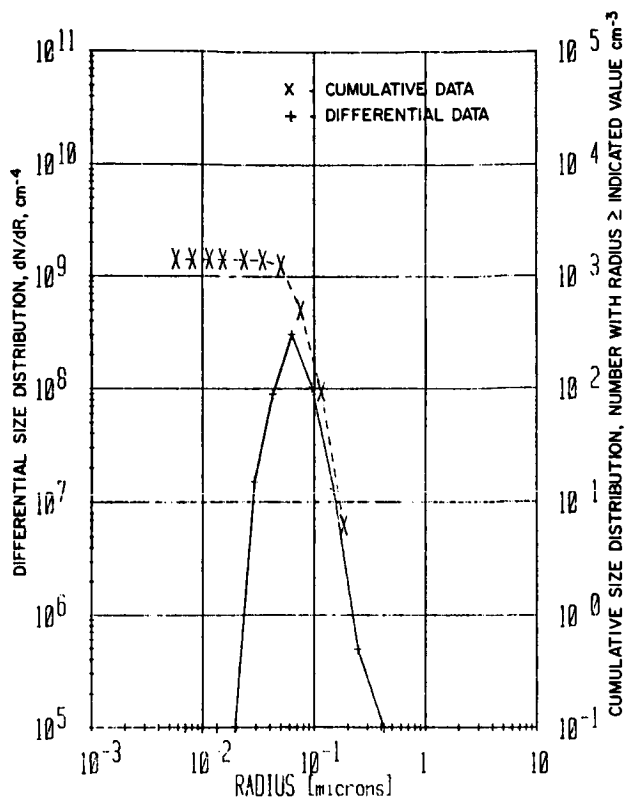
EXPERIMENT 13, POLYDISPERSE $(NH_4)_2 SO_4$
Experiment #13 - Average of 3 Runs



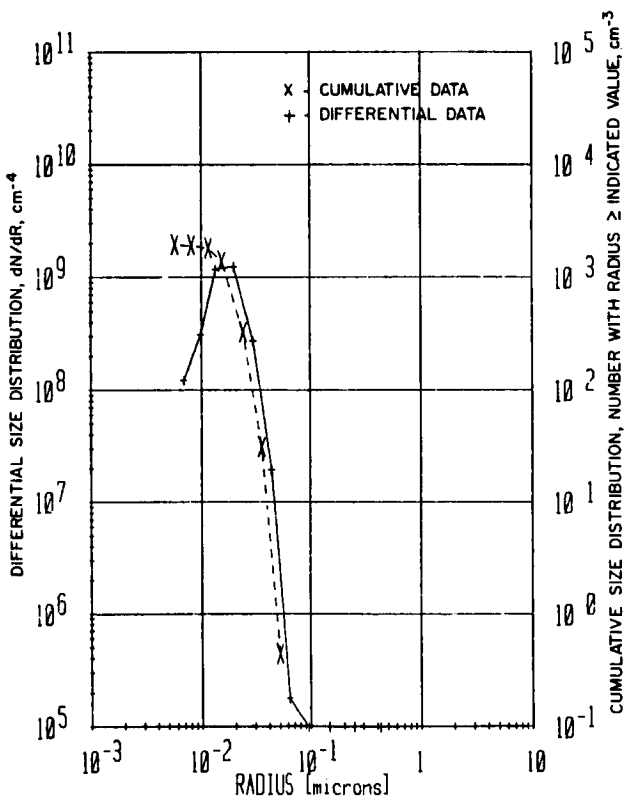
EXPERIMENT 15, $(NH_4)_2 SO_4$
Experiment #15 - Average of 2 Runs



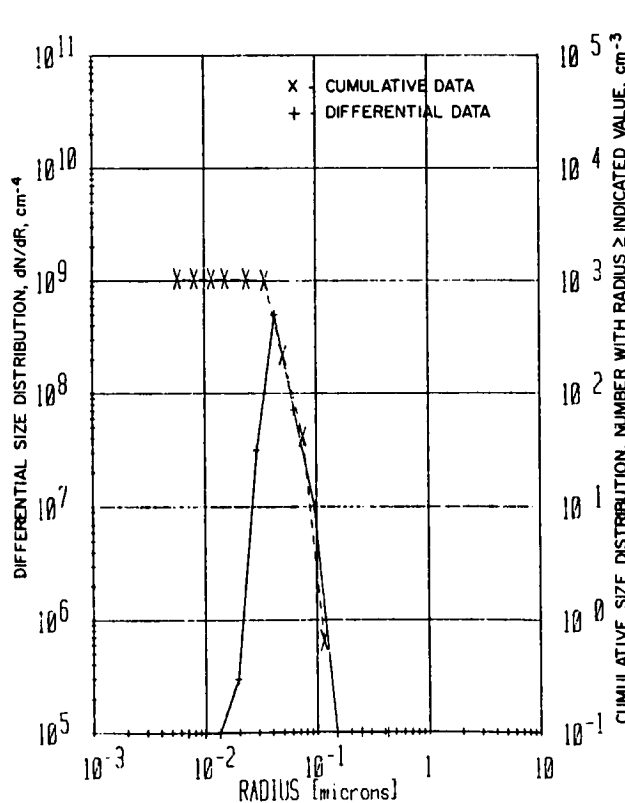
Experiment #17 - Average of 2 Runs



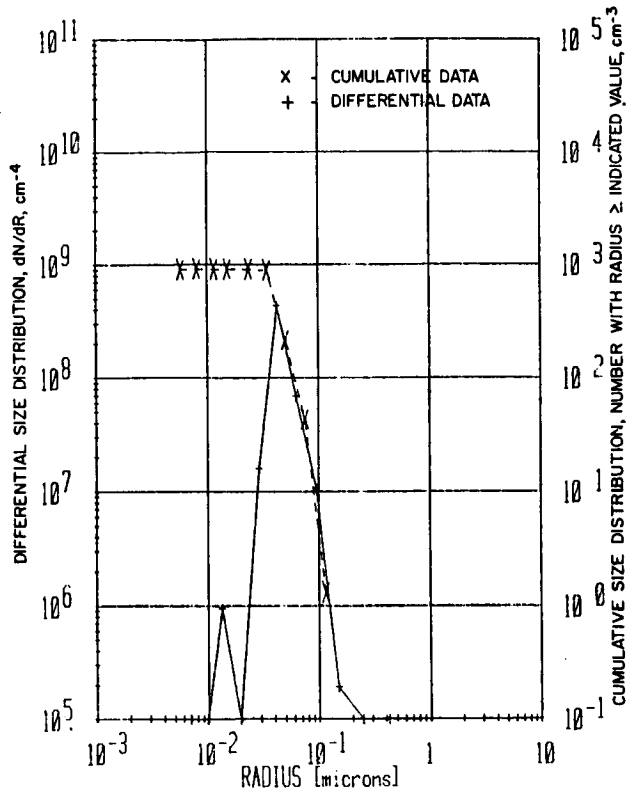
Experiment #19 - Average of 4 Runs



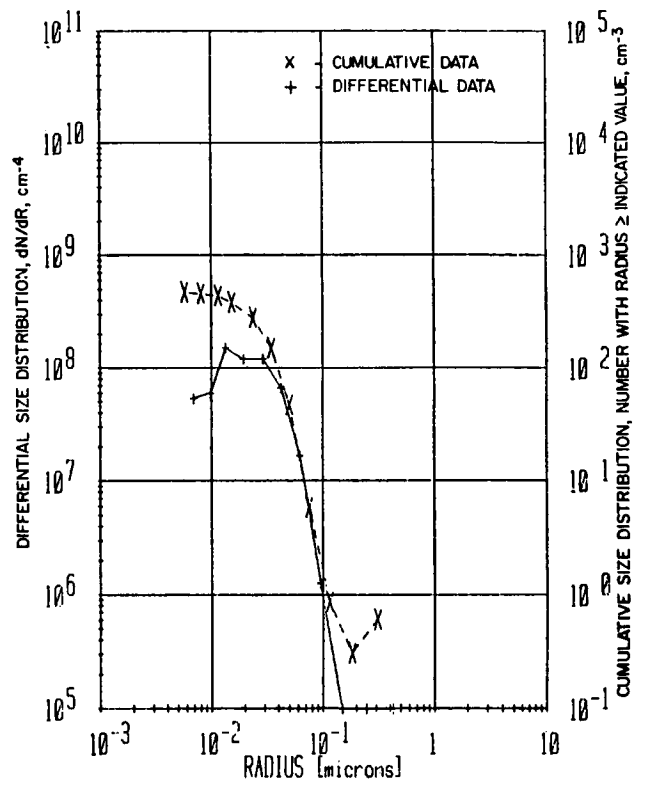
Experiment #18 - Average of 3 Runs



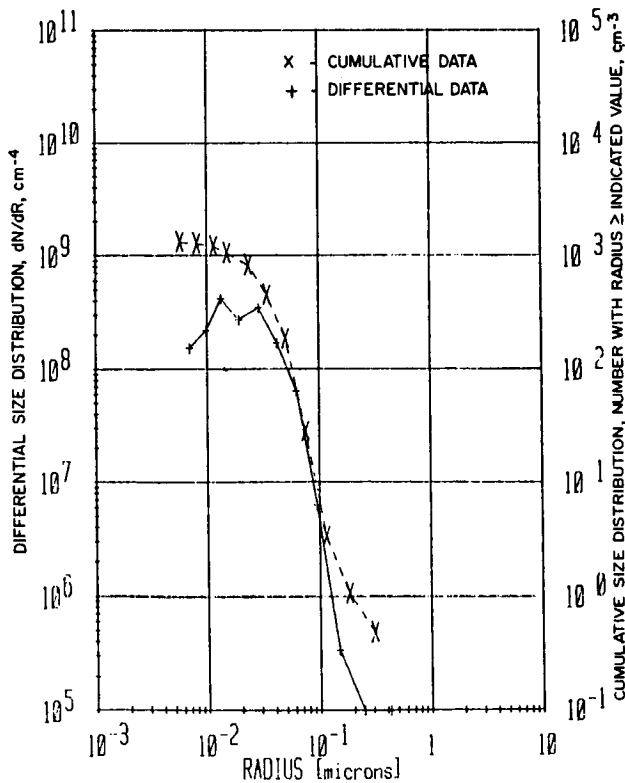
Experiment #20 - Average of 4 Runs



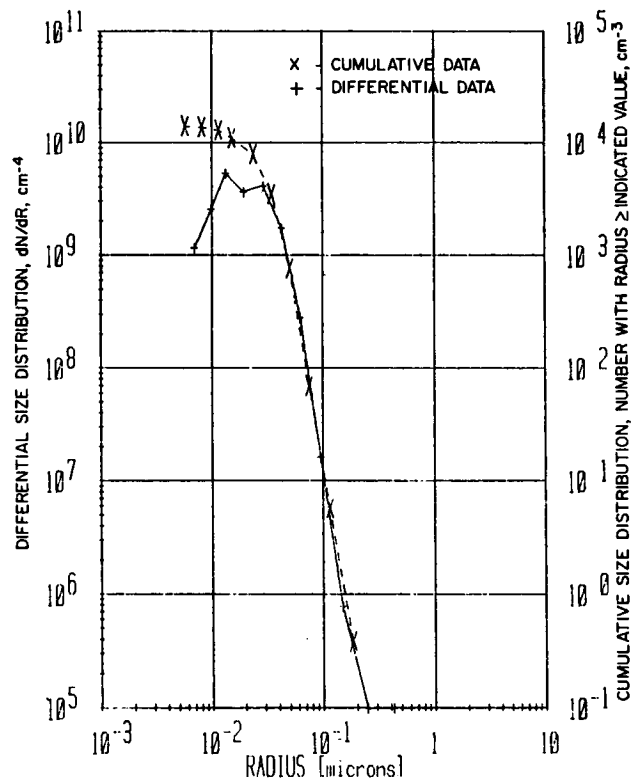
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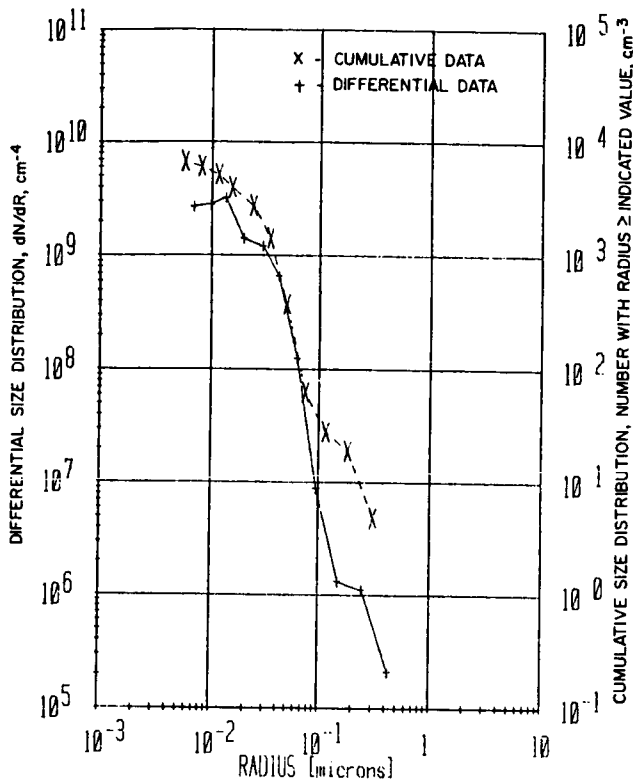
Experiment #23 - Average of 4 Runs



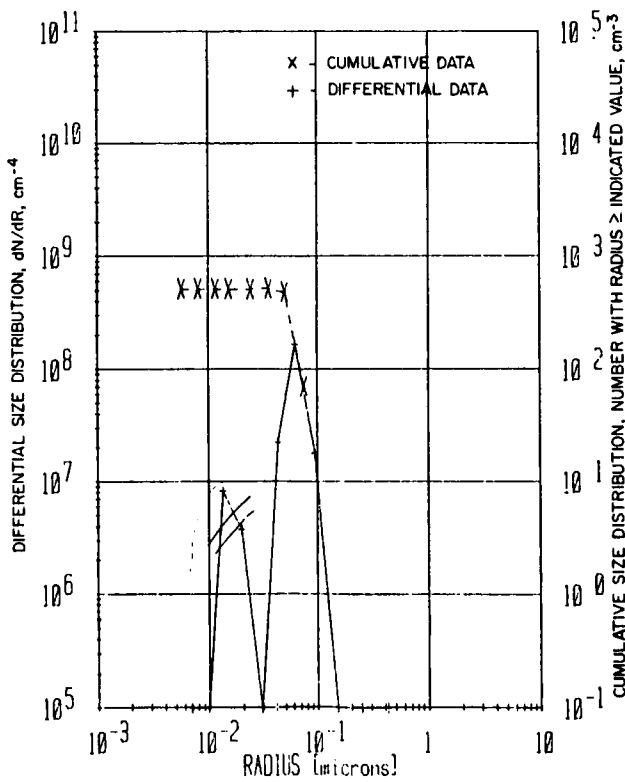
Experiment #22 - Average of 3 Runs (2,3 and 4)
[taken between 1100 and 1220; early run (#1)
was taken when the count was higher
and remains as distributed at Workshop].



Experiment #24 - Average of 4 Runs



Experiment #27 - Average of 4 Runs



Experiment #28 - Average of 2 Runs

PERFORMANCE OF THE CONTINUOUS FLOW DIFFUSION CHAMBERS

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ABSTRACT

A brief comparative description is made of the five continuous flow chambers which participated in the Workshop. Overall, comparisons for the various types of experiments - monodisperse, polydisperse and ambient aerosol - showed agreement among these chambers to within 15% in most cases. A careful analysis of the results indicated that a proper accounting of certain parameters would bring about much closer agreement among four of these instruments.

1. COMPARATIVE DESCRIPTION OF INSTRUMENTS

There were five continuous flow diffusion chambers in the Workshop. Three of these were from the Desert Research Institute (DRI), one was from the University of Missouri-Rolla (UMR), and the other was from the University of Washington. One of the DRI chambers was the conventional continuous flow diffusion (CFD) chamber, designated C. The second DRI chamber was the rapid cycle spectrometer built for NASA, designated N. The third DRI chamber was the instantaneous spectrometer, designated I. The UMR instrument acted in a dual capacity since it also was used as an isothermal haze chamber. This article will be restricted to analysis of its operation in the CFD mode. The designations M and W will, respectively, denote the UMR and University of Washington instruments.

All of these chambers moved samples of aerosol through parallel plate diffusion chambers after which the drops which formed were detected by optical particle counters. The parallel plates in all five chambers were vertically oriented although the original design of CFD chambers called for the plates to be horizontal to suppress convection. However, the widespread acceptance of vertical plates seems to indicate that this convection is not a serious problem. The direction of the sample flow was not the same in all chambers since the sample moves horizontally in the three DRI chambers (C, N and I) and vertically downward in the M and W chambers. In all five chambers, the sample was confined to a plane midway between the plates by particle-free air. The sample was spread out into a lamina in the plane midway between the plates for all chambers except UMR where the sample was an axial stream. The sample was thus confined in these chambers to keep it in the zone of maximum supersaturation which is approximately midway between the plates. In the DRI and UMR chambers, all of the sample which entered the chambers was detected by the optical particle counters (OPC's), whereas the Washington chamber detected only a small fraction of the sample which was carried through a central tube to an optical box. In all of the chambers, the time of exposure to each

supersaturation was adjusted for maximum performance by using suitable lengths of the wet zones and by adjusting the total flows. The times used in each of the chambers seemed to be consistent when similar supersaturations were considered. The wet surfaces were filter paper for the C and M chambers, metal for the N and I chambers, and felt for the W CFD.

Plate temperatures in the C and I chambers were controlled by water baths whereas the W chamber had refrigeration coils in direct contact with the cold plate. The warm plate of the W chamber was heated by the exhaust gases from the refrigerator along one edge of the warm plate. The N chamber also used circulating fluid but the regulation of that fluid was done with computer-controlled thermoelectric modules.

The M plate temperatures were monitored by checking the reservoir temperatures with mercury thermometers. The C and I CFD's used thermistors imbedded in the plates whereas the N CFD used a thermopile to measure ΔT . The W chamber also used thermistors to monitor the plate temperatures.

The M, C and N chambers obtained spectra by changing plate temperatures. The former two were changed manually while the N CFD changed temperatures according to a prearranged computer program. The I and W chambers did not normally change plate temperatures. The I chamber simultaneously maintained three supersaturations by using three chambers in series and using the droplet distributions from one OPC to deduce the concentrations for three supersaturations. The W instrument had four chambers in parallel and four detectors to monitor the concentrations at four supersaturations.

The DRI chambers (C, I and N) all used Royco 225 optical particle counters while the M chamber used a Climet 201 OPC. The W chamber used four non-commercial OPC's using a single laser for illumination and four photodiodes for detectors. This system does not give as much size discrimination as the commercial OPC's, but accurate sizing is not usually necessary at the higher supersaturations where the W instrument was normally operated.

2. PARTICIPATION AND EXPERIMENT TYPES

Of the 29 experiments, five did not yield data suitable for comparison purposes. Experiments 3, 7 and 17 were ambient aerosols which were not constant enough in time to permit useful comparisons. Experiment 25 was a noise test which all of the participating CFD's passed. Experiment 29 was a truly insoluble aerosol (paraffin) which did not show activated droplets in any of the continuous chambers.

Of the 24 useful experiments, the M instrument participated in all except Experiment 21 where its haze chamber mode was used exclusively. The C chamber was involved in all 24 experiments. The DRI spectrometers (N and I) participated in 20 of the 24 experiments as they missed the first 4 (No. 1,2,4 and 5). The W instrument participated in 10 experiments.

The most important division of the experiments is between the monodisperse aerosols on the one hand and the polydisperse and ambient aerosol experiments on the other hand. These two distinct types of aerosols require separate analysis. One conventional way of describing the CCN spectra of aerosols is by noting the concentration, C, at one reference supersaturation (e.g. 1%) and the slope, K, of the distribution. Such a characterization is not applicable to some of the Workshop aerosols, because the slope of the distribution sometimes varied greatly over the range of supersaturations.

3. MONODISPERSE AEROSOL EXPERIMENTS

A monodisperse aerosol is an extreme CCN spectrum because there is a very steep slope over a narrow range of S and a flat distribution over other parts of the spectrum. Hence, this type of spectrum would not be characterized very well by C and K. Instead, a monodisperse aerosol should be characterized by the total number of drops active on the plateau and by the supersaturation (S_C) in the very steep region.

Experiments with a monodisperse aerosol allow a determination of the counting ability of the instruments because, as long as the chamber can attain a supersaturation greater than S_C , the concentration should be constant. Thus this concentration is insensitive to supersaturation levels in the chambers. In fact there should be agreement between the CFD's and Aitken particle counters for monodisperse aerosols. On the other hand, S_C , the second parameter which characterizes the monodisperse aerosols should be a very sensitive test of the supersaturations in the chambers.

3.1 Total Concentration

There were 10 monodisperse experiments, Nos. 4,5,8,9,15,18,19,20,21, and 28. An examination of the experimental plots shows that the M, C, I and N chambers always exhibit a constant concentration for supersaturations above the critical supersaturation, S_C of the aerosol when the chambers operated at high enough supersaturations. The W chamber

observed a flat distribution for only one (No. 9) of the four monodisperse experiments in which it participated.

Table 1 shows the total CCN concentrations for each instrument; that is, the concentrations at and above the flat portion of the distribution beyond which no more nuclei are detected. Also listed is the percentage count compared to the M chamber. At the bottom is the average and standard deviation of the comparison with M. The data in Table 1 is averaged over the entire duration of the experiment. Although the aerosol generator was usually quite constant, this data is subject to error due to the fact that the instruments may have been operating at the high supersaturations during different time periods over which the concentration may have changed.

Table 2 was devised in an attempt to cover the above shortcoming. It was derived by choosing time periods during the monodisperse experiments when there was simultaneous data from at least two chambers. Simultaneous Pollak and TSI Aitken (CN) nucleus data is also presented. These should be in agreement with the CCN counters since these are monodisperse aerosols. Also listed in the table is the percentage difference of each CCN counter from the Pollak counter (or TSI when the Pollak data was not available). The last row shows the average and standard deviation of the comparison with the CN counters. These tables are quite consistent showing that Table 2 is fairly representative of the discrepancies between the chambers. This is in spite of the fact that the data in Table 2 is incomplete and rather unequally distributed. Simultaneous times could not be found for some instruments for some experiments while more than one case could be found for some experiments and instruments. Hence Table 2 is more precise, while the data from Table 1 is more representative.

Table 1 shows that the three DRI instruments were in excellent agreement with each other but that they were about 15% lower than the UMR chamber. Since the DRI instruments are so similar, this would seem to represent a systematic error between the UMR and DRI instruments, especially since the standard deviations for the DRI instruments are about equal. As pointed out above, this discrepancy in concentration for monodisperse aerosol reflects actual differences in counting rather than a problem with supersaturation. Either the sample volume or the counting efficiencies are different or the same sample is not being seen by each chamber.

TABLE 1. MONODISPERSE TOTAL COUNT

Exp. No.	M	C		N	I	W			
		N	cm ⁻³						
4	260	220	85%						
5	1300	950	73%						
8	310	295	95%	260	84%	295			
9	~610	490	79%	540	89%	535			
15	1630	1290	79%	1290	79%	1290			
18	1080	900	83%	920	85%	900			
19	1030	990	96%	990	96%	990			
20	1030	900	87%	910	88%	970			
21	no data	880				880			
28	430	370	80%	330	77%	380			
Averages		85% ± 7%		85% ± 6%		88% ± 6%		100% ± 23%	

TABLE 2. MONODISPERSE TOTAL COUNT, SIMULTANEOUS CASES

Exp. No.	Time	Pollak	TSI	M	C	N	I				
4	1010-1026	262 \pm 2	275-280	269	103%						
	1141-1156	282 \pm 6	300			230	82%				
5	1428-1440	[~1304]	2800	1405	108%						
	1537-1553	1241	2750			968	78%				
8	1005-1037	273	300-270	308 \pm 22	113%	284 \pm 17	104%	304 ψ	111%	297	109%
9	1207-1221	572	570-550	649	113%			642	112%	~616	108%
	1245-1247		510	544	*						
	1247-1302		510-470			504				530	
15	1510-1529	1683	3200-3300	1810	108%	1509	90%	1564	93%	1563	93%
18	1014-1021	1112	2400	1168	105%			965	87%		
	1044-1053	1190	2500			971	82%			968	81%
19	1145-1206	1057	910-890	1125	106%	1009	95%	1047	99%	1020	96%
20	1400-1425	1068	900-910	1137	106%			1040	97%	990	93%
	1427-1429	"	910	1117	105%	961	90%	966	90%	976	91%
	1513-1516	[1081]	900			900	84%			926	86%
28	1532-1548	392	210	438	112%	400	102%	392	100%	408	104%
Averages				108 \pm 3.6		89 \pm 9.2		98 \pm 9		96 \pm 9	
* = Haze Mode											
ψ = Abbreviated Experiment											

Figures 1-4 were plotted to see if the discrepancies were dependent on particle size. Data from Table 2 was used because of its superior precision. The data from the three DRI chambers (Figs. 2-4) is quite similar and shows a definite size dependence whereas the UMR chamber (Fig. 1) shows only weak size dependence. Loss of particles due to diffusion is a size dependent process which could be responsible for some of the discrepancies between these continuous flow chambers and the TSI. Diffusion loss also depends on the sample flow rate and the distance which the sample has to travel. Since these quantities were nearly identical for the DRI chambers, and the DRI chambers yielded similar data, loss by diffusion seems to be a good possibility. Furthermore, diffusional losses for the UMR chamber are expected to be smaller, because the smaller sample flow branched off of a larger delivery flow. Thus the UMR data reflects this fact (Fig. 1).

Figures 5 and 6 show the theoretical diffusion loss rate for the UMR and DRI chambers, respectively. The agreement between the experimental and theoretical curves indicates that diffusion losses can account for the size dependent relationship between the CFD's and the TSI and thus the differences between the UMR and DRI chambers. However, the data seems to show a linear relationship whereas the theory shows a parabolic relationship. Another possible cause of size dependent particle loss is by electrostatic effects. Since the monodisperse

particles were produced from an Electrostatic Classifier, they are all charged particles. It has been found that just touching the plastic tubing or even the conducting tubing through which the sample aerosol passed caused reductions in the apparent concentration. This is especially true for the smaller particles which have higher mobilities.

These results indicate that, if theoretical diffusional losses were accounted for, the agreement between the UMR and DRI continuous flow chambers would be much better than Table 1 would indicate. With such a correction, it would appear that the UMR and DRI chambers agree to about 5 percent on the plateau concentration for monodisperse aerosols.

3.2 Critical Supersaturation, S_c

The other parameter used for the monodisperse aerosols is S_c , the critical supersaturation, corresponding to the peak in the differential aerosol size distribution. The measured value of S_c was defined as the supersaturation which yielded one half of the concentration found at the high supersaturations where the concentrations were constant (and presumably all particles were being counted). These one-half values for each chamber and for each monodisperse experiment are listed in Table 3. Also listed are the theoretical values, based on the size from the TSI electrostatic classifier used to generate the aerosols. As to the theoretical

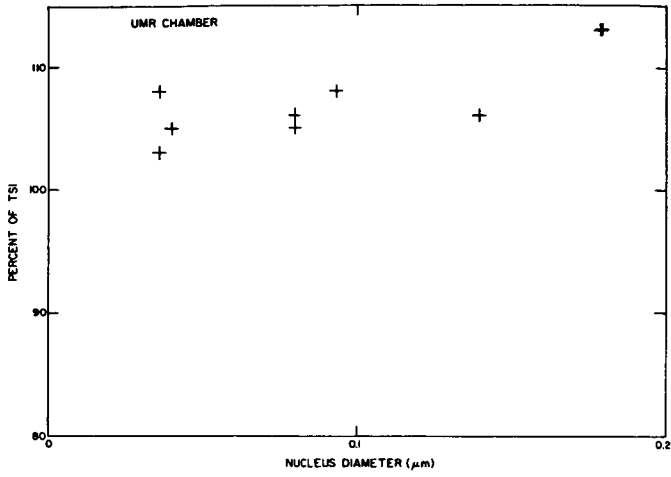


Figure 1. Ratio of concentrations in UMR chamber to those in the TSI CN counter for monodisperse particles of various sizes. Data taken from Table 2.

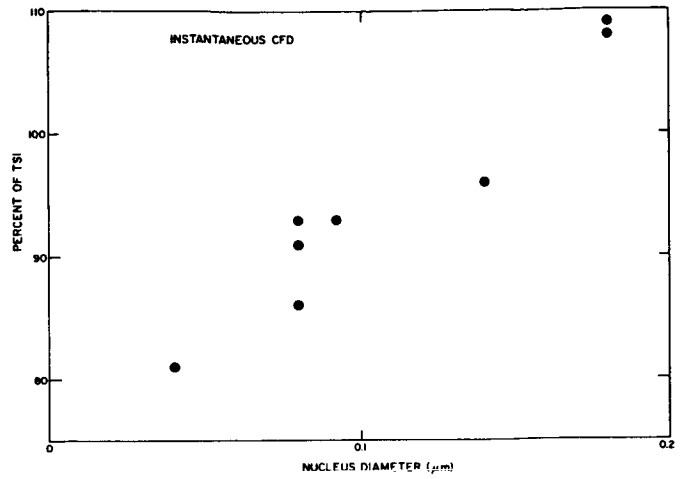


Figure 4. Same as Fig. 1, for DRJ instantaneous chamber.

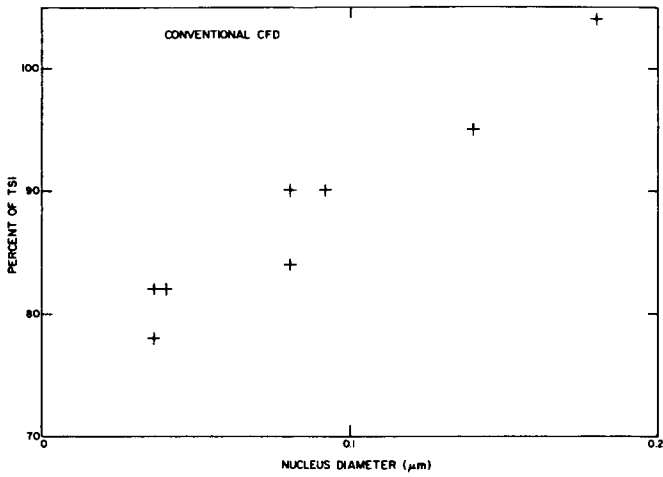


Figure 2. Same as Fig. 1, for DRJ conventional CFD.

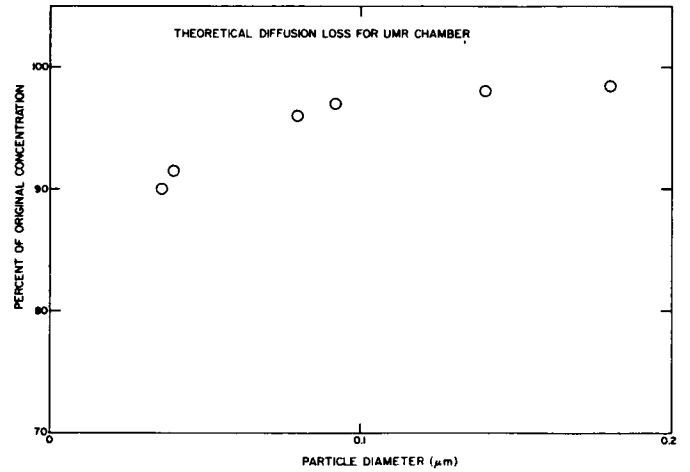


Figure 5. Calculated diffusion losses for the UMR chamber for various particle sizes.

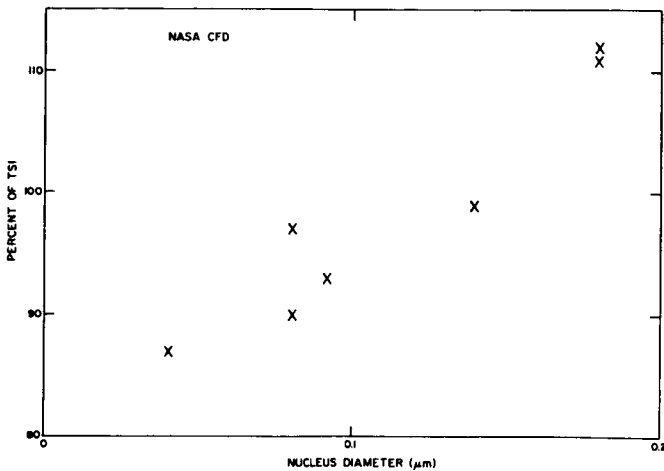


Figure 3. Same as Fig. 1, for DRJ NASA chamber.

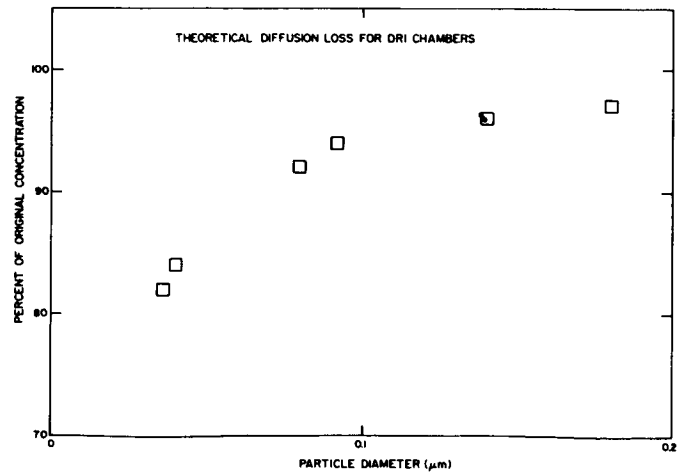


Figure 6. Same as Fig. 5, for DRJ chambers. All three DRJ chambers would have the same diffusion loss characteristics.

TABLE 3. MONODISPERSE S CUTOFF

Exp. No.	Size/Com- position	Theoreti- cal S_c	M	C	N	I	W
4	0.036 NaCl	0.50	0.52	104%	0.52	104%	
5	0.036 NaCl	0.50	0.54	108%	0.52	104%	
8	0.18 NaCl	0.045	0.052	116%	0.084	187%	
9	0.18 NaCl	0.045	0.05	111%	0.065	144%	
15	0.092 Amon	0.18	0.21	86%	0.17	81%	0.2 95%
18	0.04 Amon	0.64	0.62	85%	0.61	84%	0.54 74% 0.70 96%
19	0.14 Amon	0.091	0.11	100%	0.12	109%	0.13 118%
20	0.08 Amon	0.22	0.20	69%	0.18	69%	0.22 85%
28	AgI		0.33		0.40		0.38 0.40 0.30 115%
Averages			97% + 16%	110% + 38%	93% + 18%		
[Excluding Exp. 8,9]			92% ± 15%	92% ± 16%	93% ± 18%		

relation between dry size and S_c , we note that various sources disagree somewhat. For example, for ammonium sulfate at 20°C, Hanel (1976) lists S_c values of 1.91% and 0.0532% at dry diameters of 0.02 μm and 0.2 μm respectively. Jiusto and Lala (their equation 3, this proceedings volume) give approximate relation which, for the corresponding conditions give higher values of S_c (2.01% and 0.0635%, respectively). For sodium chloride, the two sources agree much more closely. In Table 3, we have used the values from Hanel.

It can be seen in Table 3 that the M, C and N chambers showed excellent agreement with theory and each other. The I chamber was really not participating in this aspect of the experiment since it was always operating at three fixed supersaturations throughout the Workshop. Experiments 8 and 9 were actually below the range of the DRI chambers and within the haze mode of the UMR chamber. Exclusion of these two experiments reveals a more realistic test of the diffusion chambers.

The agreement of the UMR and DRI chambers on S_c , indicates that these chambers were sensing the same supersaturations. The agreement with theory shows that the deduced supersaturations were probably accurate.

4. POLYDISPERSE AND AMBIENT EXPERIMENTS

These CCN distributions were more conventional so they could usually be characterized by more typical parameters such as C and K from $N = CS^K$, where C is the concentration at 1% supersaturation and K is the slope of the log-log distribution. However, the calculation of K rests on the assumption of a constant slope and even when this condition is fulfilled, the calculation of K is one step removed from the data. For these reasons, we decided that a better method of analysis of this data would consist of direct comparisons of all of the chambers at the two most extreme supersaturations which can be used. This would test C and K in a more direct manner without the necessity of a straight line distribution. The two supersaturations chosen were 0.70% and 0.30%.

Tables 4 and 5 show the results of this analysis. There is not a standard instrument to relate the measurements to but, since there are so many similarities between the three DRI instruments, we

have again chosen to relate all measurements to the UMR chamber. The fact that the DRI chambers show better agreement with UMR at 0.30% is consistent with the results of the monodisperse experiments. Thus the smallest CCN seemed to be undercounted in the DRI chambers, suggesting that once again diffusion losses or electrostatic charging is a problem. The magnitude of the discrepancy is also quite similar to that found in the monodisperse experiments.

Figure 7 shows that there was probably some undercounting by the DRI instruments at very high concentrations. This data is rather inconclusive except for the concentration of 10,000 cm^{-3} (Exp. 24). Such a graph tests the possibility of effects such as vapor depletion in the cloud chamber or coincidence losses in the OPC. It appears that one of these is a problem in the DRI chambers at very high concentrations.

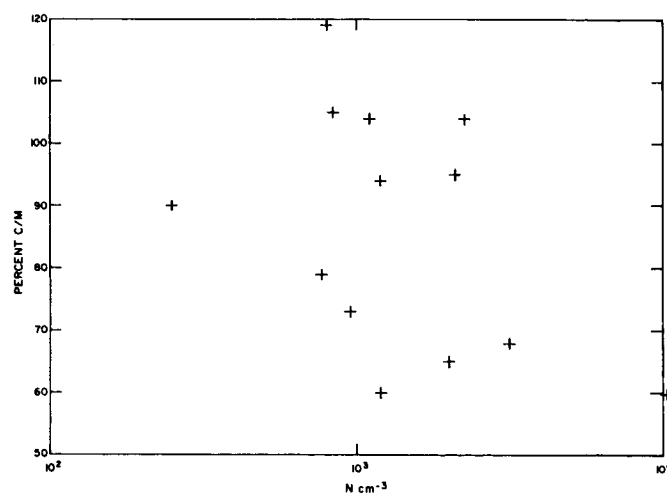


Figure 7. Ratio of concentration in DRI conventional chamber to the UMR chamber vs. concentration in the DRI chamber. Data taken from Table 4.

TABLE 4. POLYDISPERSE AND AMBIENT, 0.7%

Exp. No.	Composition	M	C	N	I	W			
1	NaCl	960	700	73%		410	43%		
2	NaCl	2000	1300	65%		820	41%		
6	Ambient	1200	1125	94%	1300	1175	98%		
10	Bimodal/NaCl	800	950	119%	800	700	88%	1150	144%
11	Ambient	1250			1100	1030	82%	1040	83%
12	Ambient	2100	2000	95%	2000	2000	95%	1000	48%
13	Ammon. Sul.	2250	2350	104%	2150	2000	89%	1200	53%
14	Ammon. Sul.	3150	2150	68%	2300	2100	67%		
16	Ambient	840	880	105%	800	820	98%		
22	Ammon. Sul.	1200	720	60%	720	700	58%		
23	Ammon. Sul.	250	225	90%	235	225	90%		
24	Ammon. Sul.	10800	6500	60%	5000	5200	48%		
26	Ambient	1100	1140	104%	1150	1140	104%		
26	Run 2/Amb.	1250			1280	1150	92%		
27	AgI	775	615	79%	400	650	84%		
27	Run 2	510			320	500	98%		
Averages			86% ± 20%		84% ± 21%	85% ± 16%		69% ± 40%	

TABLE 5. POLYDISPERSE AND AMBIENT, 0.3%

Exp. No.	Composition	M	C	N	I			
1	NaCl	500	435	87%				
2	NaCl	1010	890					
6	Ambient	940	800	85%	700	74%	940	100%
10	Bimodal/NaCl	315	260	83%	280	89%	300	95%
11	Ambient	960	820	85%	700	73%	700	73%
12	Ambient	1000	1200	120%	1200	170%	1200	170%
13	Ammon. Sul.	1300	1010	78%	1010	78%	990	76%
14	Ammon. Sul.	1500	1100	73%	1130	75%	1000	67%
16	Ambient	600	670	112%	540	90%	420	70%
22	Ammon. Sul.	525	450	86%	310	59%	415	79%
23	Ammon. Sul.	110	115	105%	100	91%	120	109%
24	Ammon. Sul.	3050	2800	92%	1300	43%	2500	87%
26	Ambient	600	600	100%	460	77%	660	110%
26	Run 2/Amb.	620			620	100%	620	100%
27	AgI	42	21	50%	65	155%	70	167%
27	Run 2	13			25	192%	42	323%
28	AgI		0.7		101			
Averages			89% ± 18%		94% ± 39%		112% ± 66%	

5. CONCLUSIONS

The five continuous flow diffusion chambers showed good agreement with each other in spite of differences such as three types of optical particle counters and three different wicking materials. With the various aerosol distributions and types of particles, the chambers nearly always agreed to within 15% in number concentrations and supersaturation determinations. Error analysis showed that diffusion losses could account for most of these differences.

6. REFERENCES

Hanel, G., 1976: The properties of atmospheric aerosol particles as functions of the relative humidity at thermodynamic equilibrium with the surrounding moist air. *Advances in Geophysics*, 19, H.E. Landsberg and J. Van Mieghem, Eds., Academic Press, New York, 312 pp.

REVIEW OF ISOTHERMAL HAZE CHAMBER PERFORMANCE

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ABSTRACT

Isothermal haze chambers (IHC) were present at the 1980 International CCN Workshop (Reno, Nevada); the theory of this method of characterizing cloud condensation nuclei (CCN) over the critical supersaturation range of about 0.01% to 0.2% is reviewed, and guidelines for the design and operation of IHC's are given. IHC data from the International Workshop are presented and critically analyzed. Two of the four IHC's agreed to about 40% over the entire range of S_C 's. A third chamber showed similar agreement with the first two over the lower part of the S_C range but only a factor of two agreement at higher S_C 's. Some reasons for the discrepancies are given.

1. INTRODUCTION

Since the Second International Workshop on Condensation and Ice Nuclei in 1970, the measurement of cloud condensation nuclei (CCN) has been extended to the range of 0.015%-0.15% supersaturation through the use of the isothermal haze chamber (IHC), a new type of CCN counter first described by Laktionov (1972). The principle of operation of the IHC derives from the almost unique relationship which exists between the critical supersaturation, S_C , of a particle and its equilibrium size, r_{100} , at 100% RH. In the IHC, nuclei are grown to their equilibrium sizes in an environment of 100% RH and are then counted as a function of size. From these data and the relationship between S_C and r_{100} , one obtains the CCN supersaturation spectrum. Alofs and Podzimek (1974) are credited with bringing Laktionov's work to the attention of the Western cloud physics community.

Four groups operated IHC's at the International CCN Workshop, Reno, NV, 6-17 October 1980: the Desert Research Institute (DRI), the University of Missouri at Rolla (UMR), the Naval Research Laboratory (NRL), and Colorado State University (CSU). All of these instruments are described in companion papers. In addition, descriptions of the DRI and UMR instruments have been published by Hudson (1980) and Alofs (1978), respectively.

The purpose of this review is threefold: (1) to describe the $S_C - r_{100}$ relationship; (2) to discuss some of the critical design aspects of the IHC; and (3) to present some results of the IHC intercomparisons conducted at the Workshop.

2. THE $S_C - r_{100}$ RELATIONSHIP

From the relationship between S_C and dry particle radius (r_0) and that between r_{100} and r_0 one obtains the following relationship between S_C and r_{100} .

$$S_C (\%) = 38.5 \left(\frac{2\sigma_C'}{\rho_W R_V T} \right) \left(\frac{\sigma_C' i_{100}}{\sigma_{100} i_C} \right)^{\frac{1}{2}} r_{100}^{-1} \quad (1)$$

In this equation σ_C' and i_C are the surface tension and van't Hoff factor of the solution droplet when it has attained its critical radius; σ_{100}' and i_{100} are the surface tension and van't Hoff factor of the droplet in equilibrium at 100% RH; ρ_W is the density of water; T is the temperature; and R_V is the gas constant of water vapor.

The relationship between S_C and r_{100} depends on particle composition only to the extent that the value of $\sigma_C' (i_{100}/i_C)^{\frac{1}{2}}$ depends on composition. If we assume that $i_{100} = i_C$ and that $\sigma_{100}' = \sigma_C' = \sigma_W$, σ_W being the surface tension of pure water, then Eq. (1) reduces to

$$S_C (\%) = 38.5 \left(\frac{2\sigma_W}{\rho_W R_V T} \right) r_{100}^{-1}, \quad (2)$$

which is the relationship used by Laktionov (1972). At $T = 20^\circ\text{C}$, and with r_{100} expressed in microns, Eq. (2) may be written

$$S_C (\%) = \frac{0.041}{r_{100}}. \quad (3)$$

Taking into account the temperature dependence of σ_W , we find that S_C varies approximately as $T^{-3/2}$ for a fixed value of r_{100} .

Corrandini and Tonna (1979) analyzed the S_C vs. r_{100} relationship for the main electrolytes composing continental and marine aerosol particles. They found that the uncertainty in the actual electrolytic composition of CCN results in a maximum deviation from Eq. (2) of approximately 6%.

Hoppel and Fitzgerald (1977) examined the effect of insoluble material in aerosol particles on the relationship between S_C and r_{100} . Their calculations showed that insoluble material will not cause a significant departure from Eq. (2) as long as soluble material accounts for at least 5% of the dry particle mass.

The largest departure from the Laktionov relationship [Eq. (2)] is likely to be caused by the presence of surface-active organic materials in the particles which will lower the droplet surface tension below the value for pure water. In the presence of organic material Eq. (2) overestimates S_C by (σ_W/σ_C') , where σ_C' is the actual surface tension of the droplet at its critical radius. The

surface tension of the droplets formed upon atmospheric aerosol particles is not well known. Hanel (1976) made a few measurements of the surface tension of dilute aqueous solutions of atmospheric aerosol samples collected in central Europe and found values 10% to 30% below the value of pure water.

3. CRITICAL DESIGN ASPECTS OF THE IHC

In addition to the errors in IHC data arising from the lack of specific knowledge about the chemical composition of the aerosol particles, the accuracy of CCN spectra obtained with the IHC also depends on the accuracy with which the equilibrium particle size distribution at 100% RH can be determined. In order to achieve an accurate measurement of this size distribution, the humidifying section of the IHC must be truly isothermal, the relative humidity of the air sample must be brought acceptably close to 100%, the aerosol particles must be allowed enough growth time to reach r_{100} and the droplets must be accurately counted and sized. We shall now discuss each of these requirements in turn.

3.1 Isothermal Operation

The diffusivities of heat and water vapor in air are not equal (at standard conditions, the latter exceeds the former by about 15%). Therefore, if an air sample containing CCN flows through a tube having wet walls, the temperature of which increases in the direction of flow, a small supersaturation will be experienced by the CCN. Conversely, if the temperature of the tube decreases along the direction of flow, undersaturation may result. Under either of these conditions, the size of the CCN can depart significantly from r_{100} . It is important, therefore, that the humidifying section of the IHC be accurately isothermal and that its temperature be known to sufficient accuracy.

It is, of course, also important to take into account the inherent dependence of r_{100} upon temperature. From Eq. (2) we find that knowledge of T to an accuracy of 1°C easily suffices for 1% accuracy in S_c .

3.2 Humidification of the Air Sample

It is not possible to humidify the air entering the IHC to exactly 100% RH since this would require an infinitely long chamber. In practice, then, the sample is brought close enough to saturation that the error in S_c resulting from the departure of the actual equilibrium particle size from r_{100} is acceptably small. Figure 1 shows the relative error in S_c as a function of the deviation in relative humidity from saturation. The quantity $1-S$ plotted along the abscissa is one minus the saturation ratio. The error has been computed for different values of dry particle radius (r_0) having the indicated values of S_c . For the purpose of these calculations it was assumed that particles are composed of pure ammonium sulfate. We see from Fig. 1, for instance, that in order to operate an IHC as low as $S_c = 0.015\%$ with an error (in S_c) of no more than 10%, the sample must be humidified to 99.99% RH. If we require an error of less than 5% then we must humidify the sample to 99.996% RH. It is also seen that, should the maximum humidity achieved be only 99.95%, then errors in S_c of as much as 40% can be expected. If the CCN spectrum

can be described by the relationship $N = CS_c^k$, then the error ΔN in particle concentrations due to an error ΔS_c in critical supersaturation is given by $\Delta N/N = -k\Delta S_c/S_c$.

We now turn to the question of how long the humidifying section must be to attain the desired relative humidity. Since most IHC's are of cylindrical geometry, we shall consider the case of air flow through a tube. For flow through a wetted tube, the increase in relative humidity depends only on the length of the tube and the volume flow rate and not on the radius (R) of the tube. The reason for this is that both the time constant for diffusion of water vapor from the walls and the residence time of the sample in the tube are proportional to R^2 . Figure 2 shows the relative humidity at the center of a wetted tube as a function of distance from the inlet and volume flow rate, ϕ , for the case of an air sample having an initial relative humidity of 50%. Figure 2 is based on analytical solutions to the diffusion equations for laminar flow through tubes (Goldstein, 1965). These solutions assume Poiseuille flow and neglect the axial diffusion term. For initial relative humidities of 30% and 70%, the tube length required to raise the relative humidity of the sample above 99.9% is about 5% greater and 10% less, respectively, than the length needed for air entering at 50% RH.

The volume flow rate through the humidifying section is usually determined by the flow requirements of the optical particle counter (OPC) used to count and size the droplets. The Royco 225 optical counter, for instance, uses a flow of $47 \text{ cm}^3\text{s}^{-1}$ (or 2.8 liters/min). For this flow rate, a tube length of 140 cm is required to humidify the sample from 50% RH to 99.99% RH.

3.3 Residence Time Needed to Achieve Droplet Equilibrium

When an aerosol particle is exposed to an environment of increasing relative humidity its actual size will lag behind its equilibrium size. The smaller the critical supersaturation of a particle, the larger is its equilibrium size, r_{100} , and the longer it takes to grow to that size. Therefore, the humidifying section of the IHC must provide enough residence time at ~100% RH so that the CCN can grow acceptably close to r_{100} .

Laktionov (1972) presented, without documentation, calculated values of the time required for droplets to grow to within 5% of r_{100} in an environment of 100% RH. These values have been used in determining the length of, or time required in, the humidifying section of the IHC. Recently, Robinson and Scott (1980) computed growth times of water solution droplets in an IHC, using a new droplet growth equation derived from first-principles kinetic theory. This equation was shown to agree with a conventional growth equation to within 5% for droplets smaller than $1 \mu\text{m}$ and to within 0.1% for large radii. Table 1 is taken directly from their paper and shows the time needed for nuclei of varying S_c to grow from their dry size to 95% and 99% of r_{100} in a saturated environment. The computations of Robinson and Scott assumed that nuclei are composed entirely of sodium chloride and that the condensation coefficient of water is 0.036. Table 1 also lists Laktionov's values of the time needed

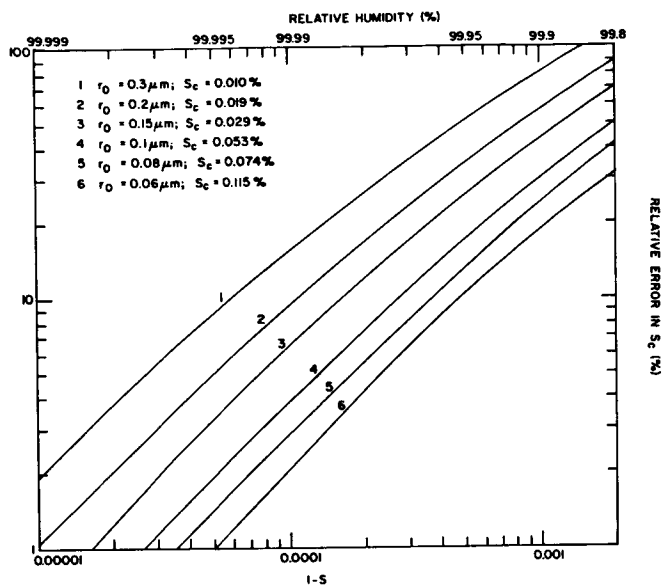


Figure 1. Relative error in S_c , in percent, as a function of the deviation in relative humidity in the IHC from saturation. The error is shown for six values of dry particle radius, r_0 , having the indicated values of S_c .

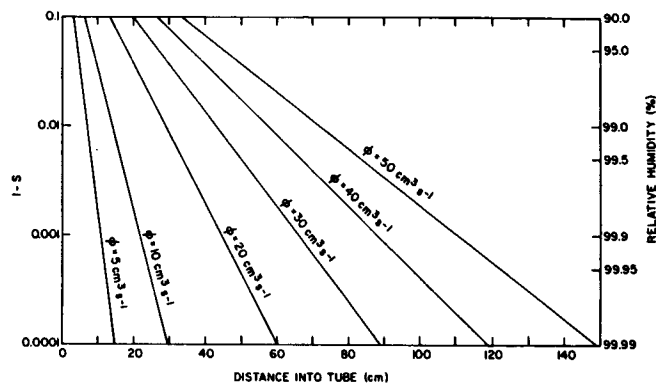


Figure 2. Relative humidity of the air at the center of a wet-walled tube as a function of distance down the tube and volume flow rate, ϕ . The initial relative humidity of the air is 50%.

TABLE 1. TIME REQUIRED FOR DROPLET GROWTH FROM DRY SIZE TO 95 AND 99 PERCENT OF r_{100} AT $S = 1.0$, $T = 20^\circ\text{C}$

S_c (%)	0.103	0.063	0.051	0.041	0.033	0.026	0.021	0.016
r_{100} (μm)	0.40	0.65	0.80	1.0	1.25	1.6	2.0	2.5
t(S) to .95 r_{100}	1.6	4.6	7.4	12.5	21.3	38.9	68.1	121
t(S) to .99 r_{100}	3.3	9.8	16.7	26.6	45.5	84.1	148	262
t(S) [Laktionov] to .95 r_{100}	4	10	15	25	40	65	100	180

to grow to 95% of $r_{p,0}$. It is seen that Laktionov's growth times are significantly greater than those computed by Robinson and Scott.

Since Laktionov did not give the details of his calculations, we do not know the reason for the discrepancy in growth times. If we accept the values of Robinson and Scott, then we see that an IHC must provide over 120 s of growth time at a relative humidity of ~100% if it is to give accurate results at supersaturations as low as 0.015%.

A conservative estimate of the total length needed for the humidifying section may be obtained by adding together the length needed to humidify the air to an acceptable closeness to 100% RH (see previous section) and the additional length necessary to provide 120 s of residence time at ~100% RH.

3.4 Droplet Sizing and Counting

The accuracy of CCN spectra obtained with the IHC also depends on the accuracy with which the droplets emerging from the isothermal (humidifying) chamber are counted and sized. In order to obtain an accurate measurement of the droplet size distribution, the droplets must enter the OPC without serious modification of their numbers or sizes, and the response curve of the OPC must be well known, both in terms of absolute accuracy and resolution.

(a) Entry of Droplets into the OPC

The IHC droplet size distribution can be modified as a result of impaction of droplets on the walls of the inlet tube of the OPC and by evaporation shrinkage of the droplets before they enter the light beam. Alofs (1978) made a thorough investigation of both of these problems as part of an analysis of the performance of his dual-range cloud nucleus counter. In Alofs' device, these two problems may be amplified, compared to other IHC's, because the droplets must pass through a tube 1.3 mm diameter by 12 cm long to reach the OPC.

Alofs (loc. cit.) checked the problem of impaction by varying the flow into his OPC (a Royco at that time) while keeping the main flow through the humidifying chamber constant. The tests were made for 2.5 μm and 5.0 μm radius droplets. For each of these sizes, it was found that a plateau of count vs. Royco flow existed (where presumably impaction is not a problem) and that at higher flow rates the count dropped off due to impaction. Losses due to impaction set in at a flow rate of ~1.0 liters/min in the case of the 2.5 μm droplets and at a flow of ~0.5 liters/min for the 5.0 μm droplets. Alofs attributed the loss in counts to the fact that the aerosol stream was not exactly lined up with the 1.3 mm inlet tube. It seems reasonable to expect that in an IHC in which the droplets do not pass through a narrow tube to reach the OPC, higher flow rates can be used before impaction becomes a problem.

The inlet tube of the OPC will normally be warmer than the isothermal chamber temperature due to electrically-generated internal heat in the OPC. This will result in some evaporation of the droplets before they are sized. Alofs (loc. cit.) calculated the change in droplet size due to evaporation in the OPC inlet tube of his device. One may conclude from these calculations that, for nuclei

with $0.01\% \leq S_c \leq 0.1\%$, evaporation will be relatively insignificant in an IHC which uses the full manufacturer's flow rate through the OPC (2.8 liters/min in the case of the Royco 225), even if the inlet tube is as much as 0.5°C warmer than the isothermal chamber. However, if the droplet residence time in the OPC inlet is significantly increased due to the use of a reduced flow rate, then it is necessary to avoid evaporation. Alofs showed that the problem of evaporation (of nuclei with $S_c \leq 0.1\%$) could be effectively eliminated by thermostatting the OPC so that the inlet tube is not more than 0.1°C warmer than the humidifying section.

The problems of impaction and evaporation are both very difficult to avoid completely by a priori design provisions. It seems best to allow enough adjustability in the IHC design, that these phenomena can be avoided by experimental adjustments.

(b) Sizing Accuracy of the OPC

In contrast with OPC use with continuous-flow, parallel-plate counters, the OPC on an IHC must not only count, but must also size accurately in order to give the concentration of nuclei as a function of S_c . This means that the response curve (i.e., signal strength as a function of particle radius) of the particular OPC used must be accurately known for the case of water spheres. OPC's are calibrated by the manufacturer with polystyrene latex spheres which have an index of refraction, m , of 1.59. Therefore, use of the OPC with the IHC to size water droplets ($m = 1.33$) means that the manufacturer's calibration curve must be corrected for index of refraction.

Cooke and Kerker (1975) used a Mie scattering computer program to predict the theoretical response curves of several widely used commercial OPC's, including Royco and Climet devices, for a range of values of refractive index. These curves, for all but the Climet and Royco 220 devices, show an initial monotonic increase in signal with increasing particle size, followed by a region of multivalued response. For particle sizes larger than about 1 μm radius, the response of OPC's again becomes single valued. The Royco 225, a popular device used by Hudson (1980) and others, is not treated by Cooke and Kerker, but its response should be close to that of the Royco 245, which is treated.

The calculated response curves of Cooke and Kerker are presently the best available basis for extrapolating the manufacturer's polystyrene latex calibration to work with water droplets. In using these curves to correct for index of refraction, a problem arises in that the response curves are calculated for refractive indices 1.33, 1.45, 1.54, and 1.70, but not for index 1.59, that of polystyrene. The operators of IHC's, then, have been forced to interpolate between the curves presented by Cooke and Kerker, a procedure which is made difficult by the oscillatory, mathematically complicated nature of these curves. As pointed out by Hodgkinson and Greenfield (1965), the initial monotonically increasing portions of the response curves represent the large-radius end of the Rayleigh scattering region, the upper limit of which occurs at a particle radius of about 0.05 μm . As long as the monotonic character of the response curves persists above the Rayleigh scattering re-

gion, the response can be expected to vary as $(m^2-1)^{1/2} (m^2+2)^{-1/2}$, and the interpolation between curves to provide response values for other indices of refraction may be carried out on that basis. However, the oscillatory (multivalued) region of many OPC response curves sets in at values of radius well below 0.5 μm , marking the end of any resemblance to the behavior in the Rayleigh region. Outside the Rayleigh region, the complicated nature of the scattering functions (wherein m appears both as the coefficient and argument of Bessel and Legendre functions) precludes easy interpolation. One then has the choice of either repeating the work of Cooke and Kerker for $m = 1.59$ or simply doing a visual interpolation based on the Cooke and Kerker curves. There may, of course, be considerable error involved in determining the response curve by the latter method.

OPC size resolution on the order of 0.1 μm is required to be able to distinguish between nuclei differing by as much as 0.02% in S_c . Whitby and Willeke (1979) have estimated that most OPC's in good adjustment will artificially broaden a monodisperse spectrum to the extent that the resulting geometric standard deviation is of order 1.1. This would imply, for example, that slightly over 68% of the count of an ideal 1.0 micron aerosol would be interpreted by the OPC as belonging to aerosol sizes between 0.91 and 1.1 μm .

The actual calibration of any OPC should be checked by the user but, in general, sizing errors are a serious concern, especially if the OPC has a multivalued response function in the radius range of interest.

4. HAZE CHAMBER DATA FROM THE INTERNATIONAL WORKSHOP

4.1 Preliminary Comments

While IHC's were operational during all 29 experiments of the International Workshop, useable data resulted from only a subset of the total number of experiments when the aerosol was large enough to register and when other problems such as fluctuations were not present. In addition, the four instruments (#11, Naval Research Laboratory; #14, Desert Research Institute; #21, University of Missouri; #26, Colorado State University) were not always simultaneously operational. Therefore, the data to be discussed is a subset of the total data file, which will generally cover those experiments where the aerosol was suitable for IHC and all four IHC's were operational.

A few remarks about the instruments themselves are worthwhile. IHC #21 is unlike the others in that it uses a Climet OPC for detection and sizing of the haze droplets. IHC #11 is actually a second-generation device, built by the Naval Research Laboratory after considerable experience had been gained with an earlier model. In particular, a very efficient sheath air pre-humidifier is found on this version. IHC #14 and #26 are related in terms of design; #26 was constructed following, to some extent, the mechanical details of #14 but was only newly-completed at the time of this Workshop. Some difficulties in the operation of #26 were experienced, but allowance should be made for the fact that it was a newly-constructed device. Finally, one should generally note that IHC's are required to respond to concentrations that span many (eg,

3-5) orders of magnitude, a considerable dynamic range which is taxing under the best of circumstances.

In the process of analyzing the data, it was discovered that somewhat different procedures were used to deal with the index of refraction problem cited in Section 3.4(b) of this report. Although DRI and NRL both used Royco OPC's it was determined that subtle differences in the procedure used to correct for the index of refraction resulted in some significant differences in the data. Specifically, this meant that different S_c 's were deduced from voltage threshold responses which would have shown identical sizes for the polystyrene latex spheres used for size calibrations. After this was discovered, the DRI data were redone using the method similar to that used by NRL. Since the CSU data were treated identically to NRL data, this modification then allowed all Royco-equipped IHC's to be compared on an equivalent basis. It is difficult to determine which procedure is best, although the NRL procedure does yield higher concentrations, at a given S_c , in better agreement with the UMR chamber and with the mobility analyzer.

The operators of the UMR chamber accounted for the index of refraction in an entirely different manner. While the procedures used for the Royco-equipped chambers were based solely on theory, the UMR IHC relied most heavily on an empirical technique. Size thresholds were determined by passing known sized NaCl nuclei through the UMR IHC; the nuclei would form drops which were assumed to be the equilibrium size of solution droplets at 100% R.H. This procedure was followed for one or two sizes and the rest of the curve was completed by using the functional shape of the Cooke and Kerker results. This circumvents the index of refraction problems which have been described in Section 3.4(b) and in the last paragraph for the Roycos.

4.2 Best and Typical Agreement Cases

In Figure 3, the IHC results of Experiment 20 (monodisperse ammonium sulfate aerosol) are shown; the data provide the best agreement between IHC's found during the Workshop. The curve labeled #14' is the original DRI data; #14 has been redone with the index of refraction correction treated similarly to the data of #11 and #26. This yielded much better agreement with #11 and #21. Note that the theoretical CCN spectra derived from the NRL mobility analyzer data (#12) are also plotted; these were obtained by using the theoretical (Kohler) relationship between S_c and dry particle size for ammonium sulfate aerosols to transform the NRL size distribution data into a CCN spectrum. (IHC #26 did not report data for this experiment.)

Figure 4 shows a count-versus- S_c plot of Experiment 9, monodisperse sodium chloride, and represents the more "typical" case from the Workshop. Once again, it is apparent that #14' shows much closer agreement with #11, #12, and #21. The difference between #14 and #14' shows the sensitivity to the method used to correct for the index of refraction. In contrast to Figure 3, it was more frequently the case that the count magnitudes were in the order shown in Figure 4 at the higher S_c 's: IHC #11 and #21 highest, #14' next, then #14, and #26 lowest. At the lowest supersaturations, however, curves 14', 11, and 21 were in no consistent

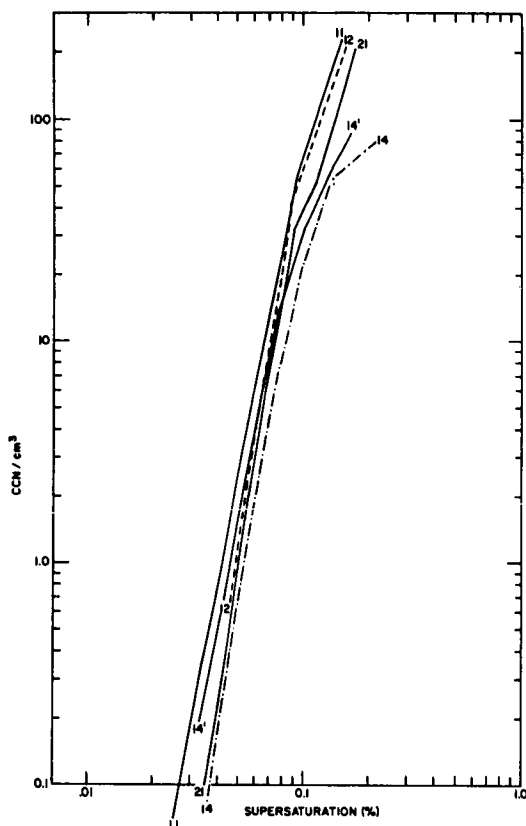


Figure 3. Concentrations registered by isothermal haze chambers and NRL mobility analyzer as a function of supersaturation setting, for Experiment 20. Aerosol was monodisperse ammonium sulfate. Mobility analyzer data are labelled by instrument Number 12. Haze chamber data are similarly labelled; see text for instrument numbers.

order and agreement between the three IHC's was much better.

Not shown in Figure 4 are the counts of the continuous-flow CCN counters. In Figure 3 this data shows a "plateau" in count; the mean S_C of this nominally monodisperse aerosol was 0.045%, with the smallest CCN in the distribution exhibiting S_C as high as 0.06% to 0.1% supersaturation. For this experiment the counts from those devices generally fell in the range 500-600, or at about the same magnitude as indicated at 0.15% by IHC #11, #21, and #14. IHC's #21 and #14 (#14' also) detected a plateau in number vs. S_C which allowed them to estimate the S_C of the aerosol. These were all overestimates of the theoretical value 0.045%; #21 saw 0.055%, #14' estimated 0.073%, #14 0.086%. Although IHC #11 did not detect a plateau this may have just been due to the OPC channel threshold selections. If 0.15% were assumed to be a plateau reading, #11 would estimate 0.073%. Experiment #8 was the only other experiment which used an S_C in the IHC range, 0.045% as Experiment 9. Although the concentration was lower in Experiment 8, the results were nearly identical to Experiment 9. IHC #26 often exhibited a kink in the region around and immediately below 0.1%; this kink may correspond to the double-valued region of the particular Royco in use on IHC #26.

The relative order (by magnitude of count at a given S_C) of the four instruments was preserved

throughout most of the experiments. Considering that three of the devices utilized a Royco OPC, the double-valued response region of which falls in the approximate S_C interval of 0.04% to 0.08% ($r_{1,00}$ of 0.5×10^{-4} cm to 1.0×10^{-4} cm), it may be inferred that at least a given Royco OPC seems to consistently place signals of the same magnitude in the same electronic channel, even though the signals may fall within the double-valued region. Otherwise, interpretation of data from the IHC's using the Royco (IHC #11, #14, #26) would be next to impossible in the 0.04% to 0.08% S_C range. (Even granting this consistency, it is not too surprising to see discrepancies in this troublesome region of the spectrum.)

4.3 Relative IHC Results as a Function of Experiment

Figures 5 and 6 simply display the relative counts of each IHC as a function of experiment number, where the relative count is shown as a percentage of the NRL mobility analyzer count at the same S_C ($S_C = 0.09\%$ for Fig. 5 and 0.15% for Fig. 6). The curves on Figs. 5 and 6 show discontinuities where data are inappropriate or missing. These S_C values were chosen somewhat arbitrarily; our intention is simply to display relative performance at representative points along the S_C spectrum.

In Figure 5 once again #14' is plotted. As in the last two figures, this represents DRI data which has been corrected for the index of refraction in a manner similar to that used by the other

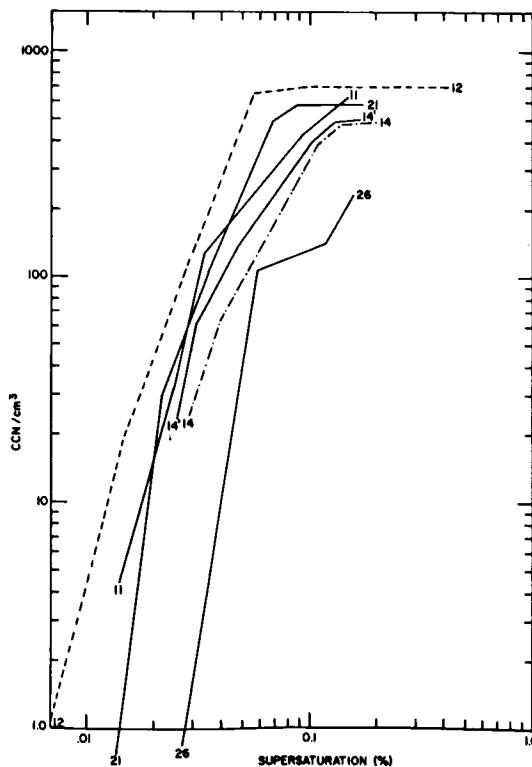


Figure 4. Same as Figure 3, but for Experiment 9. Aerosol was monodisperse sodium chloride.

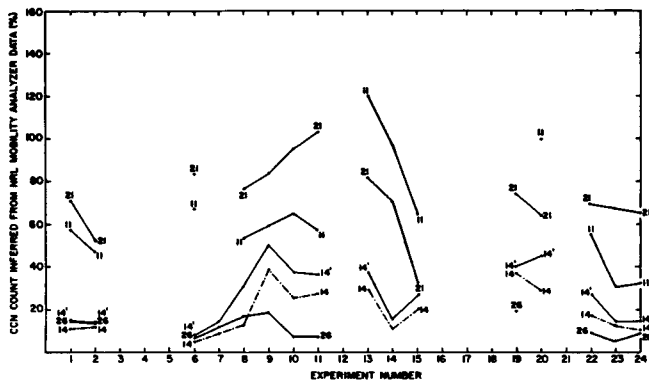


Figure 5. Isothermal haze chamber data compared to NRL mobility analyzer data, plotted as a function of experiment number, for a supersaturation setting of 0.09%.

two Royco equipped IHC's. No. 14' is not plotted in Figure 6 because at 0.15% the ratio of #14' to #14 is only about 1.15 except in Experiments #22, 23 and 24 where it is about 1.5, causing only a small change in the relative positions of the data from the various instruments.

In examining these figures, it is apparent that the relative order of the instruments is generally preserved although #11 and #21 seem to alternate as the highest reading IHC. A similar figure for $S_c = 0.04\%$ was constructed, but had very large gaps in data, and considerably more scatter than seen in Figs. 5 and 6. Nevertheless, the order was like that of Figure 5, except that #14' alternated with #11 and #21 for highest readings. These data cover ambient aerosols (Experiments 6 and 11), monodisperse sodium chloride (8 and 9), polydisperse sodium chloride (1,2,10), monodisperse ammonium sulfate (15, 19,20) and polydisperse ammonium sulfate (13, 14,22,23,24). The maxima and minima of the curves of the four instruments sometimes seem to occur in unison, but are not clearly related to any particular type of aerosol. Rather, it appears that a significant variable might be the response of the NRL mobility analyzer, more than the type of aerosol under study. From experiment to experiment, small changes were occasionally made in the sample flow rate (time of exposure of the sample to saturation) of instrument #14. Although these should have been within the region where counts have plateau values as a function of carrier flow, this may not have always been the case for all of the OPC droplet size thresholds used.

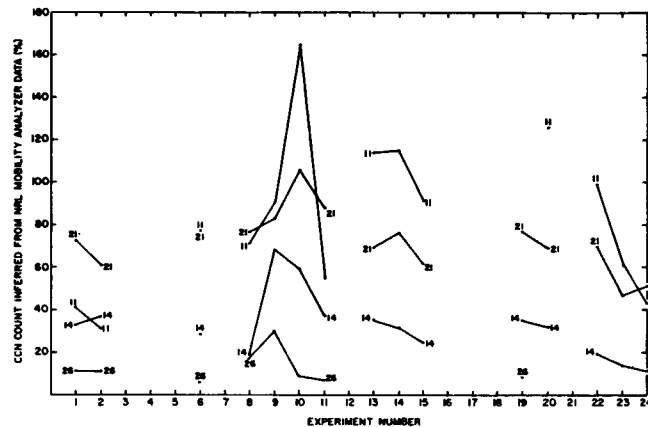


Figure 6. Same as Figure 5, but for a supersaturation setting of 0.15%.

Figure 6 takes on additional significance if we regard 0.15% as representative of the supersaturation range where haze chamber data overlaps that of the static and continuous-flow CCN counters. Inspection of the Workshop data shows that the NRL mobility analyzer usually read somewhat higher than the approximate average of the static and continuous counters, by a factor between one and two. Therefore, if those CCN counters were also shown on Figure 6, their results would fall in about the same range as given by the curve for IHC #21, a result that is not too surprising because this IHC always gave data which was continuous with the higher supersaturation data given by the dual-mode operation of chamber #21. Thus, both IHC #21 and IHC #11 tended to agree well with other chambers around 0.15% supersaturation. The other two IHC's were less often in agreement in this region, although often #14 seemed to give data which joined well with the data of a continuous CCN counter from the same institution, #15.

Although Figures 5 and 6 indicate that there is a considerable counting discrepancy between the various IHC's, it should be kept in mind that when the slope of the CCN spectrum is very steep (as was often the case in the range of S_c of the IHC's), a small absolute error in S_c can result in a large error in particle count. Thus small discrepancies in the sizes assigned to given water droplets by various OPC's could lead to large differences in count; ignoring for the moment all other sources of the discrepancies, the primary calibrations (even of OPC's from one manufacturer) of OPC's differ

from instrument to instrument. In regard to the present data, clearly the most desirable approach would have been to compare the OPC's (at least the three Roycos) on a non-water aerosol such as polystyrene latex before or during the Workshop.

4.4 Conclusions

The general tradition in comparisons of CCN counters has been to regard the instruments which register the highest counts as the most correct. That tradition should never be accepted without scrutiny, for instruments may overcount as well as undercount, although generally there are more reasons for the latter. In the case of these comparisons, there are additional measurements which drive one to the conclusion that the higher-counting instruments are indeed to be favored; those measurements are the continuous-flow and static CCN counts, in the case of monodisperse aerosols where a plateau in the count-versus- S_c is to be found, and the results of the NRL mobility analyzer. It would appear, then, that instruments #11 and #21 offer some advantages over the other devices.

The differences between #14 and #14' point out the problems involved in correcting for the index of refraction. Although these differences at first appeared to be very minor, they resulted in significant differences in the region of supersaturation between 0.04% and 0.08%, corresponding to the double valued region of the Royco. The steepness of the aerosol distribution in this region results in a very high sensitivity to this variability. Fortunately all of the IHC's usually avoided using voltage thresholds in this region of the spectrum. Hence the differences involved in correcting for index of refraction were usually not as serious as they could have been. Although the differences which resulted from variations in the procedure are significant they are not overwhelming. Nevertheless, it would have been much better to use a uniform method of correcting for index of refraction at the time of the Workshop as corrections made later are not certain to be identical. In the future, it would be desirable to look into the whole question more thoroughly.

IHC #21 is different in that it uses a Climet OPC. However, it is difficult to compare the treatment applied by its operator to the index of refraction problem, which unlike the Royco-equipped chambers, does not strictly rely on theoretical considerations. Hence it is difficult to determine if the Climet OPC offers advantages in sizing water droplets based upon calibration with polystyrene latex spheres of a different index of refraction. Nonetheless, the UMR IHC does show good number concentration agreement with the NRL mobility analyzer as well as better agreement with the theoretical S_c 's in the monodisperse experiments (8 and 9). This is understandable since the UMR chamber was essentially calibrated with a mobility classifier, which may be a better method for applied CCN counting. Nevertheless even though the Royco equipped IHC's have considerable calibration problems they do at least use an independent method of calibration. There are more occasions when IHC #11 counts higher than the NRL mobility analyzer, than is the case with IHC #21, but such occasions are rare enough that no conclusion can be drawn.

All IHC's in this study were basically stable, rugged devices which at least showed an output data

signal which was proportional to the input aerosol concentration, despite the taxing dynamic range and requirement that OPC accuracy and resolution be pushed to the limit. These devices clearly have a promising future in certain measurements of hygroscopic aerosols, such as the counting of fog condensation nuclei, or even in new applications such as studies of aerosol retention in the human lungs.

5. REFERENCES

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6. ACKNOWLEDGEMENTS

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AEROSOL GENERATION AND DISTRIBUTION SYSTEM
FOR THE THIRD INTERNATIONAL CLOUD CONDENSATION NUCLEI WORKSHOP

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ABSTRACT

In order to obtain identical samples participating CCN instruments and aerosol characterizing equipment were located along and connected to a 8.2 cm diameter aluminum tube through which the test aerosols were pumped directly from the source at a rate of 200 to 600 μmin^{-1} and at very slight overpressure.

Of the total of 29 experiments, 18 were carried out with artificial NaCl or $(\text{NH}_4)_2\text{SO}_4$ aerosols. These were generated from salt solutions by pneumatic atomizers of special design (by DRI) to ensure high constancy of the aerosol output concentration. In three experiments with insoluble CCN (AgI, paraffin wax) the aerosols were generated thermally. In some of the tests, an electrostatic classifier was used for narrowing the particle size distributions.

1. INTRODUCTION

Comparisons or cross-calibrations of small particle instrumentation are often performed, for convenience's sake, with the aid of most easily obtainable aerosols, such as the ones present in room or atmospheric air, the particles of which are usually not well characterized and are mostly of a rather complex nature. However, since different aerosol instruments generally differ in their response spectra, these differences can only be assessed and ultimately understood if aerosols of distinct characteristics are utilized. Thus, in order to achieve the objectives of the Workshop, it was imperative to provide test aerosols of high monodispersity and chemical purity, and to take the utmost care in conveying the sample to the instruments.

2. AEROSOL DISTRIBUTION SYSTEM

Based on the favorable experience in previous Workshops (WS), the primary requirement of supplying all instruments simultaneously with as nearly identical samples as possible was met by locating the instruments along a sample duct (or manifold) in which a flow rate was maintained that was high compared to the sample flow extracted from the duct by the instruments.

A further requirement was to provide a steady flow of aerosol in which variations of particle concentration with time were less than a few percent ($\leq \pm 3\%$, according to recommendations from the Steering Committee). In order to achieve this criterion, the choice was to either use a large storage bag (as had been the case at the Ft. Collins WS) or to operate a very constant aerosol source.

Several considerations led to the decision to deliver aerosols directly into the sampling duct instead of using a large bag. With regard to artificial aerosols, generation techniques at the DRI had improved over the years to the point where runs of one to two hours could be achieved with about 90% probability of keeping the particle concentration within $\pm 5\%$ of a mean value whereby the short term fluctuations (which are generally more detrimental for instrument comparisons) usually were even smaller; superimposed on this was a slow drift in the particle concentration. Furthermore, examination of the Ft. Collins WS data (Grant, 1971) indicated that samples stored in the 54 m^3 bag were decaying at a considerable rate; total NaCl particle concentration, $\sim 10\% \text{ hr}^{-1}$; large sub-micron particles, $\sim 50\% \text{ hr}^{-1}$; and ~ 20 to $\sim 30\% \text{ hr}^{-1}$ for the CCN of atmospheric samples active at 1% supersaturation. These values did not point to an advantage of a bag system, not even for the case of ambient aerosol when compared with the low level of fluctuations in the prevailing situations of westerly winds usually experienced at the DRI during routine direct measurements of the atmospheric aerosol.* Another disadvantage of a bag system is its finite volume dictating the duration of an experiment (which may be too short for certain investigations; some participants had indicated the desirability of prolonging some experiments well beyond an hour).

Since the DRI aerosol laboratory was already equipped with a duct system to aspirate outside air through an inlet 9 m above the roof (i.e., ~ 25 m above ground level) for distribution to instruments, it was relatively easy to extend that system to the WS room. Figure 1 shows the floor plan of the 150 m^2 WS area and the layout of the sample duct. The numbers 1 to 28 refer to the sampling outlets spaced about 0.6 m apart along the 8.2 cm ID aluminum duct. The sampling outlets were of 5 mm stainless steel tubing having a 90° bend inside the duct, pointing into the air flow; this resulted in isokinetic sampling conditions for instrument sampling rates of 0.0037 times the flow rate in the duct (which was varied between 200 and $700 \mu\text{min}^{-1}$ from one experiment to another). Also shown are the sampling locations of all the instruments (identified by type and participating organization). In order to minimize the risk of losing particles

* Unfortunately, unusually calm weather persisting during the period of the WS, combined with local industrial sources, was causing considerable fluctuations in aerosol concentration.

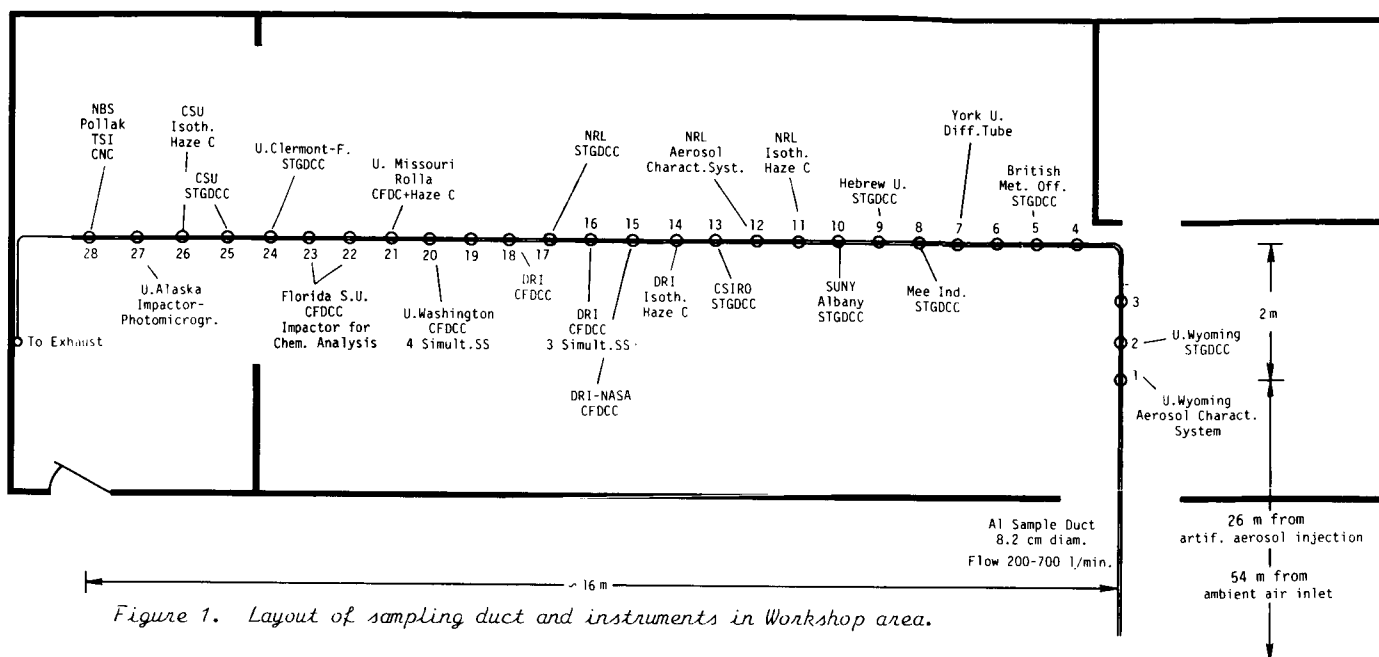


Figure 1. Layout of sampling duct and instruments in Workshop area.

through charge accumulation on surfaces, all instruments were connected to the sample duct by electrically conductive latex rubber tubing which had been tested previously and found to have no adverse effects on aerosol sampling.

Although the distance between the first and last sample outlet was 18 meters, the sample time delay between first and last instrument was only 8 to 25 seconds depending on the sample flow velocity of 0.6 to 2 m s⁻¹. Since significant changes in aerosol characteristics (mainly concentration) generally were associated with considerably longer time constants, simultaneous sampling by all instruments can be assumed for all but a few circumstances. By measuring the total particle concentration at sample outlets 3 and 28 with the same instrument (alternating between the two outlets) it was determined that no significant decay of the aerosol along the duct was taking place.

In order to prevent contaminants from entering the sampling duct, the latter was kept at slightly higher than room pressure, i.e., 0.05 to 0.5 mb, depending on sample flow rate. When using artificial aerosols, the compressed air generation system provided the needed overpressure while, in the case of ambient air sampling, an axial fan (Rotron Model TN3A2) built into the sampling duct upstream of the WS area produced the pressure difference. It was determined experimentally that the fan caused a reduction in total particle concentration in the ambient aerosol by about 5% but, more importantly, that its motor did not generate any particles.

3. AEROSOL GENERATION SYSTEM

For reasons mentioned previously, more emphasis was placed on work with artificial, well defined CCN which were used in 21 of the experiments while the ambient aerosol served in only eight comparison tests.

In order to cover the basic types of soluble, insoluble and hydrophobic aerosols, NaCl and (NH₄)₂SO₄

were selected for the first category due to their relevance to the real atmospheric aerosol. The choice for the water insoluble CCN material fell on pure AgI, mainly due to the relative ease with which this highly insoluble substance can be dispersed; it has to be pointed out that these AgI particles most likely do not resemble the ones generated for cloud seeding purposes which usually possess a water soluble component. Originally, the candidate material for hydrophobic CCN was Teflon (briefly used at the Ft. Collins WS), but the lack of a satisfactory dispersal method led to the use of paraffin wax instead.

Parameters to be varied other than the chemical make-up were the total particle number concentration (a few hundred to several thousand per cm³), and the particle size distribution ("monodisperse", polydisperse and mode at several different sizes).

The experimental set-up for producing these aerosols is shown schematically in Figure 2. The aerosol generator, in this case a pneumatic atomizer for the production of water soluble CCN from salt solutions, is depicted at the top right of the figure. While many types of atomizers, especially inhalation therapy nebulizers, could, in principle, have been used for the present purpose, the requirements of constant output and prolonged operation demanded the design of a specialized and more versatile device as described in detail by Dea and Katz (1981) in a subsequent article.

In order to minimize contamination while providing a wide range of pressures needed for control of output parameters, the atomizer was operated with high grade compressed air from commercial tanks. The salt solutions were prepared from J.T. Baker and Fisher Scientific reagent grade salts and HPLC water (J.T. Baker Chemical Co.). The concentration of the salt solutions was varied according to the particle size requirements of individual experiments. After emerging from the atomizer, the fine mist of solution droplets was diluted about

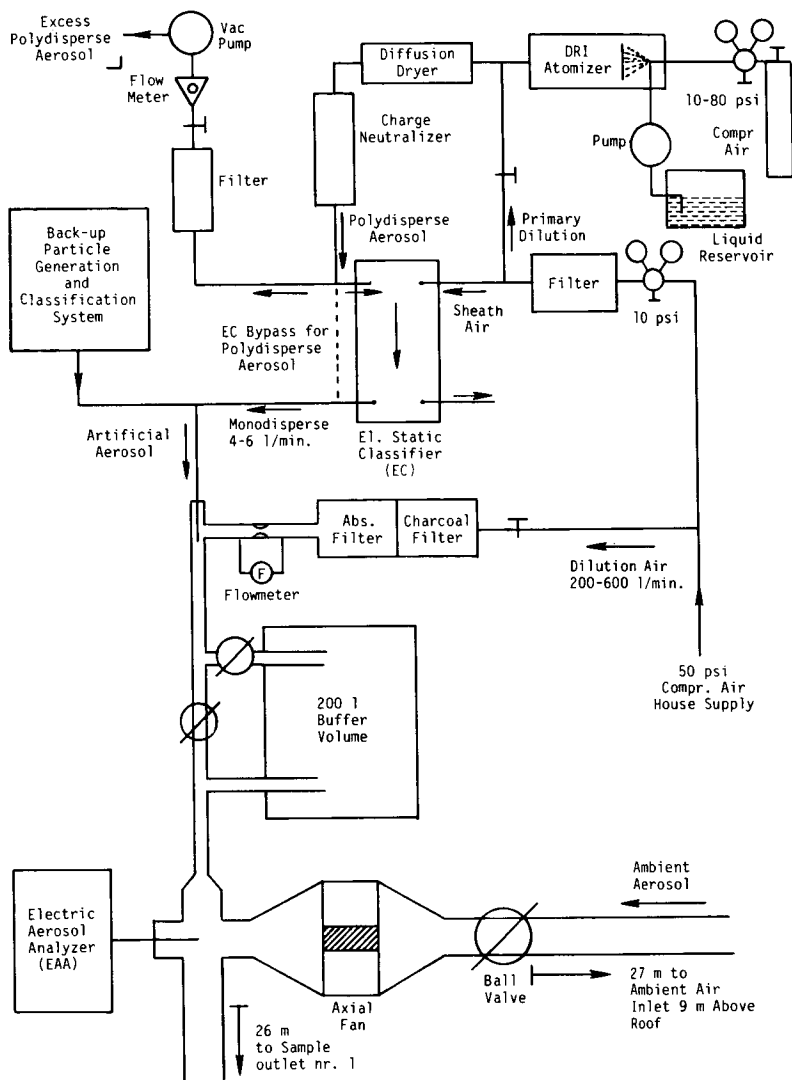


Figure 2. Schematic of Aerosol Generation System.

1:1 with dry filtered air, mainly to promote rapid evaporation of the droplets, but also to help reduce coagulation which is considerable at typical number densities of the order of 10^8cm^{-3} .

After complete removal of liquid water from the particles in the diffusion dryer (TSI Model 3062), the aerosol was brought to charge equilibrium by passing it through a charge neutralizer equipped with a 10 mCi Kr^{85} source. While this procedure helped reduce particle losses caused by charge effects, it was mainly preparatory to extracting a quasi-monodisperse fraction from the total atomizer output by means of an electrostatic classifier (EC, TSI Model 3071). Since theory and operation of this device are discussed extensively in a variety of papers (Knutson, 1975, 1976; Hoppel, 1978), only the following points most relevant to the WS application will be mentioned: The fact that the particles are classified by electrical mobility means that, in addition to particles of the desired size, a small but not negligible percentage of larger, multiply-charged particles were included in the nominally monodisperse aerosol leaving the EC. Also, the population of particles having essentially the selected size is not truly monodisperse but has a

size distribution of finite width which depends on the flow conditions through the EC - a typical geometric standard deviation being about 1.2. Since laminar flow inside the EC is essential, the aerosol flow in and out of the EC is limited, in the present case, to about $100 \text{cm}^3 \text{s}^{-1}$. Due to this condition, it was necessary to discard excess aerosol prior to entering the EC (the filter shown in that line on Figure 2 serves to protect the flow control valve and flow meter from salt deposits). Furthermore, this flow limitation in the EC, combined with a maximum particle number concentration imposed by coagulation rates, resulted in a maximum rate on the order of 10^7s^{-1} at which "monodisperse" particles could be introduced into the sampling duct.

In order to achieve the necessary flow in the sampling duct (cf. previous section), the aerosol was injected axially into a turbulent stream of 200 to $600 \text{m} \text{min}^{-1}$ clean dilution air. The 3 cm diameter dilution and mixing section shown in the center of Figure 2 incorporated an optional 200 l buffer volume to smoothen high frequency fluctuations in aerosol concentration. Due to the large volume required, the dilution air had to be drawn from the house compressed air system which, in turn, necessitated particularly careful purification with oversized absolute and charcoal filters, especially in view of the fact that the dilution air constituted 97 to 99% of the air in the final aerosol delivered to the sampling duct. For flows under $600 \text{m} \text{min}^{-1}$, it was possible to keep the rate constant within two percent.

As the bottom part of Figure 2 indicates, the artificial aerosol was fed into the sampling duct 26 m from the first sampling outlet, just downstream of the axial fan. A 6 cm ball valve was used to seal off the ambient aerosol branch of the duct during artificial aerosol experiments.

The upper left-hand side of Figure 2 displays the location of an auxiliary aerosol generation system which was essentially a duplicate of the one described above. The auxiliary system was equipped with commercial inhalation type nebulizers and operated with filtered compressed air from the house supply. This system was used during some preliminary WS experiments and also during Experiment No. 10 where two "monodisperse" aerosols with different particle size were mixed.

In those experiments where no monodisperse aerosol was desired, the EC was by-passed as indicated by the dashed line in Figure 2. In contrast to the monodisperse case, the problem facing us here was to produce a sufficiently low particle concentration to be meaningful and acceptable for CCN instruments. Reduction of the air pressure on the atomizer to only a few psi drastically lowered the number output but, at the same time, reduced the output stability markedly. Similarly, when nearly all the aerosol originating from the atomizer was removed by aspiration into the excess aerosol line, the small differential flow into the

sampling system underwent amplification of the small irregularities in the atomizer or excess flows. However, by applying both methods in moderation it was possible to provide a final particle concentration in the sampling duct as low as 400 cm^{-3} .

For the few experiments carried out with other than soluble CCN, the atomizer in Fig. 2 was replaced with the devices shown in Figs. 3 and 4.

AgI (purified, Fisher Scientific) was aerosolized thermally by bringing a small amount of the substance ($\sim 1 \text{ g}$) slightly above its melting point (550°C) by means of the arrangement shown in Fig. 3. The AgI was placed in a tantalum boat which was mounted on two copper electrodes inside a Pyrex flask. The temperature of the Ta-boat was controlled by manually regulating the low voltage heating current. A stream of dry N_2 (several l min^{-1}) directed at the hot AgI served to quench the vapors thus inducing recondensation of AgI particles. The resulting aerosol left the generator through the side tubulation and subsequently underwent the same handling as the soluble aerosols. In order to provide assurance that no parts of the generator other than the AgI were participating in aerosol formation, the device was tested without AgI; not until reaching temperatures well above 600°C did particle generation from the hot Ta set in. While temporal stability of the polydisperse AgI aerosol was quite within the required tolerances, the monodisperse case suffered from deviations in concentration of up to 10% from the mean.

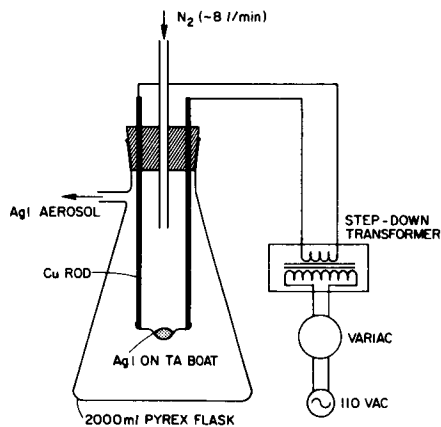


Figure 3. AgI Aerosol Generator

Preparation of the paraffin wax aerosol followed similar procedures. About 50 ml of paraffin were kept molten in a Pyrex flask at constant temperature by immersion in a bath of boiling water. A jet of dry N_2 from a capillary tube was blown vertically onto the liquid paraffin surface, again triggering particle formation through a quenching action. The polydisperse aerosol fluctuated in concentration by about 5%. Since the particle concentration was only in the vicinity of 400 cm^{-3} and most CCN instruments were able to detect only a small percentage of the hydrophobic particles, no experiment with monodisperse paraffin aerosol was carried out.

In order to facilitate interpretation of the CCN instruments' response to the test aerosols, a few WS participants measured size distributions

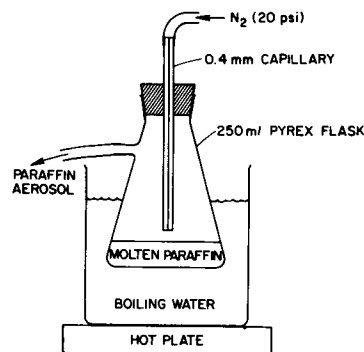


Figure 4. Paraffin Aerosol Generator

(Hoppel, 1981; Rogers, 1981) and total number concentrations (Rogers and McKenzie, 1981), while Mach and Hucek (1981) performed some chemical analyses; the reader is referred to the respective papers for additional details.

4. ACKNOWLEDGEMENTS

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ELEMENTAL COMPOSITION OF AEROSOLS IN FOURTEEN EXPERIMENTS OF THE
CLOUD CONDENSATION NUCLEI WORKSHOP

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ABSTRACT

International Cloud Condensation Nuclei (CCN) Workshop aerosols were collected with two Ci Impactors and analyzed with proton induced X-ray emission (PIXE) for chemical composition and to detect if contamination was present. One of the impactors sampled the generated aerosols; the other impactor sampled droplets from a diffusion cloud chamber. The purpose of the experiments was to test the feasibility of a study of the transfer of chemical elements from the fine particle sizes to the coarse particle sizes, after CCN are activated and cloud droplets are formed. The data indicated that sulfur-containing aerosols did exhibit the expected transfer.

1. INTRODUCTION

The aerosols generated during fourteen experiments of the International Cloud Condensation Nuclei Workshop were collected with seven stage, single orifice, Battelle-type cascade impactors (Mitchell and Pilcher, 1959) and analyzed with proton induced X-ray emission (PIXE) (Johansson et al, 1975) for chemical composition. Seven experiments were associated with the generation of aerosols of ammonium sulfate, three with sodium chloride, two with silver iodide, and two with ambient aerosols. A sample of AgI powder used to generate the aerosols was also analyzed with PIXE. Katz and Dea (1980) described the source of this AgI powder. The aerosols were analyzed for the mass in nanograms of Na, Mg, Al, Si, P, S, Cl, K, Ca, Ti, Cr, V, Mn, Fe, Ni, Cu, Zn, Br, Pb, Ag. The aerosols generated during the workshop did not always contain these elements; however, the aerosols were analyzed to detect any contamination. Several experiments were combined into one sampling period with the impactors. The aerosols were sampled in the experiments: 8-9, 10, 13-14, 15, 18-19, 20, 22, 24, 26, 27, 28.

2. DESCRIPTION OF INSTRUMENT AND ANALYSIS

Two cascade impactors were used to collect the aerosols. One impactor was connected to the tube that supplied aerosols to each experimenter, and a second impactor was placed behind a continuous flow diffusion (CFD) chamber that is similar to the CFD described by Hudson and Squires (1973, 1976). Both impactors have six stages with a Mylar film for collecting the aerosols and one stage with a 0.4 μm Nuclepore filter. The six stages separate the sampled aerosol into six size ranges with > 8 , 8-4, 4-2, 2-1, 1-0.5, 0.5-0.25 μm aerodynamic diameter. Aerosols with aerodynamic diameters less than 0.25 μm were collected on the seventh stage

with the Nuclepore filter. This Nuclepore filter limited the flow through the impactor to approximately 1 l/min. The first five Mylar films were coated with Vaseline and the sixth Mylar film was coated with paraffin. Each set of six coated Mylar films and one Nuclepore filter were matched with Mylar films and Nuclepore filter that were not exposed to the aerosol but were analyzed with PIXE to establish the background chemical composition of the filters.

The continuous flow diffusion chamber has two aluminum plates that are 50 cm long, 33 cm wide, and separated by 1.5 cm. The temperature of each plate is maintained with a circulating water bath, and the inner surface of each plate is covered with a black cotton cloth. Each cloth is wetted with water from the appropriate water bath. The chamber was normally operated with a temperature difference across the plates that produced a supersaturation of approximately 1%. The aerosol is sampled at the entrance of the chamber through a capillary tube that restricted the flow to about 0.25 l/min. A sheath flow of air maintained the aerosol in the middle of the chamber and provided air to supply the 1 l/min required for the cascade impactor at the exit port of the chamber.

In the analysis of the aerosols with PIXE, the aerosol sample is exposed to 5 MeV protons. The resulting X-ray spectrum is decomposed with a modified version of a computer program described by Kaufmann et al, 1977. Constants in this program are specified after the analysis of the spectra from a set of standards, which are samples of compounds or elements whose mass is known to within $\pm 5\%$. The analysis by PIXE can provide a mass measurement to within ± 5 nanograms.

3. DISCUSSION OF EXPERIMENTS

The purpose of these experiments with the two cascade impactors was to test the feasibility of a study of the transfer of chemical elements from the fine particle sizes, where $d < 0.5 \mu\text{m}$, to the coarse particle sizes, where $d > 4 \mu\text{m}$, after condensation of water vapor produces droplets. The chemical elements in those aerosol particles that are cloud condensation nuclei (CCN) will be transferred to the larger sizes; those elements in the aerosol particles that are not CCN, however, will remain at the smaller sizes. Several experiments with the two cascade impactors did indicate a possible transfer of sulfur. For example, a comparison of the sulfur collected on each stage of the impactor during experiments 13 and 14 is shown in Figure 1. These two experiments were sampled with the same filters to obtain more mass on each stage of the

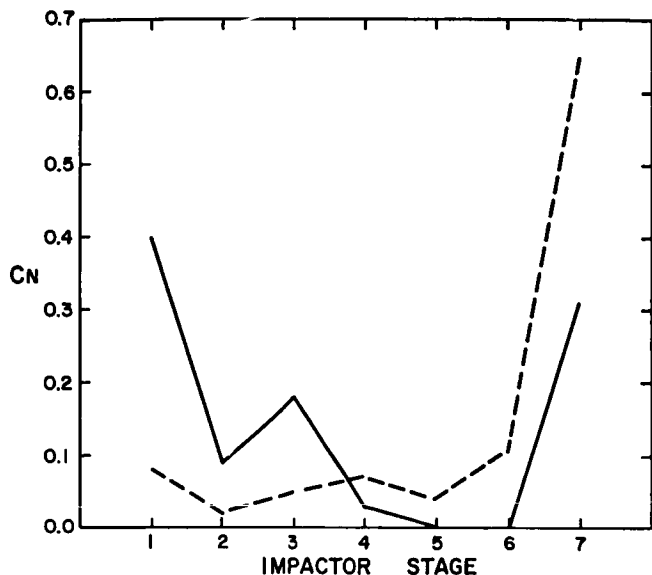


Figure 1. Comparison of mass of sulfur on seven stages of two Cascade Impactors. Stage 1 collects aerosol particles with an aerodynamic diameter $d < 0.25 \mu\text{m}$, and stage 7 collects particles with $d > 8 \mu\text{m}$. The mass collected by the impactor behind the CFD is shown with the dashed line, and the mass supplied to the CFD is shown with the solid line. The mass collected by each impactor has been normalized with the total mass collected on the impactor.

impactors. Stage 1 is the Nuclepore filter that collects aerosol particles with $d < 0.25 \mu\text{m}$, and stage seven collects particles with $d > 8 \mu\text{m}$. There is a smaller amount of sulfur, where the amount is expressed as a percent of the total mass of sulfur collected on all stages, on the Nuclepore filter and a larger amount on the seventh stage of the impactor behind the CFD.

Two problems made these experiments with the cascade impactors difficult. There was usually insufficient mass of a chemical element on a filter to permit a good determination of the total mass of each element on the filter. And some sulfur, as well as other elements, was usually found on stage seven of the impactor connected to the tube that supplied the aerosol from the aerosol generator. The CFD will be redesigned to provide more aerosol for the impactor. Furthermore, a filter can be placed in front of the CFD to prevent aerosol with $d > 8 \mu\text{m}$ from entering the chamber.

The analysis of experiment 26, which was conducted with ambient aerosols that were in the room normally used to generate the aerosols, indicated 296 nanograms of bromine on the Nuclepore of the impactor connected to the tube. The Nuclepore on the CFD had 264 nanograms. The bromine was not found on the filters that were analyzed to provide an analysis of the chemical elements in the filters. The bromine, therefore, was on aerosols with $d < 0.25 \mu\text{m}$; and these aerosols were neither deliquescent nor sufficiently hygroscopic to be CCN at the imposed supersaturation of approximately 1%.

The sample of AgI powder used to generate the AgI aerosols was analyzed with PIXE to search for contamination with trace elements. No contamination was found in the analysis of this AgI powder.

4. ACKNOWLEDGEMENT

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COMPARISON BETWEEN TWO AITKEN COUNTERS
AND WITH CLOUD CONDENSATION NUCLEI COUNTERS
AT THE 1980 INTERNATIONAL CCN WORKSHOP

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ABSTRACT

Activities at the 1980 International CCN Workshop (Reno, Nevada) included using a Pollak counter and a TSI Model 3020 Condensation Nucleus Counter to monitor the test aerosol concentration. The performance of these two counters has been intercompared and, when monodisperse CCN were used as the test aerosol, these two counters have also been compared to one of the CCN counters which gave consistently good performance and is representative of the CCN counters involved in the Workshop. The Pollak and the TSI 3020 counters agreed to within about 10% at concentrations below 1000 cm^{-3} , whereas above that concentration, substantial systematic differences were observed. When the test aerosol was monodisperse and the CCN counters were operated at supersaturations that should nucleate all the aerosol, the Pollak counter read about 6% to 20% lower than the CCN counter in the concentration range from 250 cm^{-3} to 1800 cm^{-3} . This discrepancy is similar to that reported by Emmanuel and Squires (1969). The TSI 3020 read about 20% lower than the CCN counter at concentrations less than 1000 cm^{-3} , and a factor of about two higher at concentrations above 1000 cm^{-3} . Post-workshop calibration of the Pollak counter indicates that its performance was essentially in agreement with expectations for the instrument. Post-workshop evaluation of the TSI 3020 performance suggests the cause of the poor performance above 1000 cm^{-3} and indicates the care one must use to be sure the instrument is operated in such a way as to fulfill all the theoretical assumptions its proper performance is based upon.

1. INTRODUCTION

The main purpose of the October 6-17, 1980 International Cloud Condensation Nuclei Workshop, held at Reno, Nevada, was the intercomparison and performance evaluation of instruments which utilize applied supersaturations of the order of 1%. Such supersaturations should activate most cloud condensation nuclei (CCN), i.e., those nuclei capable of initiating cloud droplet growth in atmospherically-realistic conditions. Two inherently different instruments - Aitken counters - were also operated during the Workshop; these devices detect particles which nucleate at supersaturations of the order of 200% (which includes the special class of CCN). The Aitken counter data were used to monitor the stability of the output of the Workshop's test aerosol generation facility. The main purpose of this paper, however, is to discuss the intercomparison of the two Aitken counters - a Pollak counter and a TSI Model 3020 Condensation Nucleus Counter -

and to compare them with the CCN counters when the test aerosol was a monodisperse aerosol under experimental conditions where it should be activated and detected by all the various devices.

The Nolan-Pollak Aitken counter has, in its present configuration, been in use about 20 years (Pollak and Metnieks, 1960). Detailed descriptions of this device and of its immediate predecessors are readily available (e.g., Metnieks and Pollak, 1959; Pollak and Metnieks, 1960; Pollak and Metnieks, 1957). [We shall follow the custom of referring to this device in an abbreviated manner as the "Pollak" counter, although the name of P.J. Nolan was also associated with the critical, early years of development of the instrument; see for example Nolan and Pollak (1946) - a work establishing the technical relationship and comparison of the Pollak counter to the classical Aitken counter.]

Subsequent workers examined various aspects of the operation and calibration of the Pollak counter. Kassner, et al. (1968) responded to earlier work criticizing the adiabaticity of the device, but concluded the reported discrepancies were a function of inappropriate diagnostic techniques rather than the Pollak counter itself. Emmanuel and Squires (1969) used a laboratory version Aitken counter in which activated droplets were recorded photographically to calibrate a Pollak counter over the approximate concentration range from zero to 1000 cm^{-3} . The resulting calibration curve gave concentrations about 30% higher than the curve of Pollak and Metnieks (1960). Liu, et al. (1975), responding to reports that the Pollak counter did not agree well with aerosol concentration measurements by the electrical mobility analyzer, reported careful comparison studies which indicated agreement to better than 10% between these two vastly different measurement techniques for concentrations less than 10⁴ cm^{-3} . In general, then, the Pollak counter has found widespread acceptance and use, particularly at concentrations below about 10⁴ cm^{-3} where vapor depletion by growing droplets does not significantly depress the applied supersaturation. The Pollak counter used in this study is No. 7 of the series manufactured by R. Gussman.*

The newer Aitken counter at the Workshop was a TSI Model 3020 Condensation Nucleus Counter**

* BGI Incorporated, Waltham, MA 02154

** TSI Incorporated, St. Paul, MN 55164

(Serial No. 12). This counter samples a continuous, uninterrupted flow of aerosol through a saturation chamber (using butyl alcohol as the working fluid) and a chilling section, 25°C colder than the saturator, where the supersaturation is achieved. The commercial instrument was developed directly from the laboratory design described by Bricard, et al. (1976). The principle of operation and performance were discussed at the Ninth International Nucleation Conference (Argawal, et al., 1977). To the best of our knowledge, this report is the first published critical evaluation of the TSI Model 3020.

2. EXPERIMENTAL ARRANGEMENT

The layout of the Workshop laboratory is shown in Figure 1, indicating the arrangement of the various participants along the sample supply duct. The two Aitken counters received their sample from the same sample port, which was the last port on the supply line. Other papers in this volume discuss the details and results of the other instruments, and the nature of the aerosol generation and supply system. Twenty-nine separate experiments were carried out during the course of the Workshop (one of which had to be terminated due to an accidental duct blockage). A chronological listing and description of the type of aerosol used in each experiment is indicated in Table 1.

Excluding Experiment "0", for which no Aitken data were taken, there were ten experiments involving artificial monodisperse aerosols, ten involving artificial polydisperse aerosols, and eight involving natural (ambient) polydisperse aerosols. This comparison is mainly concerned with the monodisperse aerosol experiments, in which the aerosols were obtained by passing the polydisperse output from an atomizer through an "electrostatic classifier" which selects a monodisperse fraction of aerosol based upon electrical mobility. Aerosols thus prepared are not strictly monodisperse; however,

examination of the results of these experiments in the form of cumulative CCN counts plotted as a function of individual CCN counter supersaturation setting usually shows a "plateau" or region where counts do not increase above a certain supersaturation, indicating that the size distribution is in fact very narrow. The results of these kinds of measurements are discussed in more depth in the papers describing the performance and intercomparison of the continuous-flow and static diffusion chambers. For our purposes, it is sufficient to note that when such a "plateau" exists, the nuclei concentration measured with an Aitken counter should be equal to the concentration measured on the "plateau" by a CCN counter, i.e., the existence of the "plateau" demonstrates that all the nuclei in the sample have been activated at the supersaturation setting of the CCN counter at which the "plateau" occurs and all nuclei will, of course, have been activated in the Aitken counters which operate at much higher supersaturations.

The operation of the Aitken counters was generally according to standard practice. The Pollak counter was operated according to the recommended procedure of Metnieks and Pollak (1959, op. cit.) with two exceptions: the light beam was slightly convergent (as recommended by Pollak and Metnieks (1960, op. cit.)), and 152 mm Hg over-pressure was used instead of the sea level value of 160 mm Hg in order to compensate for the effect of laboratory altitude above sea level upon the amount of liquid water released during the expansion (cf. Emmanuel and Squires, 1969, op. cit.).

The TSI 3020 CNC was operated according to manufacturer's instructions, with extra care given to maintain the sample flow rate through the instrument at the prescribed value of 5 cm³/sec. Because the instrument uses a mass flow meter to measure and control flow, it is necessary to adjust the instrument flow control in response to altitude

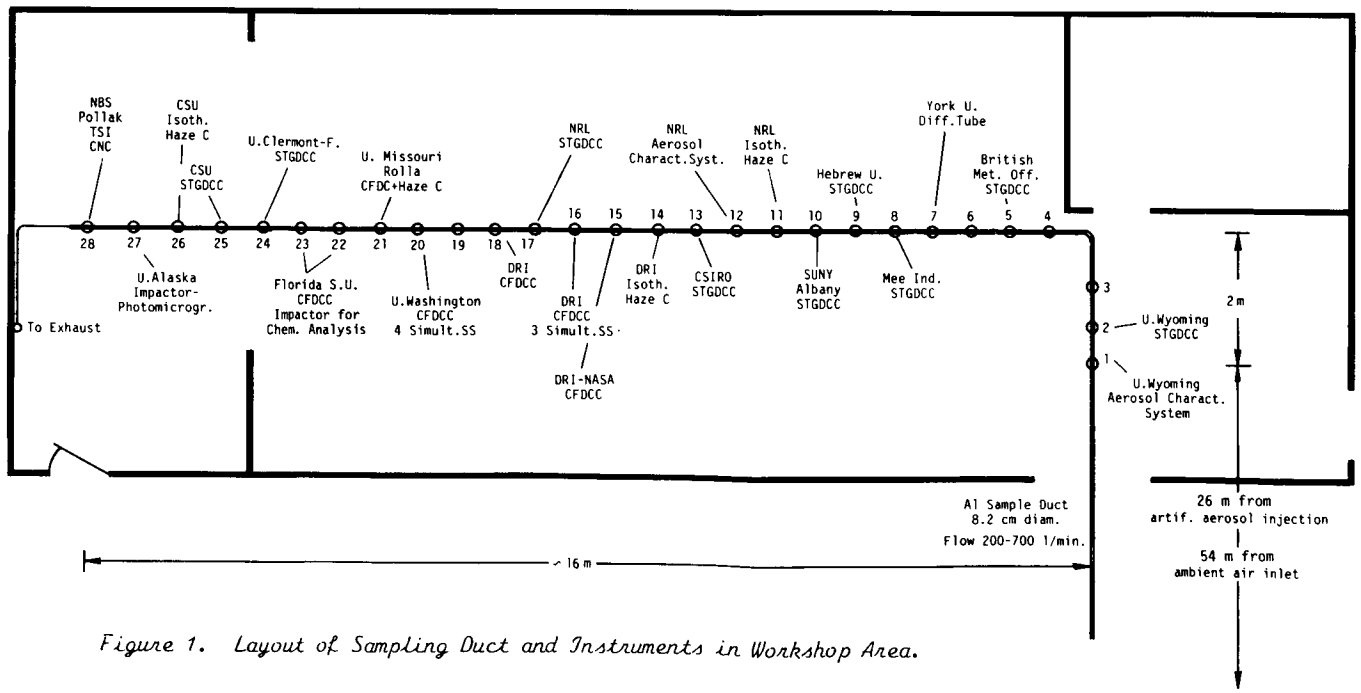


Figure 1. Layout of Sampling Duct and Instruments in Workshop Area.

TABLE 1. LIST OF EXPERIMENTS

No.	Date	Aerosol	
		A=Ambient; M=Monodisperse; P=Polydisperse	
0	Tues 7 Oct	AM	M - $(\text{NH}_4)_2\text{SO}_4$
1	Tues	PM	P - NaCl - oscillating concentration
2	Tues	PM	P - NaCl - higher concentration
3	Wed 8 Oct	AM	A - quite fluctuating
4	Wed	AM	M - NaCl - low concentration
5	Wed	PM	M - NaCl - medium concentration
6	Wed	PM	A
7	Thurs 9 Oct	AM	A - aborted - duct blockage
8	Thurs	AM	M - NaCl - slight drift down
9	Thurs	AM	M - NaCl - higher concentration
10	Thurs	PM	Bimodal - NaCl - "flat k"
11	Thurs	PM	A
12	Fri 10 Oct	AM	A
13	Fri	AM	P - $(\text{NH}_4)_2\text{SO}_4$
14	Fri	PM	P - $(\text{NH}_4)_2\text{SO}_4$
15	Fri	PM	M - $(\text{NH}_4)_2\text{SO}_4$
16	Fri	PM	A
17	Mon 13 Oct	AM	A
18	Mon	AM	M - $(\text{NH}_4)_2\text{SO}_4$
19	Mon	AM	M - $(\text{NH}_4)_2\text{SO}_4$
20	Mon	PM	M - $(\text{NH}_4)_2\text{SO}_4$
21	Mon	PM	M - $(\text{NH}_4)_2\text{SO}_4$ - time variations
22	Tues 14 Oct	AM	P - $(\text{NH}_4)_2\text{SO}_4$ - medium concentration
23	Tues	PM	P - $(\text{NH}_4)_2\text{SO}_4$ - low concentration
24	Tues	PM	P - $(\text{NH}_4)_2\text{SO}_4$ - high concentration
25	Wed 5 Oct	AM	Filtered air - noise check
26	Wed	AM	A
27	Wed	PM	P - AgI, "insoluble"
28	Wed	PM	M - AgI, "insoluble"
29	Wed	PM	P - paraffin, hydrophobic

effects on the density of air if one wants to maintain the volumetric flow rate at 5 cm³/sec. When checked with a volumetric flowmeter (a bubble meter), the volumetric flow rate through the instrument was 4.92 cm³/sec. The TSI 3020 CNC continuous real-time output, displayed on a chart recorder, provided a valuable monitor of the Workshop aerosol generation and delivery system stability.

3. EXPERIMENTAL RESULTS

A. Intercomparison of the Pollak and TSI CNC

The results of simultaneous measurements of the concentration of both polydisperse and monodisperse aerosols by the two instruments are shown in Figure 2. The results obtained with the TSI CNC are dependent upon the operating mode of the instrument, i.e., whether it was in the single-count mode or the photometric mode. The agreement between the two instruments is quite good when the TSI was operating in the single-count mode (lower curve in Figure 2), with the Pollak reading consistently about 10% lower. When the TSI operated in the photometric mode (upper curve) a nonlinear response occurred in the concentration range from 1000 cm⁻³ to 2000 cm⁻³, and above that concentration the TSI gave values twice those of the Pollak. The discontinuity in the results of the TSI between the two operating modes and the nonlinear portion of the response curve will be discussed below in the section on TSI Performance Evaluation. We have post-Workshop evidence to suggest that the Pollak performed according to expectations and we conclude that the upper curve in Figure 2 indicates that the TSI performance suffered from a systematic calibration error in the photometric mode. This error did not seriously detract from the value of the instrument as a continuous monitor of the relative aerosol concentration during the course of the various experiments performed during the Workshop.

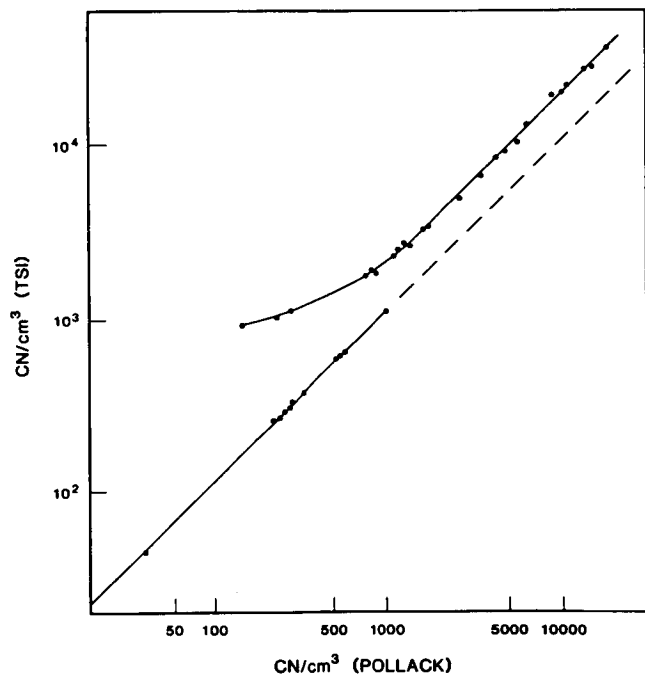


Figure 2. Comparison of the Pollak and TSI 3020 CNC Counters.

We can also compare the Aitken counters with a representative CCN counter when all three instruments were sampling monodisperse aerosols. Monodisperse sodium chloride of nominal diameter 0.036 μm (Experiments 4 and 5) and 0.18 μm (Experiments 8 and 9) and monodisperse ammonium sulfate of nominal diameters 0.092, 0.04, 0.14 and 0.08 μm (Experiments 15, 18, 19, and 20, respectively) were the test aerosols. Monodisperse silver iodide of nominal diameter 0.12 μm was the test aerosol in Experiment 28. (The tenth monodisperse aerosol experiment was Experiment 21, a repeat of No. 20 for the purpose of evaluating the measurement reproducibility of the various instruments; it is not included here because the CCN counter selected for comparison to the Aitken counters was operated as an isothermal haze chamber during that experiment.)

B. Intercomparison of the Pollak and TSI with a CCN Counter

It has not been a trivial task to select a "representative" CCN counter for use in this comparison. Many devices, both static diffusion and continuous-flow diffusion types, appeared to operate well during the course of the Workshop. We have selected the continuous-flow diffusion (CFD) chamber from the University of Missouri at Rolla for the comparison, as it consistently provided concentration values that were higher in comparison to the average of all CCN counters, but not so high as to indicate a malfunction. Results from this chamber on monodisperse aerosols usually exhibited a well-defined "plateau" in the plot of cumulative CCN concentration as a function of supersaturation value. The higher-than-overall-average count values for this chamber may suggest it had fewer sampling and detection losses. In another paper in this volume, comparing this chamber (hereinafter referred to as the UMR CFD) with the other CFD chambers in the Workshop, it was identified as showing good comparative performance (Hudson and Alofs, 1981).

Figure 3 shows a comparison of the Pollak counter and the UMR CFD using count values taken from the "plateau" regions. Error bars on the data simply indicate the magnitude of the statistical counting error. Only the lowest value shows perfect agreement (Exp. 24). The UMR CFD gives higher values by 20% at the lower concentration experiments to 6% at the higher concentration experiments. It is interesting to recall that Emmanuel and Squires (1969, op. cit.) reported that the Pollak using the 1960 calibration values, undercounted in comparison to their photographic device, by about 30% over the concentration range from about 200 cm⁻³ to 1000 cm⁻³. In a post-Workshop comparison of the DRI Pollak with another Pollak counter maintained as a standard by Dr. Austin Hogan, the DRI Pollak read high by some 10% to 20% (to be discussed below). Taking this discrepancy into consideration, the UMR CFD would be in agreement with the photographic device results reported by Emmanuel and Squires. These data, then, support the suggestion that a good CFD chamber will typically indicate on the order of 10% to 30% higher values than a Pollak counter when sampling monodisperse aerosol under conditions where the CFD is operating in the "plateau" for aerosols of varied chemical composition and of moderate concentration. (The activity of silver iodide as CCN is still a matter of some debate; here we present the one available data point (Experiment 28) and note that it is unremarkable in position on the plot.)

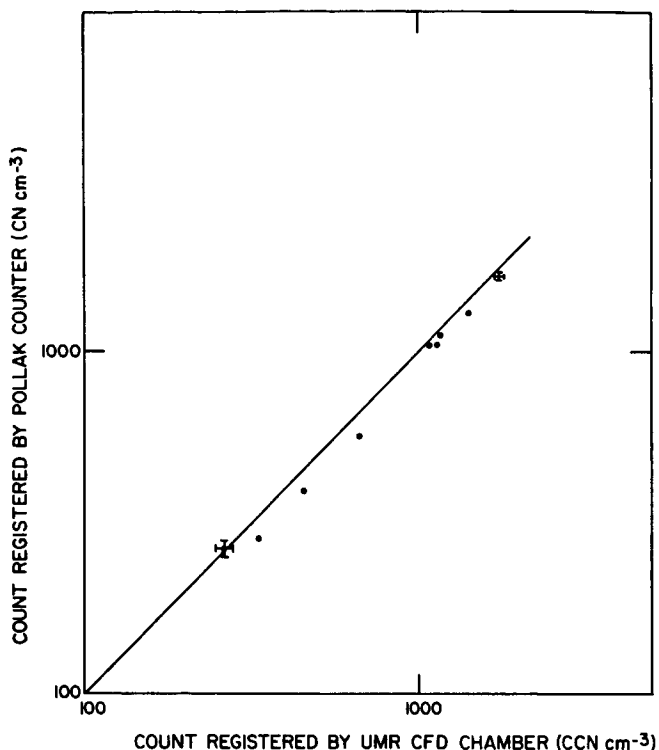


Figure 3. Comparison of the Pollak Counter and the UMR CFD CCN Counter.

A comparison of the results of the TSI CNC and the UMR CFD is shown in Figure 4 where, again, the values for the UMR CFD are the cumulative concentration values determined from the "plateau". The UMR CFD results correlate very well with the TSI (with the possible exception of silver iodide), being consistently 20% higher over the range up to 1000 cm^{-3} , where the TSI is operating in the single-count mode. As one would anticipate from the data presented in Figure 2, the TSI reads higher by a factor of about 2 when operating in the photometric mode.

4. POST-WORKSHOP CALIBRATION OF THE POLLAK COUNTER

The DRI Pollak counter (No. 7) was built and calibrated according to the published procedures of its inventors, but had not been compared against another Pollak counter in recent years before the Workshop. At the kind invitation of Drs. Jim Jiusto and Austin Hogan of the Atmospheric Sciences Research Center, State University of New York at Albany, a post-Workshop calibration was performed in February, 1981 in their laboratory. Pollak counter No. 7 was first compared to No. 12 "as is", and then was dismantled and the internal alignment of the light beam was checked. The "as is" calibration gave the result that the DRI Pollak No. 7 consistently read 10% to 20% higher than No. 12 over the concentration range for 400 cm^{-3} to 70,000 cm^{-3} . Upon dismantling and inspecting No. 7, it was found that although the light beam was of the proper diameter at the photocell end of the fog tube, it was off-axis by about 0.4 cm. Correction of this misalignment did not, however, change the systematic discrepancy that existed between the two instruments.

Several possible sources of such a systematic difference were explored, for example, by interchanging the photocells and the microammeters which provide the output signals and readings of the Pollak counters, and by interchanging operators. None of these changes removed the discrepancy. A potentially important parameter suggested in the paper of Pollak and Metnieks (1960, op.cit.) led us to investigate the rapidity with which the clean-air overpressure was established in the two instruments. Due to minor differences in the plumbing of these instruments, No. 12 achieved overpressure at a slower rate than No. 7. Standardizing both devices to the overpressure rate of No. 12 led to excellent agreement, with differences now less than would be anticipated due to statistical counting errors. It appears that the time to achieve overpressure in No. 7 (the Pollak used in the Workshop) was less than that recommended by Pollak and Metnieks (1960, op.cit.), whereas the time required for No. 12 was longer than the recommended time. We can only estimate that Pollak No. 7 was reading higher during the Workshop than an "ideal" Pollak would have, by an amount less than 10 to 20% of the number concentration.

5. TSI PERFORMANCE EVALUATION

A brief description of the operational modes and the associated underlying assumptions of the TSI CNC is necessary in order to evaluate its performance during the Workshop. The sample stream is first saturated with alcohol and then passed through a chilled condenser tube where supersaturation occurs; the nuclei are activated and the resulting droplets grow to 5 μm to 10 μm diameter. The sample of droplets then passes into a viewing/detection volume. There are two modes of operation

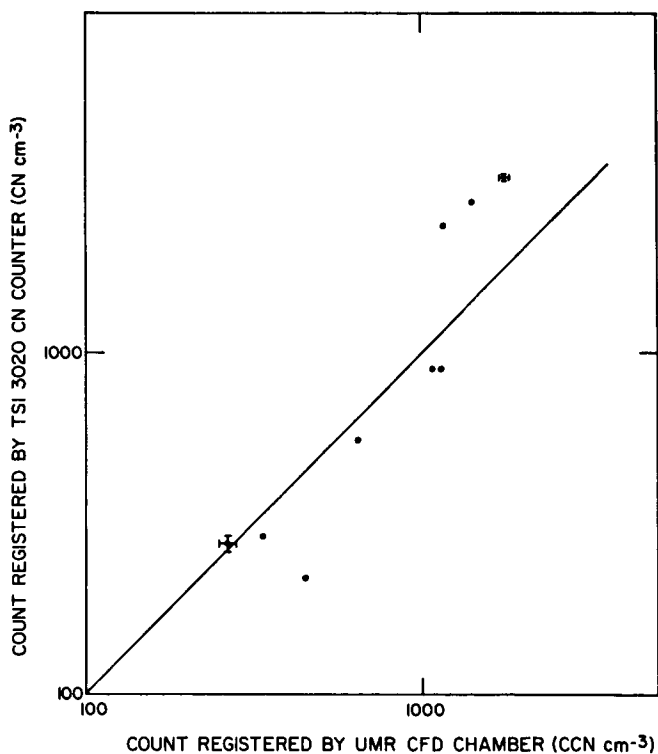


Figure 4. Comparison of the TSI 3020 CN Counter and the UMR CFD CCN Counter.

for detecting the droplets. At concentrations below 1000 cm^{-3} , electrical pulses generated by light scattered from individual droplets are counted and this count rate is converted to particle concentration and displayed on the front panel and is also available as an analog output. At particle concentrations above 10^3 cm^{-3} , the photodetector circuit measures the light scattered from all the droplets present in the viewing volume at any given time. The photodetector output is calibrated as a function of concentration, and the results of the calibration are incorporated into the electronics of the instrument so that it displays the correct concentration.

In the single count mode the instrument operates in an absolute counting mode and the accuracy of the concentration is determined only by the accuracy of the flow rate of the sample and correction for coincidence losses occurring due to two or more droplets passing through the viewing volume simultaneously. (Statistical errors of counting for low count values can be reduced to desired levels by increasing counting times.) Because the flow monitor and control system is based upon a mass flow meter, the volumetric flow rate through the instrument will be sensitive to altitude and must be corrected if the instrument is operated at altitudes other than that at which it was set up. One can either reset the flow to give $5 \text{ cc}^3/\text{sec}$ or calculate the actual volumetric flow and apply an appropriate correction factor to the indicated concentration (i.e., multiply by $[5 \text{ cm}^3 \text{ sec}^{-1} / (\text{actual volumetric flow in } \text{cm}^3 \text{ sec}^{-1})]$). The instrument we used in the Workshop was adjusted to correct the flow to $4.92 \text{ cm}^3 \text{ sec}^{-1}$.

In the photometric mode additional conditions must be met to assure that the instrument performs in calibration. Because the final droplet size determines the amount of light scattered per original activated nuclei, any parameter that affects droplet size must be controlled to maintain the conditions for which the photometric mode was calibrated. Changes in altitude in this case will affect not only the flow rate (and therefore the growth time in the condenser) but may also affect vapor diffusivity and thermal diffusivity. These effects have not yet been well studied for this instrument. A second precaution one must be aware of is the effect of dirty optics in the photodetector. In the single count mode, this will have no effect because one is just counting pulses, not scattered intensity. However, in the photometric mode all scattered light contributes to the count rate, whether it is scattered from dirty optics or from droplets in the viewing volume. The error will be most pronounced at low values in the photometric mode and will become increasingly less important as the concentration increases. We believe that this is the cause of the non-linear portion of the TSI response when compared to the Pollak in the log-log plot in Figure 2. Dirty optics will produce a constant, small D.C. offset that will be integrated into the signal in the photometric mode signal. As the total signal increases, it becomes a negligible contribution to the signal and the response then appears linearly related to the Pollak response. (Unfortunately there has not been opportunity since the Workshop to inspect the optics as they were during the Workshop in order to confirm this explanation of that portion of the curve.) The linear portion of the photometric mode response is parallel to the single count mode re-

sponse when extrapolated to the photometric mode concentration range. This suggests that the discrepancy is due to an error in the calibration factor for the instrument under the operating conditions encountered at the Workshop. One instrumental parameter that is useful in helping determine if the calibration may be in error is the photodetector pulse height. For the instrument we used, the value when the CNC was calibrated was 0.30 Volts peak-to-peak. A post-Workshop reading indicated it was 0.55 to 0.6 Volts, indicating that the instrument would not be expected to be in proper calibration. It is tempting to note that the ratio of the actual pulse height to the correct pulse height is the same as the difference between the photometric mode results and the extrapolated single count mode values, i.e., a factor of about 2. This cannot, however, be applied in such a straightforward manner because calibration will also depend on other parameters. It is sufficient here to note that there is reasonable evidence to indicate that the photometric mode results are high because the calibration is not applicable under the conditions in which we used the instrument, but that the linear response with concentration indicates that the TSI CNC will still give reliable relative concentration values under these conditions. These results again demonstrate the importance of understanding the underlying principles of the detection/measurement process of the instrument.

6. CONCLUSIONS

The performance of a Pollak counter (No. 7) of nearly standard specifications was compared with that of a set of CCN counters as represented by the UMR CFD under conditions where both were sampling the same monodisperse aerosol. Nine experiments are represented in the comparison data which indicates that the Pollak underestimates the concentration by 10% to 30% over the concentration range from about 260 cm^{-3} to 1800 cm^{-3} in comparison to the UMR CFD. This discrepancy is of the same sign, but of slightly smaller magnitude than that reported earlier by Emmanuel and Squires (1969, op.cit.) for a comparison of the Pollak counter with an "absolute" Aitken counter which recorded the droplet concentration photographically for subsequent counting.

A TSI 3020 CNC was also compared with the Pollak and the UMR CFD counters. It gave concentration values 20% lower than the UMR CFD in the concentration range from 260 cm^{-3} to 1000 cm^{-3} (i.e., when the TSI was in the single-count mode). It gave values 10% higher than the Pollak over the same concentration range. Above 1000 cm^{-3} , when the TSI operated in the photometric mode, it gave values about 2 times higher than the Pollak or the UMR CFD.

This work, then, has demonstrated very good agreement between three instruments of vastly different design and detection/measurement principles - the Pollak counter, a continuous-flow CCN counter, and a continuous-flow CNC. The present results essentially confirm the earlier work of Emmanuel and Squires (1969, op.cit.) which indicated a small positive correction to the calibration of the Pollak at aerosol concentrations below 1000 cm^{-3} . The apparent discrepancy between the Pollak and the UMR CFD is somewhat accentuated by the fact that the UMR CFD consistently read higher than the average of the other CCN counting devices in the Workshop.

The comparison study by Hudson and Alofs (1981, op.cit.) indicates that the UMR CFD was operating properly and providing accurate data. The excellent agreement with the TSI, which is a self-calibrating or absolute detector when in the single-count mode, is particularly gratifying and provides additional confidence in the validity of the large store of Aitken concentration values that have been obtained through the use of the Pollak counter over the past 20 some years.

The TSI 3020 CNC exhibited very good performance in the single-count mode and, despite loss of absolute accuracy in the photometric mode (as discussed in the text), it still provided valuable real-time relative concentration measurements which were of great utility in assessing the stability of the test aerosol generation system.

7. ACKNOWLEDGEMENTS

The authors wish to thank Dr. Austin Hogan, Mr. Stephen Barnard, and Mr. Mike Meyer for their very careful efforts in the post-Workshop calibration of the Pollak counter. This work was supported by the National Aeronautics and Space Administration under Grant No. NAS8-33820, and the National Science Foundation under Grant No. ATM79-21558.

Certain commercial equipment, instruments or materials are identified in this paper in order to adequately specify the experimental procedure. Such identification does not imply recommendation or endorsement by the National Bureau of Standards nor does it imply that the materials or equipment are necessarily the best available for the purpose.

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COMPARING DRY AEROSOL SIZE MEASUREMENTS

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Laramie, Wyoming

Two instruments were used to size dry aerosols for the CCN experiments. The Naval Research Laboratory (NRL) instrument uses an electrical mobility analyzer (Hoppel, 1978) to separate the aerosol size fractions and a diffusion chamber (Hoppel, et al., 1979) to grow and count the particles. The University of Wyoming (WYO) Aerosol Monitoring System (Rogers and Vali, 1981) consists of three devices which are monitored and controlled by a mini-computer. The devices are: a Thermo-systems Electrical Aerosol Size Analyzer (Model 3030) which covers the size range 0.005 to 0.18 μm radius, an Environment One (E1) Condensation Nucleus Monitor (Model Rich 100) for Aitken particles, and an optical particle counter (Climet Model 6064A optics and in-house electronics) to cover the size range 0.15 to 8 μm radius.

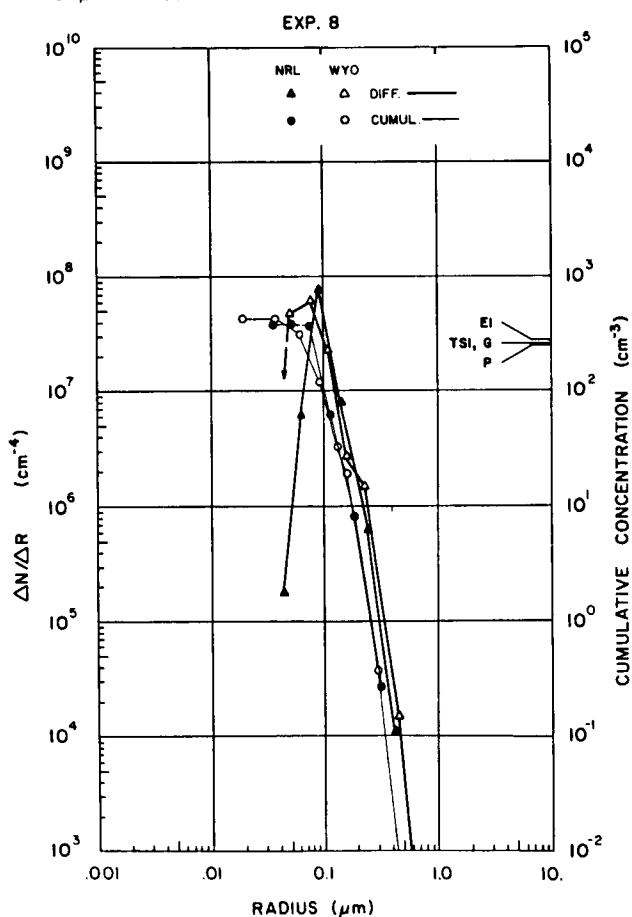


Figure 1. Aerosol size distributions of Experiment 8: average of two Naval Research Laboratory (NRL) measurements and of three University of Wyoming (WYO) measurements. Also shown are Aitken counter values (E1 = Environment One, Model Rich 100, TSI = Thermo-systems Model 3020, P = Pollak Counter and G = Gardner Counter).

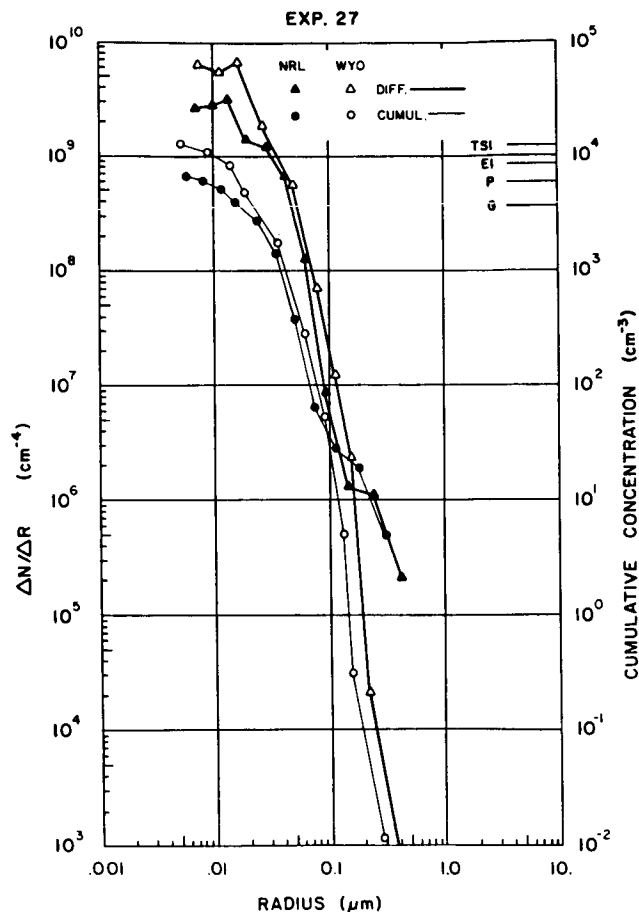


Figure 2. As for Figure 1, but aerosol from Experiment 27: average of four NRL measurements and of two WYO measurements.

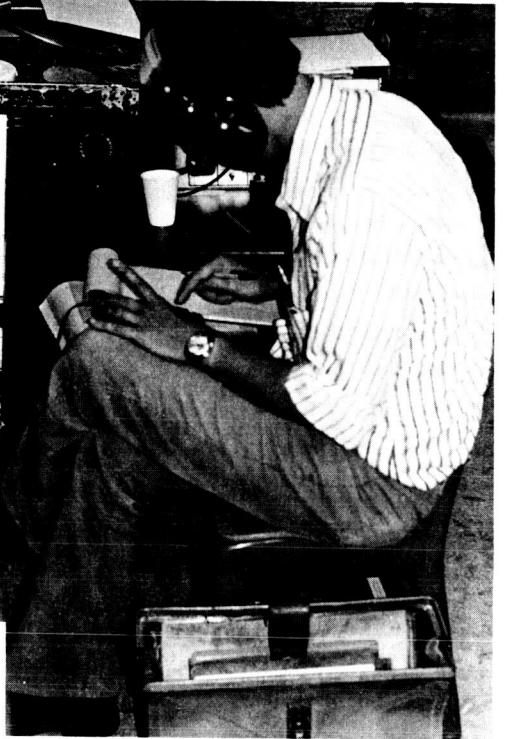
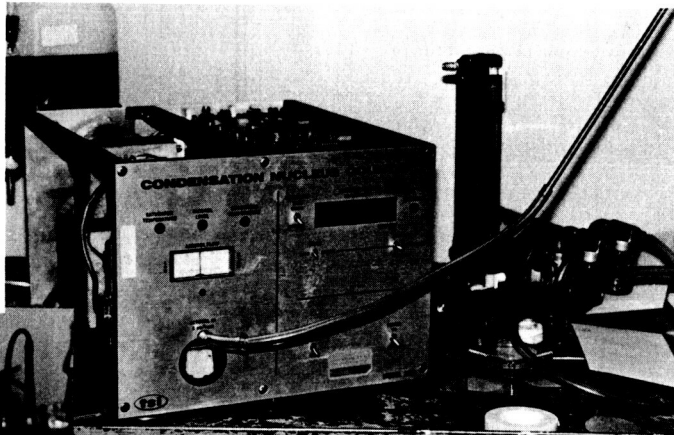
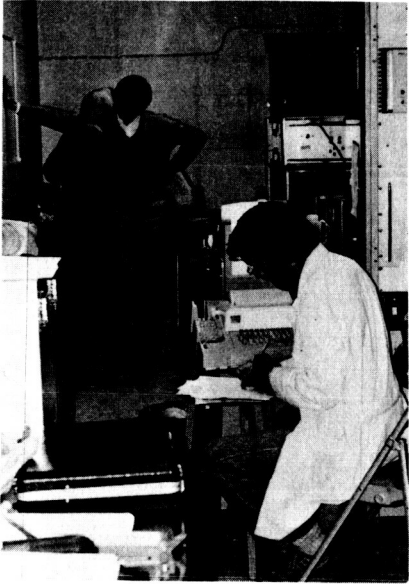
Measurements from these two systems were usually found to be in agreement during the CCN Workshop. Two examples are presented here to compare aerosol size distribution measurements of the two instruments: Experiment 8 (monodisperse NaCl) and Experiment 27 (polydisperse AgI). Differential (dN/dR) and cumulative plots are shown for both instruments for Experiment 8 in Figure 1 and for Experiment 27 in Figure 2; also shown are Aitken particle measurements for comparison. The monodisperse peak of Experiment 8 was well defined by both instruments, although information below 0.05 μm was not discernible for the WYO system; this is likely due to the low concentration, lower than the mobility analyzer is expected to detect reliably. Both instruments indicated peak concentrations in the range 0.08 to 0.1 μm radius and total particle concentrations of 400 to 500 cm^{-3} . The total concentrations from the mobility analyzers are ap-

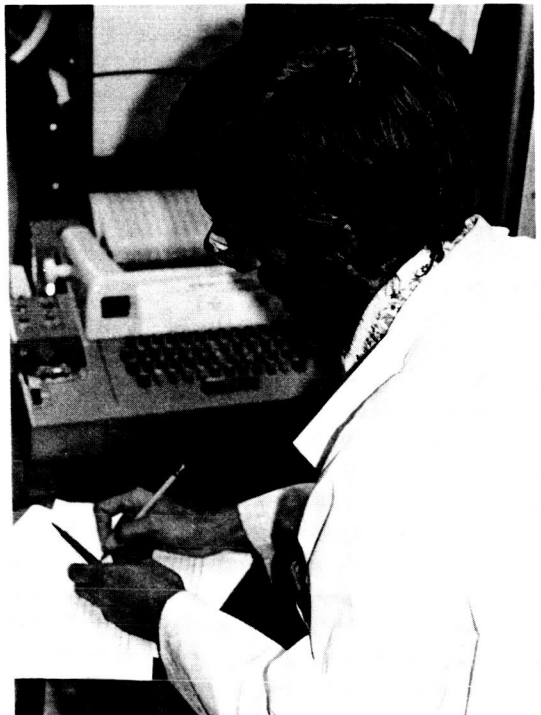
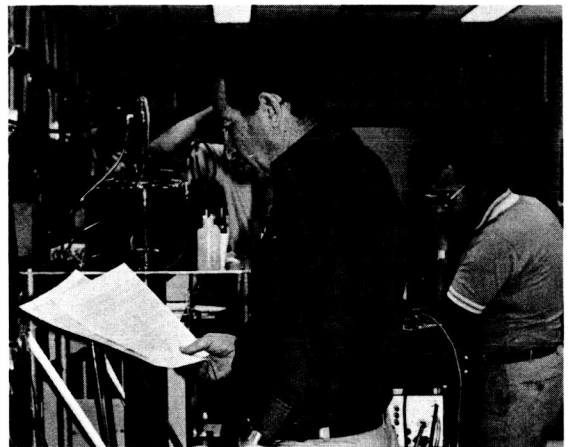
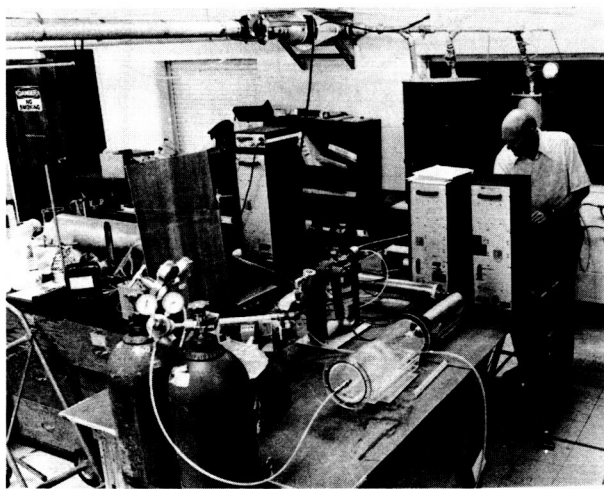
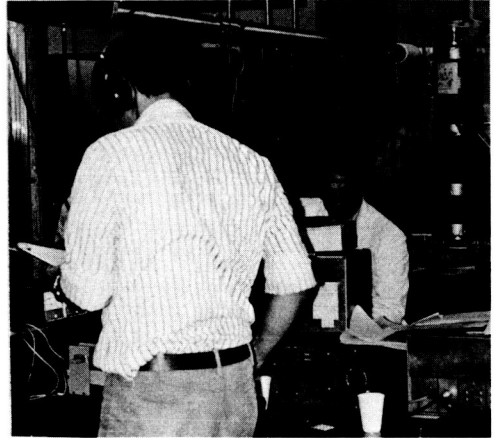
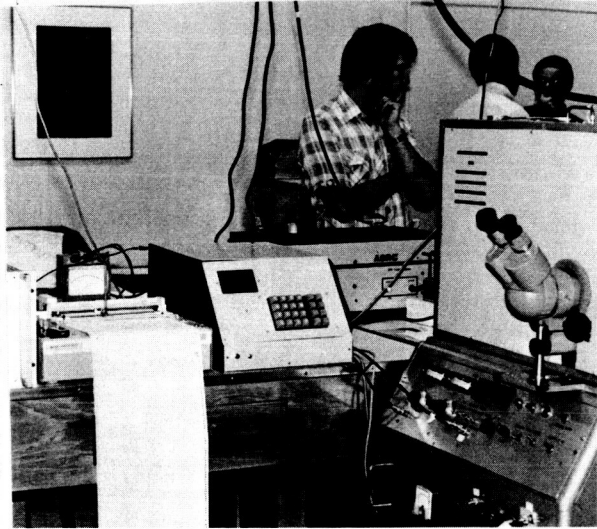
proximately twice the values obtained from the Aitken counters.

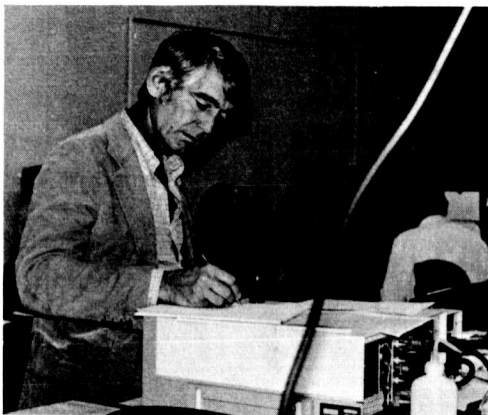
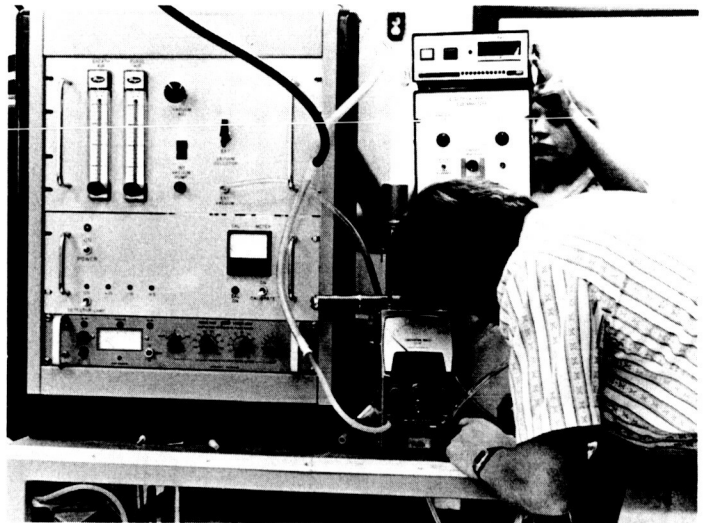
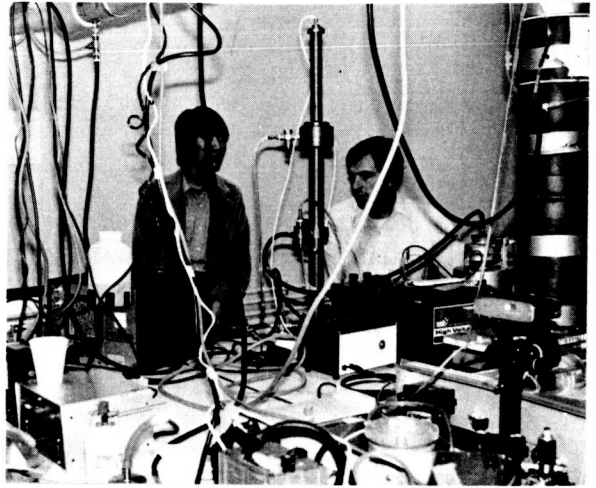
Measurements of the shape of the aerosol size distribution of Experiment 27 were in substantial agreement below $0.1 \mu\text{m}$, even to detecting a slight peak at $0.2 \mu\text{m}$ radius. At larger sizes, the NRL instrument measured a higher concentration than the WYO optical counter, and at smaller sizes, the WYO mobility analyzer measured higher concentrations than NRL. Both instruments indicated that the greatest concentrations occurred at the small sizes, but the Aitken particle counters offer little help in resolving the discrepancy in integrated total concentration, since their values range over a factor of 3. Perhaps the chemical composition and shape of the aerosol was partly responsible, as thermally generated AgI is expected to be wettable but insoluble and may have a linear rather than globular shape.

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APPENDIX A

TABULATED DIFFERENTIAL SIZE SPECTRA RESULTS FROM NRL MOBILITY ANALYZER

FOR FURTHER DETAILS, REFER TO WILLIAM A. HOPPEL PAPER,
"MEASUREMENT OF THE AEROSOL SIZE DISTRIBUTION WITH NRL'S MOBILITY ANALYZER"
CONTAINED IN SECTION VI.

NOTE THAT ABSENCE OF AN ENTRY IN DN/DR COLUMN INDICATES CONCENTRATION WAS VERY LOW OR ZERO.

Experiment #1 - Average of 3 Runs

<u>Channel</u>	<u>Cumulative Radius</u>	<u>DN/DR</u>
1	3.09×10^{-5} cm	1.69×10^5
2	1.83×10^{-5} cm	3.24×10^5
3	1.15×10^{-5} cm	4.19×10^6
4	7.53×10^{-6} cm	1.87×10^7
5	5.03×10^{-6} cm	5.80×10^7
6	3.46×10^{-6} cm	1.55×10^8
7	2.37×10^{-6} cm	2.53×10^8
8	1.52×10^{-6} cm	3.09×10^8
9	1.15×10^{-6} cm	6.05×10^8
10	8.10×10^{-7} cm	3.12×10^8
11	5.75×10^{-7} cm	2.23×10^8

Experiment #4 - Average of 4 Runs

<u>Channel</u>	<u>Cumulative Radius</u>	<u>DN/DR</u>
1	3.09×10^{-5} cm	
2	1.83×10^{-5} cm	
3	1.15×10^{-5} cm	
4	7.53×10^{-6} cm	
5	5.03×10^{-6} cm	
6	3.46×10^{-6} cm	2.70×10^5
7	2.37×10^{-6} cm	3.26×10^7
8	1.52×10^{-6} cm	3.57×10^8
9	1.15×10^{-6} cm	3.68×10^8
10	8.10×10^{-7} cm	3.54×10^7
11	5.75×10^{-7} cm	

Experiment #2 - Average of 2 Runs

<u>Channel</u>	<u>Cumulative Radius</u>	<u>DN/DR</u>
1	3.09×10^{-5} cm	2.11×10^4
2	1.83×10^{-5} cm	1.72×10^6
3	1.15×10^{-5} cm	1.07×10^7
4	7.53×10^{-6} cm	4.41×10^7
5	5.03×10^{-6} cm	1.15×10^8
6	3.46×10^{-6} cm	2.67×10^8
7	2.37×10^{-6} cm	5.14×10^8
8	1.52×10^{-6} cm	7.05×10^8
9	1.15×10^{-6} cm	1.35×10^9
10	8.10×10^{-7} cm	8.22×10^8
11	5.75×10^{-7} cm	7.14×10^8

Experiment #5 - Average of 3 Runs

<u>Channel</u>	<u>Cumulative Radius</u>	<u>DN/DR</u>
1	3.09×10^{-5} cm	
2	1.83×10^{-5} cm	
3	1.15×10^{-5} cm	
4	7.53×10^{-6} cm	
5	5.03×10^{-6} cm	
6	3.46×10^{-6} cm	3.54×10^6
7	2.37×10^{-6} cm	1.51×10^8
8	1.52×10^{-6} cm	1.55×10^9
9	1.15×10^{-6} cm	1.60×10^8
10	8.10×10^{-7} cm	6.88×10^6
11	5.75×10^{-7} cm	

Experiment #3 - One Run Only

<u>Channel</u>	<u>Cumulative Radius</u>	<u>DN/DR</u>
1	3.09×10^{-5} cm	6.40×10^6
2	1.83×10^{-5} cm	3.33×10^7
3	1.15×10^{-5} cm	9.17×10^7
4	7.53×10^{-6} cm	1.40×10^8
5	5.03×10^{-6} cm	4.44×10^8
6	3.46×10^{-6} cm	7.06×10^8
7	2.37×10^{-6} cm	8.23×10^8
8	1.52×10^{-6} cm	2.81×10^9
9	1.15×10^{-6} cm	8.97×10^9
10	8.10×10^{-7} cm	3.41×10^9
11	5.75×10^{-7} cm	9.41×10^8

Experiment #6 - Average of 3 Runs

<u>Channel</u>	<u>Cumulative Radius</u>	<u>DN/DR</u>
1	3.09×10^{-5} cm	1.06×10^5
2	1.83×10^{-5} cm	2.66×10^6
3	1.15×10^{-5} cm	3.06×10^7
4	7.53×10^{-6} cm	1.17×10^8
5	5.03×10^{-6} cm	1.74×10^8
6	3.46×10^{-6} cm	4.19×10^8
7	2.37×10^{-6} cm	9.75×10^8
8	1.52×10^{-6} cm	2.15×10^9
9	1.15×10^{-6} cm	7.26×10^9
10	8.10×10^{-7} cm	8.03×10^9
11	5.75×10^{-7} cm	2.54×10^9

Experiment #8 - Average of 2 Runs

<u>Channel</u>	<u>Cumulative Radius</u>	<u>DN/DR</u>
1	3.09×10^{-5} cm	1.08×10^4
2	1.83×10^{-5} cm	6.22×10^5
3	1.15×10^{-5} cm	7.83×10^6
4	7.53×10^{-6} cm	7.67×10^7
5	5.03×10^{-6} cm	6.13×10^6
6	3.46×10^{-6} cm	1.77×10^5
7	2.37×10^{-6} cm	
8	1.52×10^{-6} cm	8.27×10^4
9	1.15×10^{-6} cm	2.54×10^4
10	8.10×10^{-7} cm	1.76×10^2
11	5.75×10^{-7} cm	

Experiment #11 - Average of 3 Runs

<u>Channel</u>	<u>Cumulative Radius</u>	<u>DN/DR</u>
1	3.09×10^{-5} cm	1.90×10^5
2	1.83×10^{-5} cm	1.77×10^6
3	1.15×10^{-5} cm	2.81×10^7
4	7.53×10^{-6} cm	8.86×10^7
5	5.03×10^{-6} cm	1.03×10^8
6	3.46×10^{-6} cm	2.89×10^8
7	2.37×10^{-6} cm	2.35×10^8
8	1.52×10^{-6} cm	1.59×10^9
9	1.15×10^{-6} cm	5.69×10^8
10	8.10×10^{-7} cm	1.36×10^9
11	5.75×10^{-7} cm	3.95×10^9

Experiment #9 - Average of 3 Runs

<u>Channel</u>	<u>Cumulative Radius</u>	<u>DN/DR</u>
1	3.09×10^{-5} cm	4.86×10^4
2	1.83×10^{-5} cm	1.49×10^6
3	1.15×10^{-5} cm	1.58×10^7
4	7.53×10^{-6} cm	1.39×10^8
5	5.03×10^{-6} cm	8.82×10^6
6	3.46×10^{-6} cm	6.89×10^6
7	2.37×10^{-6} cm	
8	1.52×10^{-6} cm	2.21×10^5
9	1.15×10^{-6} cm	4.72×10^4
10	8.10×10^{-7} cm	3.19×10^2
11	5.75×10^{-7} cm	

Experiment #13 - Average of 3 Runs

<u>Channel</u>	<u>Cumulative Radius</u>	<u>DN/DR</u>
1	3.09×10^{-5} cm	5.49×10^4
2	1.83×10^{-5} cm	2.65×10^5
3	1.15×10^{-5} cm	4.76×10^6
4	7.53×10^{-6} cm	4.98×10^7
5	5.03×10^{-6} cm	1.93×10^8
6	3.46×10^{-6} cm	4.93×10^8
7	2.37×10^{-6} cm	1.02×10^9
8	1.52×10^{-6} cm	1.63×10^9
9	1.15×10^{-6} cm	3.39×10^9
10	8.10×10^{-7} cm	2.44×10^9
11	5.75×10^{-7} cm	2.26×10^9

Experiment #10 - Average of 2 Runs

<u>Channel</u>	<u>Cumulative Radius</u>	<u>DN/DR</u>
1	3.09×10^{-5} cm	
2	1.83×10^{-5} cm	
3	1.15×10^{-5} cm	1.33×10^6
4	7.53×10^{-6} cm	1.19×10^7
5	5.03×10^{-6} cm	3.38×10^7
6	3.46×10^{-6} cm	4.75×10^7
7	2.37×10^{-6} cm	5.06×10^7
8	1.52×10^{-6} cm	1.07×10^9
9	1.15×10^{-6} cm	8.61×10^8
10	8.10×10^{-7} cm	1.58×10^7
11	5.75×10^{-7} cm	

Experiment #14 - Average of 3 Runs

<u>Channel</u>	<u>Cumulative Radius</u>	<u>DN/DR</u>
1	3.09×10^{-5} cm	4.22×10^4
2	1.83×10^{-5} cm	6.92×10^5
3	1.15×10^{-5} cm	1.07×10^7
4	7.53×10^{-6} cm	5.86×10^7
5	5.03×10^{-6} cm	1.50×10^8
6	3.46×10^{-6} cm	6.23×10^8
7	2.37×10^{-6} cm	1.16×10^9
8	1.52×10^{-6} cm	1.54×10^9
9	1.15×10^{-6} cm	3.04×10^9
10	8.10×10^{-7} cm	1.98×10^9
11	5.75×10^{-7} cm	2.13×10^9

Experiment #15 - Average of 2 Runs

Channel	Cumulative Radius	DN/DR
1	3.09×10^{-5} cm	
2	1.83×10^{-5} cm	3.09×10^4
3	1.15×10^{-5} cm	9.70×10^5
4	7.53×10^{-6} cm	3.26×10^7
5	5.03×10^{-6} cm	1.84×10^8
6	3.46×10^{-6} cm	5.43×10^8
7	2.37×10^{-6} cm	
8	1.52×10^{-6} cm	
9	1.15×10^{-6} cm	1.93×10^6
10	8.10×10^{-7} cm	
11	5.75×10^{-7} cm	5.20×10

Experiment #19 - Average of 4 Runs

Channel	Cumulative Radius	DN/DR
1	3.09×10^{-5} cm	
2	1.83×10^{-5} cm	4.95×10^5
3	1.15×10^{-5} cm	1.29×10^7
4	7.53×10^{-6} cm	1.03×10^8
5	5.03×10^{-6} cm	3.04×10^8
6	3.46×10^{-6} cm	8.90×10^7
7	2.37×10^{-6} cm	1.49×10^7
8	1.52×10^{-6} cm	
9	1.15×10^{-6} cm	4.04×10^4
10	8.10×10^{-7} cm	6.39×10^2
11	5.75×10^{-7} cm	

Experiment #17 - Average of 2 Runs

Channel	Cumulative Radius	DN/DR
1	3.09×10^{-5} cm	1.48×10^5
2	1.83×10^{-5} cm	7.51×10^5
3	1.15×10^{-5} cm	6.71×10^6
4	7.53×10^{-6} cm	3.53×10^7
5	5.03×10^{-6} cm	1.77×10^8
6	3.46×10^{-6} cm	4.49×10^8
7	2.37×10^{-6} cm	9.97×10^8
8	1.52×10^{-6} cm	1.43×10^9
9	1.15×10^{-6} cm	4.67×10^9
10	8.10×10^{-7} cm	4.12×10^9
11	5.75×10^{-7} cm	4.23×10^9

Experiment #20 - Average of 4 Runs

Channel	Cumulative Radius	DN/DR
1	3.09×10^{-5} cm	
2	1.83×10^{-5} cm	
3	1.15×10^{-5} cm	9.62×10^4
4	7.53×10^{-6} cm	1.03×10^7
5	5.03×10^{-6} cm	7.06×10^7
6	3.46×10^{-6} cm	4.96×10^8
7	2.37×10^{-6} cm	3.14×10^7
8	1.52×10^{-6} cm	3.00×10^5
9	1.15×10^{-6} cm	
10	8.10×10^{-7} cm	
11	5.75×10^{-7} cm	

Experiment #18 - Average of 3 Runs

Channel	Cumulative Radius	DN/DR
1	3.09×10^{-5} cm	
2	1.83×10^{-5} cm	
3	1.15×10^{-5} cm	
4	7.53×10^{-6} cm	
5	5.03×10^{-6} cm	1.77×10^5
6	3.46×10^{-6} cm	1.93×10^7
7	2.37×10^{-6} cm	2.71×10^8
8	1.52×10^{-6} cm	1.23×10^9
9	1.15×10^{-6} cm	1.17×10^9
10	8.10×10^{-7} cm	3.10×10^8
11	5.75×10^{-7} cm	1.21×10^8

Experiment #21 - Average of 4 Runs

Channel	Cumulative Radius	DN/DR
1	3.09×10^{-5} cm	
2	1.83×10^{-5} cm	
3	1.15×10^{-5} cm	1.92×10^5
4	7.53×10^{-6} cm	1.05×10^7
5	5.03×10^{-6} cm	6.84×10^7
6	3.46×10^{-6} cm	4.29×10^8
7	2.37×10^{-6} cm	1.59×10^7
8	1.52×10^{-6} cm	
9	1.15×10^{-6} cm	9.44×10^5
10	8.10×10^{-7} cm	
11	5.75×10^{-7} cm	

Experiment #22 - Average of 3 Runs (2,3 and 4)
 [taken between 1100 and 1220; early run (#1)
 was taken when the count was higher
 and remains as distributed at Workshop].

Channel	Cumulative Radius	DN/DR
1	3.09×10^{-5} cm	2.11×10^4
2	1.83×10^{-5} cm	4.59×10^4
3	1.15×10^{-5} cm	3.37×10^5
4	7.53×10^{-6} cm	6.21×10^6
5	5.03×10^{-6} cm	6.29×10^7
6	3.46×10^{-6} cm	1.67×10^8
7	2.37×10^{-6} cm	3.43×10^8
8	1.52×10^{-6} cm	8.69×10^8
9	1.15×10^{-6} cm	4.13×10^8
10	8.10×10^{-7} cm	2.15×10^8
11	5.75×10^{-7} cm	1.52×10^8

Experiment #27 - Average of 4 Runs

Channel	Cumulative Radius	DN/DR
1	3.09×10^{-5} cm	2.11×10^5
2	1.83×10^{-5} cm	1.11×10^6
3	1.15×10^{-5} cm	1.32×10^6
4	7.53×10^{-6} cm	8.76×10^6
5	5.03×10^{-6} cm	1.24×10^8
6	3.46×10^{-6} cm	6.60×10^8
7	2.37×10^{-6} cm	1.20×10^9
8	1.52×10^{-6} cm	1.41×10^9
9	1.15×10^{-6} cm	3.16×10^8
10	8.10×10^{-7} cm	2.76×10^9
11	5.75×10^{-7} cm	2.66×10^9

Experiment #23 - Average of 4 Runs

Channel	Cumulative Radius	DN/DR
1	3.09×10^{-5} cm	2.64×10^4
2	1.83×10^{-5} cm	
3	1.15×10^{-5} cm	8.20×10^4
4	7.53×10^{-6} cm	1.25×10^6
5	5.03×10^{-6} cm	1.64×10^7
6	3.46×10^{-6} cm	6.55×10^7
7	2.37×10^{-6} cm	1.19×10^8
8	1.52×10^{-6} cm	1.19×10^8
9	1.15×10^{-6} cm	1.50×10^8
10	8.10×10^{-7} cm	5.93×10^7
11	5.75×10^{-7} cm	5.32×10^7

Experiment #28 - Average of 2 Runs

Channel	Cumulative Radius	DN/DR
1	3.09×10^{-5} cm	
2	1.83×10^{-5} cm	
3	1.15×10^{-5} cm	
4	7.53×10^{-6} cm	1.78×10^7
5	5.03×10^{-6} cm	1.65×10^8
6	3.46×10^{-6} cm	2.23×10^7
7	2.37×10^{-6} cm	
8	1.52×10^{-6} cm	3.72×10^6
9	1.15×10^{-6} cm	8.17×10^6
10	8.10×10^{-7} cm	
11	5.75×10^{-7} cm	

Experiment #24 - Average of 4 Runs

Channel	Cumulative Radius	DN/DR
1	3.09×10^{-5} cm	
2	1.83×10^{-5} cm	3.09×10^4
3	1.15×10^{-5} cm	7.78×10^5
4	7.53×10^{-6} cm	1.59×10^7
5	5.03×10^{-6} cm	2.78×10^8
6	3.46×10^{-6} cm	1.72×10^9
7	2.37×10^{-6} cm	4.04×10^9
8	1.52×10^{-6} cm	3.60×10^9
9	1.15×10^{-6} cm	5.28×10^9
10	8.10×10^{-7} cm	2.52×10^8
11	5.75×10^{-7} cm	1.15×10^9

APPENDIX B

In this section, the complete 1980 International CCN Workshop data file is presented, in the form of both tables and graphs of CCN counted as a function of supersaturation setting, for all experiments. The reader is urged to consult the companion review papers (covering the static diffusion chambers, continuous-flow chambers, and isothermal haze chambers by generic type) for important commentaries and points necessary to interpret these data. For example, as discussed in "Review of Isothermal Haze Chamber Performance", the interpretation of haze chamber results is dependent upon understanding the method used to calibrate the chamber optical counters for the sizing of water droplets. Useful summaries of the data will also be found in the review papers; discussions of the methods of obtaining data from the various instruments will generally be found in the relevant instrument description papers in Section V.

The data file is presented exactly as the data were received during the Workshop, with the exception of three post-Workshop revisions, all of which were cases of straightforward errors in the reporting of data. These revisions occurred in the cases of data from instruments #13 (CSIRO, G. Ayers), #14 (DRI, J. Hudson), and #25 (CSU, R. Borys); below we summarize, in the words of each instrument operator, the change in data together with reasons.

Instrument #13 - Ayers

I learned upon my return to CSIRO that the newly-generated calibration curves relate "film count" to "peak reading", and that to get to an estimate of the "true" CCN concentration, the "film

count" must be increased by 50%. Thus all the concentration values I fed into your data system should be multiplied by a factor of 1.5. This factor is explained at more length in the notes. I regret that I was unaware of it and hope that it does not cause too much inconvenience.

Instrument #14 - Hudson

In the process of writing the review paper on the IHC's, I looked back into my original Workshop notebook. I discovered that, from experiment #12 to the end of the Workshop, the threshold supersaturations for channel 3 were mislabeled in the data file. According to the original notebook, this should have been 0.11%. However, it was written into the Workshop data sheets as 0.085% which it had actually been set at for a couple of earlier experiments.

Instrument #25 - Borys

CCN concentrations were revised for the CSU static diffusion chambers for experiments 17-20. The changes were made because of errors in reading the original photographic film (i.e., counting the droplet images visually) by being out of sequence one frame for a 36 exposure roll of film. The results were spectra shifted to the left on plots done at DRI, thus artificially increasing the concentrations over all supersaturations. This was easily done since there were plateaus in the curves and the error wasn't noticed until the data were reviewed in detail after returning to CSU. The net effect is to simply correct a "human error". There was no physical instrumental correction made.

CCN WORKSHOP
NUCLEI MEASUREMENTS DATA SHEET

Experiment No: _____

Date: _____ / _____ / _____
(yr) (mo) (day)

Purpose: _____

Run No: ¹ _____ Instrument No: _____ Duct No: _____ Observer: _____

NUCLEI MEASUREMENTS

Sample Time		Plate Temp °C		Volume Sampled (ℓ)	S _c (%)	Nuclei Count	\bar{N}	Est K	Remarks ²	
Start	End	Cold	Hot							
										1
										2
										3
										4
										5
										6
										7
										8
										9
										10

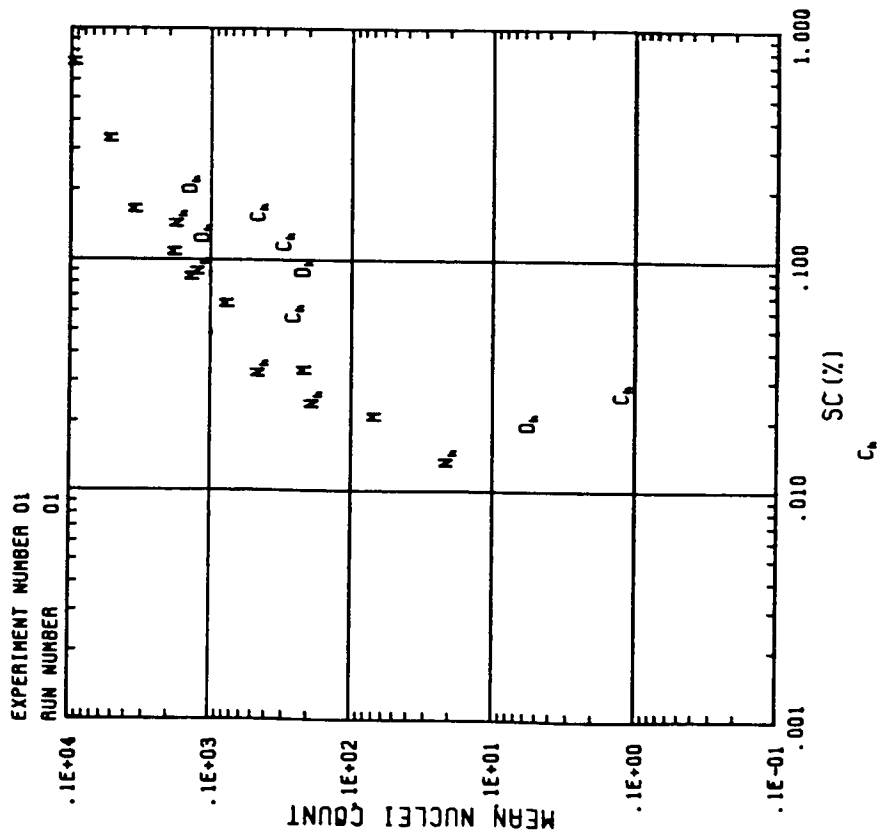
¹If more than one spectrum is run, please record the data on another sheet with a new run number.

²Only the first 25 characters of the remarks column will be recorded and displayed via computer processing.

CCN WORKSHOP DATA CODE

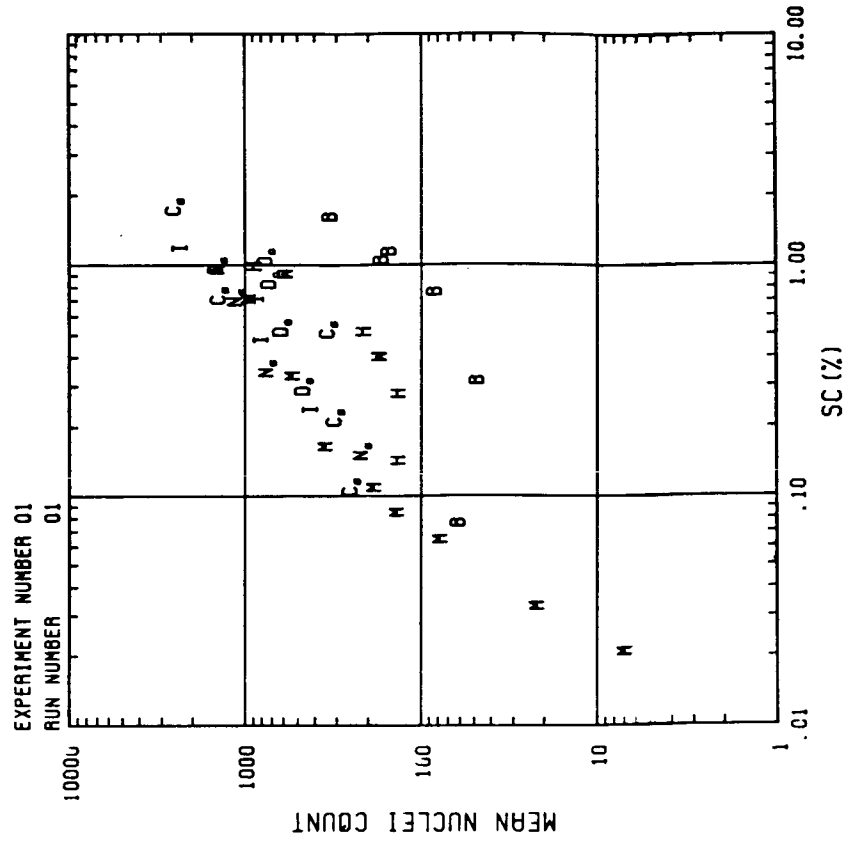
<u>Inst. No.</u>	<u>Group</u>	<u>Graphics Code</u>
2	Wyoming	O
5	British Met Office	B
7	York Univ.	Y
9	Hebrew Univ.	H
10	SUNY	S
11	NRL	Nh (Haze)
13	CSIRO	I
14	DRI	Dh (Haze)
15	DRI	Dn (Spec. NASA)
16	DRI	Di (Spec. Instant)
17	NRL	Ns (SDC)
18	DRI	Dc (CFD)
20	Washington	W
21	Missouri	M
22	Florida	
23	Florida	
24	France	F
25	CSU	Cs (SDC)
26	CSU	Ch (Haze)
27	Alaska	A
28	NBS	Gt
29	NBS	Gp

INSTRUMENT COMPARISON



HAZE CHAMBER RESULTS

INSTRUMENT COMPARISON



SIDC/CFD RESULTS

EXPERIMENT # 01

PURPOSE

INSTRUMENT COMPARISON WITH POLYDISPERSE ARTIFICIAL CCN

DESCRIPTION OF EXPERIMENT

DATE 07 OCT 1980 1458 1615

NUCLEI TYPE NA CL, POLYDISPERSE

GENERATION METHOD

DRI ATOMIZER, 18 PSI

SIZE DISTRIBUTION SHAPING

NONE

REMARKS

OUTPUT FLUCTUATES +- .8% SAMPLE WET BULB 55 (F); DRY BULB 81 (F), DIL. AIRFLOW 670 L/MIN.

WEATHER SYNOPSIS

NA

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS		OUTSIDE CONDITIONS			
AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)
NA	NA	NA	NA	NA	NA
					AVG WD (DEG)
					NA

PROCESSING DATE: 8/10/729

NUCLEI MEASUREMENTS

EXPERIMENT #: 01

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	SAMPLE TIME END	COLD	PLATE TEMP HOT (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	ORS N	EST K	REMARKS
RUN: 1															
5	5	KITCHEN		1	1509		25.60	27.00	1.70	1.70	.08	62			
5	5	KITCHEN		2	1531		24.70	27.50	3.10	1.70	.33	64			
5	5	KITCHEN		3	1532		23.80	28.00	4.20	1.20	1.2	90			
5	5	KITCHEN		4	1533		22.60	28.50	5.60	1.20	1.2	160			
5	5	KITCHEN		5	1534		21.50	28.50	5.90	1.20	1.1	177			
5	5	KITCHEN		6	1535		20.20	28.50	5.30	1.20	1.1	177			
5	5	KITCHEN		7	1537		22.00	28.50	3.20	1.20	.33	67		1.10	
5	5	KITCHEN		8	1537		26.00	28.70	2.70	1.20	.08	67			
RUN: 2															
5	5	KITCHEN		1	1538		26.00	29.00	3.00	1.20	.33	51			
5	5	KITCHEN		2	1539		24.70	29.00	4.30	1.20	1.2	114			
5	5	KITCHEN		3	1540		22.30	29.00	5.60	1.20	1.7	250			
5	5	KITCHEN		4	1541		22.30	29.00	6.70	1.20	1.7	463			
5	5	KITCHEN		5	1542		22.30	29.00	5.50	1.20	1.1	232			
5	5	KITCHEN		6	1543		22.60	29.00	4.20	1.20	.33	77			
5	5	KITCHEN		7	1544		22.70	28.70	3.00	1.20	.08	70		1.10	
5	5	KITCHEN		8	1546		26.80	28.50	1.70	1.20	.08	70			
RUN: 1															
9	9	GAGIN		1	1541		20.30	25.40	5.10	1.20	1.04	920			VOLUME SAMPLE ALWAYS THE SAME - CONSTANT
9	9	GAGIN		2	1542		21.90	25.50	3.60	1.20	.54	220			
9	9	GAGIN		3	1543		23.70	25.50	1.80	1.20	.15	140			
9	9	GAGIN		4	1544		20.40	25.40	5.00	1.20	.51	890			
9	9	GAGIN		5	1545		22.20	25.70	2.70	1.20	.16	170			
9	9	GAGIN		6	1546		23.80	25.60	1.80	1.20	.16	180			
RUN: 3															
9	9	GAGIN		1	1545		20.50	25.30	4.90	1.20	.93	620			
9	9	GAGIN		2	1545		22.80	25.30	2.70	1.20	.33	150			
RUN: 4															
9	9	GAGIN		1	1602		20.70	25.40	4.70	1.20	.88	730			
9	9	GAGIN		2	1607		22.10	25.70	3.40	1.20	.43	170			
9	9	GAGIN		3	1610		22.50	25.60	3.20	1.20	.18	140			
9	9	GAGIN		4	1610		24.00	25.60	1.60	1.20	.18	150			
RUN: 1															
11	11	HOPPEL		1	1516	1533					.15	186.5	186.5		ISOTHERMAL HAZE CHAM
11	11	HOPPEL		2	1516	1533					.094	131.2	131.2		
11	11	HOPPEL		3	1516	1533					.094	148.1	148.1		
11	11	HOPPEL		4	1516	1533					.025	19.9	19.9		
11	11	HOPPEL		5	1516	1533					.014	2.2	2.2		
RUN: 2															
11	11	HOPPEL		1	1500	1556					.15	168.1	168.1		ISOTHERMAL HAZE CHAM
11	11	HOPPEL		2	1500	1556					.094	122	122		
11	11	HOPPEL		3	1500	1556					.094	47.3	47.3		
11	11	HOPPEL		4	1500	1556					.025	17.6	17.6		
11	11	HOPPEL		5	1500	1556					.014	1.9	1.9		
RUN: 1															
13	13	AYERS		1	1521	1522	25.64	28.10	2.46		.25	480			NOISE ON COUNT RESET

PROCESSING DATE: 81/07/29

NUCLEI MEASUREMENTS

EXPERIMENT #: 01

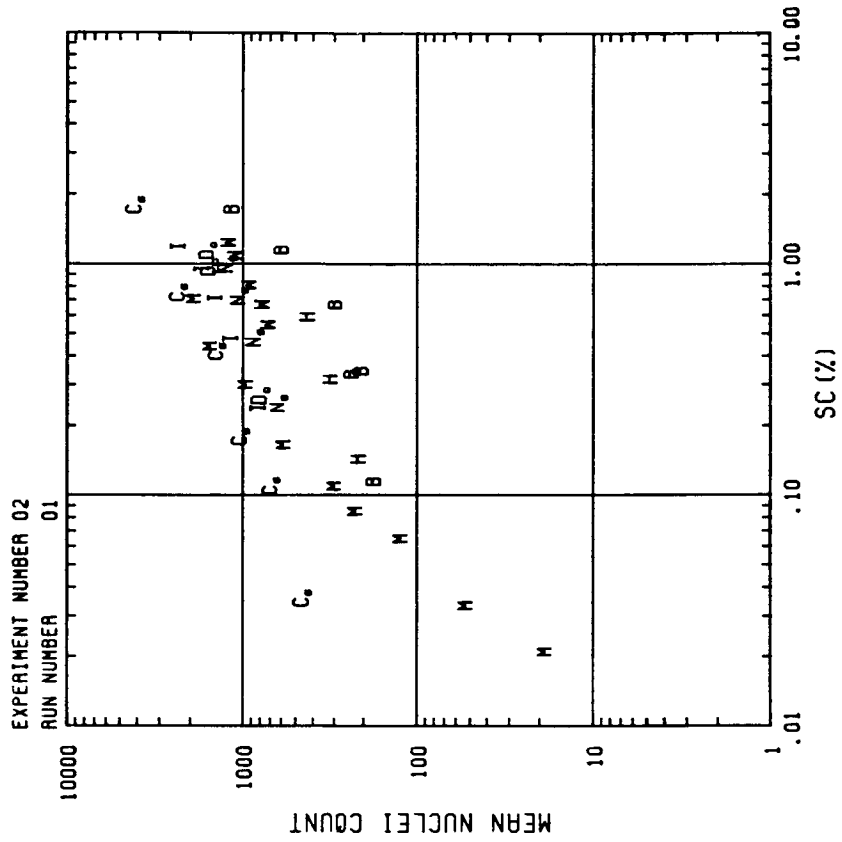
MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	COLD	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLFI COUNT	OPS - N	EST K	REMARKS
								HOT (C)	HOT (C)							
13	13	AVERS			1523	1604	24.00	28.10	5.50		.5	825			LIVE OCCASIONALLY GIVE CYCLE READING EITHER INCREASES WITH DECREASING SC.	
13	13	AVERS			1527	1608	23.00	28.20	4.97		1.23	2700			SC VALUES LISTED ARE PRESET ON INSTR. AND ARE APPROXIMATE ONLY	
13	13	AVERS			1530	1630	23.00	28.20	3.57		1.23	2700				
13	13	AVERS			1533	1633	23.00	28.20	4.97		1.23	2700				
13	13	AVERS			1533	1633	23.00	28.20	4.97		1.23	2700				
13	13	AVERS			1534	1634	23.00	28.20	5.47		1.25	1150				
13	13	AVERS			1540	1640	23.00	28.20	5.60		1.25	1150				
13	13	AVERS			1546	1646	23.00	28.20	2.55		1.25	805				
13	13	AVERS			1557	1657	23.00	28.20	5.60		1.25	2780				
13	13	AVERS			1603	1604	24.40	28.60	4.31		.75	1005				
14	14	HUDSON		9	1532	1612				.02			150			
14	14	HUDSON		9	1532	1612				.02			123			
14	14	HUDSON		9	1532	1612				.02			23.5			
17	17	WOJCIECHOWSKI		27	1542	1544							229.4		TDCC	
17	17	WOJCIECHOWSKI		27	1542	1544							791.4			
17	17	WOJCIECHOWSKI		27	1542	1544							1191.4			
18	18	HUDSON		9	1523	1527	22.00	25.70	3.75	.0024	.54	652	490			
18	18	HUDSON		9	1535	1609	21.55	26.25	4.70	.0024	.87	760	794			
18	18	HUDSON		9	1557	1607	21.20	26.45	5.25	.003	1.09					
20	20	RADKE		21	1540							180	49		HIGH SS CHANNEL FAILED RESULTS APPRX	
21	21	TRABELLO		26	1524	1525	20.70	25.00	4.30	.000	.75	1005	180			
21	21	TRABELLO		26	1529	1530	22.00	25.00	3.00	.000	.75	365	610			
21	21	TRABELLO		26	1544	1544	25.00	25.00	0.00	.000	.114	1143				
21	21	TRABELLO		26	1547	1548	25.00	25.00	0.00	.000	.189	1144				
21	21	TRABELLO		26	1554	1554	25.00	25.00	0.00	.000	.685	80.8	88			
21	21	TRABELLO		26	1554	1554	25.00	25.00	0.00	.000	.685	27.2	28			
25	25	BORYS		3	1518	1522	17.90	24.40	6.50	1.06	1.8	2688	2688		VOL SAMP X 10**5 PROBLEMS WITH LEAKS	
25	25	BORYS		3	1531	1532	20.70	24.90	4.20	2.12	.74	1485	1485			
25	25	BORYS		3	1544	1545	21.90	25.50	3.60	5.3	.53	354	354			
25	25	BORYS		3	1600	1605	23.80	26.20	2.40	10.6	.22	325	325			
26	26	HINDMAN		6	1525	1605	25.00	25.00	0.00	.83	.16	49	49		E=1.0. 0.5 UM	

EXPERIMENT #: 01
 PROCESSING DATE: 81/07/29

NUCLEI MEASUREMENTS

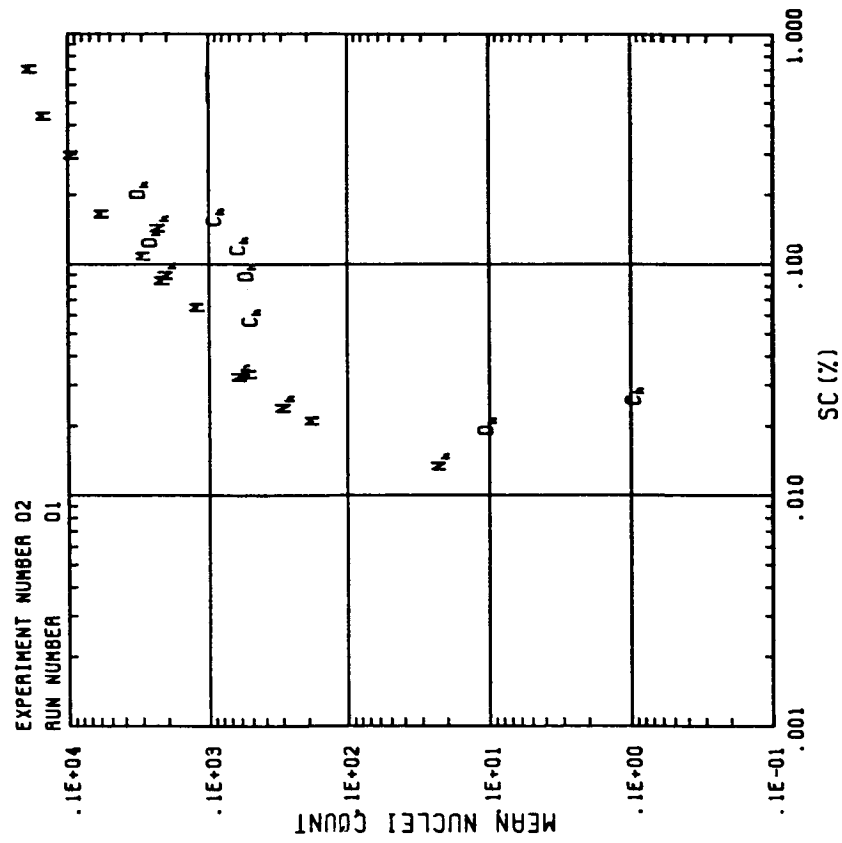
MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	DRS N	EST K	REMARKS
							COLD	HOT (C)							
26	26	HINDMAN		6	1525	1605	25.00	25.00	0.00	.83	.12	32	32		E=0.70, 0.7 UM
26	26	HINDMAN		6	1525	1603	25.00	25.00	0.00	.83	.059	27	27		E=0.64, 1.4 UM
26	26	HINDMAN		6	1525	1603	25.00	25.00	0.00	.83	.027	13	13		E=0.023, 3.0 UM
26	26	HINDMAN		6	1525	1603	25.00	25.00	0.00	.83	.016	.0024	.0024		E=0.0016, 5.0 UM
RUN: 1															
27	27	OHTAKE		19	1059	1100				8.5		.473			90% R.H.
										8.5		.508			100% R.H.

INSTRUMENT COMPARISON



SDC/CFD RESULTS

INSTRUMENT COMPARISON



HAZE CHAMBER RESULTS

EXPERIMENT # 02

PURPOSE

INSTRUMENT COMPARISON WITH POLYDISPERSE ARTIFICIAL CCN

DESCRIPTION OF EXPERIMENT

DATE	TIME START	END
07 OCT 1980	1640	1740

NUCLEI TYPE

NACL POLYDISPERSE

GENERATION METHOD

DRI ATOMIZER, 30 PSI

SIZE DISTRIBUTION SHAPING

NONE

REMARKS

STABLE OUTPUT ONLY AFTER 1700. BETTER STABILITY THAN #1 DIL. AIRFLOW 570 L/MIN.

WEATHER SYNOPSIS

NA

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS		OUTSIDE CONDITIONS			
AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)
NA	NA	NA	NA	NA	NA
					AVG WD (DEG)
					NA
					AVG WS (M/S)
					NA

PROCESSING DATE: 81/07/29

NUCLEI MEASUREMENTS

EXPERIMENT #: 02

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	SAMPLE TIME END	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OPS N	EST K	REMARKS
							COLD	HOT (C)							
RUN: 1															
5	5	KITTCHE		2	1656		27.60	29.40	1.80	.12	191				
5	5	KITTCHE		2	1657		27.90	29.50	3.10	.36	749				
5	5	KITTCHE		2	1659		27.90	29.50	6.80	1.8	1202				
5	5	KITTCHE		2	1701		22.40	29.60	5.50	1.2	1534				
5	5	KITTCHE		2	1703		22.50	29.60	4.30	.7	243				
5	5	KITTCHE		2	1704		26.70	29.70	3.00	.35	243				
5	5	KITTCHE		2	1705		27.90	29.70	1.80	.12	175			.90	
RUN: 2															
5	5	KITTCHE		2	1717		23.10	29.90	6.80	1.8	1432				
5	5	KITTCHE		2	1718		27.40	30.00	5.50	1.2	930				
5	5	KITTCHE		2	1719		27.00	30.00	3.00	.35	312				
5	5	KITTCHE		2	1721		28.20	30.00	1.80	.12	222				
5	5	KITTCHE		2	1722		26.90	30.00	3.10	.35	331				
5	5	KITTCHE		2	1723		26.70	30.00	4.30	1.2	649				
5	5	KITTCHE		2	1724		24.50	30.00	2.50	.7	421				
5	5	KITTCHE		2	1725		23.20	30.00	4.30	1.2	1451				
5	5	KITTCHE		2	1726		25.70	30.00	4.30	.7	1538			.90	
RUN: 1															
9	9	GAGIN		5	1705		20.40	25.40	5.00	1.2	1410				VOLUME SAMPLE ALWAYS THE SAME - CONSTANT
9	9	GAGIN		5	1710		23.70	25.50	2.80	.15	450				
9	9	GAGIN		5	1713		23.60	25.60	2.00	.15	320				
RUN: 2															
9	9	GAGIN		5	1717		20.00	25.00	5.00	1.2	1360				
9	9	GAGIN		5	1719		21.60	25.60	4.00	.65	390				
9	9	GAGIN		5	1721		22.20	25.40	3.20	.4	310				
RUN: 3															
9	9	GAGIN		5	1726		20.30	25.30	5.00	1.2	1690				
9	9	GAGIN		5	1729		21.60	25.40	3.80	.59	320				
9	9	GAGIN		5	1731		22.60	25.30	2.70	.73	270				
9	9	GAGIN		5	1735		23.80	25.50	1.70	.16	260				
RUN: 1															
11	11	HOPPEL		7	1704		1721				246.8			ISOTHERMAL HAZE CHAM	
11	11	HOPPEL		7	1704		1721				215				
11	11	HOPPEL		7	1704		1721				64.2				
11	11	HOPPEL		7	1704		1721				31.1				
11	11	HOPPEL		7	1704		1721				32.4				
RUN: 2															
11	11	HOPPEL		7	1728		1733				257.4			ISOTHERMAL HAZE CHAM	
11	11	HOPPEL		7	1728		1733				225				
11	11	HOPPEL		7	1728		1733				47				
11	11	HOPPEL		7	1728		1733				33.6				
11	11	HOPPEL		7	1728		1733				33.6				
RUN: 1															
13	13	AYERS		2	1704		25.70	28.20	50	.35	870			SC VALUES LISTED ARE PRESURE ON INSTR. AND ARE APPROXIMATE ONLY.	
13	13	AYERS		2	1706		25.74	28.10	3.50	.35	975				
13	13	AYERS		2	1708		24.60	28.10	3.50	.35	1290				
13	13	AYERS		2	1712		23.66	28.00	4.30	.75	1500				
13	13	AYERS		2	1714		23.70	28.00	4.30	.75	1425				

PROCESSING DATE: 81/07/29

NUCLEI MEASUREMENTS

EXPERIMENT #: 02

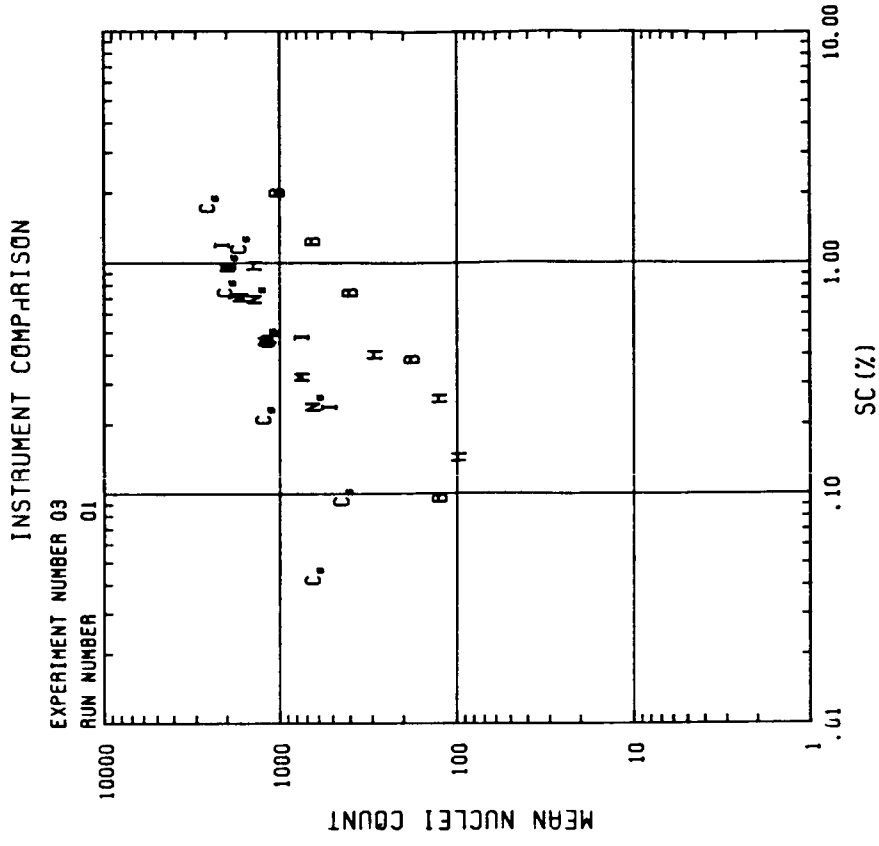
MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	SAMPLE END	COLD	HOT (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	ORS N	EST K	REMARKS
13	13	AYERS		2	1716	1717	22.91	27.90	4.99			1830			
13	13	AYERS		2	1720	1721	22.26	27.90	4.97			2010			
13	13	AYERS		2	1723	1723	22.26	27.80	5.54			2445			
13	13	AYERS		2	1725	1726	22.07	27.70	5.53			2445			
13	13	AYERS		2	1729	1730	22.62	27.60	4.98			1575			
13	13	AYERS		2	1731	1732	23.30	27.60	4.30			1575			
13	13	AYERS		2	1733	1734	24.09	27.60	3.51			1125			
13	13	AYERS		2	1735	1736	24.12	27.60	3.48			1200			
13	13	AYERS		2	1737	1737	25.20	27.70	2.50			825			
RUN: 1															
14	14	HUDSON		9	1706	1725				0.15			339		
14	14	HUDSON		9	1706	1725				0.15			379		
14	14	HUDSON		9	1706	1725				0.15			579		
14	14	HUDSON		9	1706	1725				0.15			1.11		
RUN: 1															
17	17	WOJCIECHOWSKI		27	1702	1703						660			TGDCC
17	17	WOJCIECHOWSKI		27	1702	1703						910.2			
17	17	WOJCIECHOWSKI		27	1702	1703						1117.4			
17	17	WOJCIECHOWSKI		27	1702	1703						1310			
RUN: 1															
18	18	HUDSON		9	1700	1706	21.10	26.50	5.40	0.036	1.15		1691		
18	18	HUDSON		9	1711	1714	21.30	26.25	4.95	0.024	0.97		1645		
18	18	HUDSON		9	1730	1735	22.60	25.20	2.60	0.003	0.27		840		
RUN: 1															
20	20	RADKE		21	1715	1730						750			LARGE SUPPLY VARIATN
20	20	RADKE		21	1715	1730						810			FLIGHT CHANNEL
20	20	RADKE		21	1715	1730						950			FAILED
20	20	RADKE		21	1715	1730						1250			
RUN: 1															
21	21	ALORS		1	1704	1705	20.70	25.00	4.30		74	1987			
21	21	ALORS		1	1704	1705	22.20	25.00	2.80		4.4	1503			
21	21	ALORS		1	1723	1724	22.20	25.00	0.00		3.73	612			
21	21	ALORS		1	1723	1724	22.20	25.00	0.00		1.14	231			
21	21	ALORS		1	1724	1724	22.20	25.00	0.00		0.89	231			
21	21	ALORS		1	1727	1728	22.20	25.00	0.00		0.65	54.9			
21	21	ALORS		1	1727	1728	22.20	25.00	0.00		0.22	19.5			
RUN: 1															
25	25	BORYS		3	1704	1704	19.30	26.00	6.70	1.06	1.18	4385		.62	VOL SAMP * 10**5
25	25	BORYS		3	1710	1710	21.30	25.70	4.40	1.12	1.76	2476			
25	25	BORYS		3	1715	1716	22.30	25.60	3.30	4.24	0.42	1485			
25	25	BORYS		3	1722	1723	23.30	25.50	2.20	6.36	0.16	1085			
25	25	BORYS		3	1732	1735	23.70	25.40	1.70	8.48	0.725	725			
25	25	BORYS		3	1738	1744		25.30		15.9	0.036	481			
RUN: 1															
26	26	HINDMAN		6	1708	1737	25.00	25.00	0.00	1.6	0.16	97			E=0.31, 0.5 UM

EXPERIMENT #: 02
 PROCESSING DATE: 81/07/29

NUCLEI MEASUREMENTS

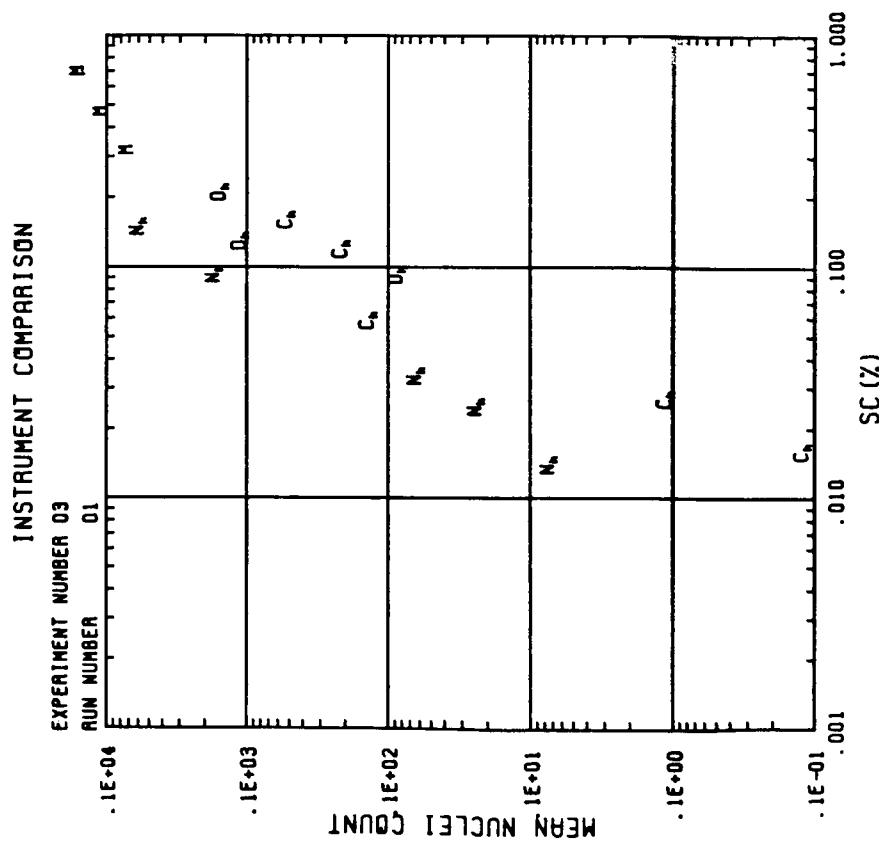
MACH NO.	DUCT NO.	OBSERVER	NO.	SAMPLE START	SAMPLE TIME END	COLD	HOT (C)	PLATE TEMP	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	ORS N	EST	REMARKS
26	26	HINDMAN	6	1708	1737	25.00	25.00	0.00	0.00	1.6	.12	66	66		E=1.2, 0.7 UM
26	26	HINDMAN	6	1708	1737	25.00	25.00	0.00	0.00	1.6	.059	54	54		E=1.1, 1.4 UM
26	26	HINDMAN	6	1708	1737	25.00	25.00	0.00	0.00	1.6	-.027	1	1		E=0.020, 3.0 UM
26	26	HINDMAN	6	1708	1737	25.00	25.00	0.00	0.00	1.6	-.016	.0013	.0013		E=0.0088, 5.0 UM
RUN: 1															
27	27	OHTAKE	19	1500	1500					8.5		.542			90% R.H.
27	27	OHTAKE	19	1500	1500					8.5		.926			100% R.H.
27	27	OHTAKE	19	1703						4.75		.202			80% R.H.
27	27	OHTAKE	19	1703						4.75		.192			90% R.H.
27	27	OHTAKE	19	1703						4.75		.253			100% R.H.
27	27	OHTAKE	19	1703						4.75		.293			101% R.H. NOMINAL

INSTRUMENT COMPARISON



SOC/CFD RESULTS

INSTRUMENT COMPARISON



HAZE CHAMBER RESULTS

PURPOSE

INSTRUMENT COMPARISON WITH AMBIENT AEROSOL

DESCRIPTION OF EXPERIMENT

DATE DATE TIME START END
 08 OCT. 1980 0805 0930

NUCLEI TYPE

AMBIENT

GENERATION METHOD

NA

SIZE DISTRIBUTION SHAPING

NA

REMARKS

HIGHLY FLUCTUATING

WEATHER SYNOPSIS

NA

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS

AVG TEMP (C) 21.0
 AVG RH (%) 20.0
 AVG PRESS (MB) NA

OUTSIDE CONDITIONS

AVG TEMP (C) NA
 AVG RH (%) NA
 AVG PRESS (MB) NA
 AVG WS (M/S) NA
 AVG WD (DEG) NA

EXPERIMENT #: 03

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/07/29

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	SAMPLE END	COLD	HOT	PLATE TEMP (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	ORS N	EST K	REMARKS
RUN: 1																
S	S	KITTCHE		1	0817	15:40	22:20	23:20	6:80	1.3	1.2	1.3	1020			
S	S	KITTCHE		1	0819	16:10	22:30	23:30	5:20	1.4	1.2	1.4	372			
S	S	KITTCHE		1	0820	19:40	22:40	23:40	3:00	1.4	1.2	1.4	131			
S	S	KITTCHE		1	0821	20:70	22:50	23:50	1:80	1.3	1.2	1.3	136			
S	S	KITTCHE		1	0823	19:60	22:60	23:60	3:50	1.3	1.2	1.3	176			
S	S	KITTCHE		1	0824	17:10	22:60	23:60	5:50	1.3	1.2	1.3	119			
S	S	KITTCHE		1	0825	15:90	22:60	23:60	6:70	1.3	1.2	1.3	64			
S	S	KITTCHE		1	0826	17:70	22:70	23:70	3:00	1.3	1.2	1.3	154			
S	S	KITTCHE		1	0827					1.3	1.2	1.3	4			
RUN: 2																
S	S	KITTCHE		2	0854	22:20	23:90	24:90	1:70	1.1	1.2	1.1	93			
S	S	KITTCHE		2	0856	21:00	24:00	25:00	3:00	1.1	1.2	1.1	11			
S	S	KITTCHE		2	0857	19:70	24:00	25:00	4:30	1.1	1.2	1.1	14			
S	S	KITTCHE		2	0858	19:60	24:10	25:10	5:30	1.3	1.2	1.3	118			
S	S	KITTCHE		2	0859	18:50	24:20	25:20	6:50	1.3	1.2	1.3	124			
S	S	KITTCHE		2	0901	20:20	24:30	25:30	4:20	1.3	1.2	1.3	238			
S	S	KITTCHE		2	0902	23:00	24:30	25:30	3:70	1.3	1.2	1.3	171			
S	S	KITTCHE		2	0905				1:70	1.3	1.2	1.3	171			
RUN: 1																
9	9	GAGIN		5	0831	20:10	25:10	26:10	5:00	1.02	1.2	1.02	1450			
9	9	GAGIN		5	0834	21:40	25:30	26:30	3:20	1.02	1.2	1.02	300			
9	9	GAGIN		5	0837	22:30	25:30	26:30	2:00	1.15	1.2	1.15	130			
9	9	GAGIN		5	0839					1.15	1.2	1.15	100			
RUN: 2																
9	9	GAGIN		5	0842	20:10	25:10	26:10	5:00	1.02	1.2	1.02	1340			
9	9	GAGIN		5	0845	21:40	25:30	26:30	3:40	1.02	1.2	1.02	290			
9	9	GAGIN		5	0848	22:40	25:20	26:20	2:20	1.11	1.2	1.11	180			
9	9	GAGIN		5	0851	23:40	25:20	26:20	1:80	1.11	1.2	1.11	180			
RUN: 3																
9	9	GAGIN		5	0915	20:20	25:20	26:20	5:00	1.01	1.2	1.01	1650			
9	9	GAGIN		5	0918	21:60	25:30	26:30	3:60	1.15	1.2	1.15	360			
9	9	GAGIN		5	0921	22:30	25:40	26:40	2:00	1.15	1.2	1.15	180			
9	9	GAGIN		5	0924	23:40	25:40	26:40	2:00	1.15	1.2	1.15	220			
RUN: 4																
9	9	GAGIN		5	0927	20:20	25:30	26:30	5:10	1.01	1.2	1.01	1320			
9	9	GAGIN		5	0930	21:70	25:30	26:30	3:70	1.15	1.2	1.15	300			
9	9	GAGIN		5	0932	22:30	25:30	26:30	2:00	1.15	1.2	1.15	40			
RUN: 1																
11	11	HOPPEL		7	0859	09:33	09:33	09:33		1.5	1.2	1.5	650			
11	11	HOPPEL		7	0859	09:33	09:33	09:33		1.03	1.2	1.03	187			
11	11	HOPPEL		7	0859	09:33	09:33	09:33		1.03	1.2	1.03	189			
11	11	HOPPEL		7	0859	09:33	09:33	09:33		1.03	1.2	1.03	62			
11	11	HOPPEL		7	0859	09:33	09:33	09:33		1.03	1.2	1.03	268			
RUN: 1																
13	13	AYERS		2	0842	21:87	24:30	24:30	2:46	1.35	1.2	1.35	540			
13	13	AYERS		2	0843	20:43	24:30	24:30	3:43	1.35	1.2	1.35	1790			
13	13	AYERS		2	0844	19:97	24:20	24:20	4:23	1.35	1.2	1.35	1830			
13	13	AYERS		2	0846	19:35	24:20	24:20	4:85	1.35	1.2	1.35	2100			

ALL SC VALUES ARE PRESET ON INSTR. AND ARE APPROXIMATE ONLY

EXPERIMENT #: 03

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/07/79

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	COLD	HOT	PLATE TEMP	DELTA	VOLUME SAMPLED (L)	SUPER SF (%)	NUCLEI COUNT	ORS N	EST K	REMARKS
13	13	ALERS		2	0853	0853	18.58	24.00	24.00	5.42	1.25	1.25	275			
13	13	ALERS		2	0854	0854	18.59	24.00	24.00	5.47	1.25	1.25	190			
13	13	ALERS		2	0855	0856	19.13	24.00	24.00	4.87	1.25	1.25	190			
RUN: 2																
13	13	ALERS		2	0906	0907	21.56	24.00	24.00	2.44	.25	.25	60			
13	13	ALERS		2	0910	0911	19.13	24.00	24.00	4.87	.25	.25	177			
13	13	ALERS		2	0911	0912	18.69	24.00	24.00	5.40	1.25	1.25	190			
13	13	ALERS		2	0915	0916	18.70	24.00	24.00	5.40	1.25	1.25	195			
13	13	ALERS		2	0918	0919	19.89	24.00	24.00	4.21	.75	.75	190			
13	13	ALERS		2	0920	0921	20.02	24.00	24.00	3.41	.5	.5	190			
13	13	ALERS		2	0922	0923	20.79	24.00	24.00	3.43	.5	.5	78			
13	13	ALERS		2	0923	0924	20.77	24.00	24.00	3.43	.5	.5	78			
RUN: 1																
14	14	HUDSON		9	0922	0922					.05	.21		170		
14	14	HUDSON		9	0922	0922					.05	.093		120		
RUN: 1																
17	17	BOUCHE	BOUCHE	27	0844	0846					.25	.48	673			
17	17	BOUCHE	BOUCHE	27	0844	0846					.25	.71	1258			
17	17	BOUCHE	BOUCHE	27	0844	0846					.25	.71	1443			
17	17	BOUCHE	BOUCHE	27	0844	0846					.25	.71	2065			
RUN: 1																
20	20	RADKE		21	0925	0927							17			
20	20	RADKE		21	0925	0927							72			
20	20	RADKE		21	0925	0927							774			
20	20	RADKE		21	0925	0927							1433			
20	20	RADKE		21	0925	0927							1725			
20	20	RADKE		21	0925	0927							152			
20	20	RADKE		21	0925	0927							1076			
RUN: 1																
21	21	ALGES		1	0827	0829	20.70	25.00	25.00	4.30	.74	.74	1755			
21	21	ALGES		1	0827	0829	22.10	25.00	25.00	2.90	.336	.336	175			
RUN: 2																
21	21	ALGES		1	0846	0847	25.00	25.00	25.00	0.00	.173	.173	1412			
21	21	ALGES		1	0846	0847	25.00	25.00	25.00	0.00	.189	.189	3258			
21	21	ALGES		1	0846	0847	25.00	25.00	25.00	0.00	.068	.068	1008			
21	21	ALGES		1	0850	0851	25.00	25.00	25.00	0.00	.022	.022	144			
21	21	ALGES		1	0850	0851	25.00	25.00	25.00	0.00	.022	.022	144			
RUN: 1																
25	25	BORS		3	0624	0630	22.80	23.90	23.90	1.10	10.6	10.6	679		.48	VOL SAMP X 10**5
25	25	BORS		3	0623	0623	20.80	23.70	23.70	1.60	8.5	8.5	467			
25	25	BORS		3	0658	0655	19.40	23.80	23.80	2.40	4.2	4.2	1291			
25	25	BORS		3	0912	0914	19.30	23.60	23.60	4.30	1.7	1.7	1222			
25	25	BORS		3	0923	0923	15.60	23.10	23.10	6.50	1.81	1.81	2688			
RUN: 1																
26	26	HINDMAN		6	0643	0931	22.60	22.60	22.60	0.00	1.5	.16	57			E=2.1, 0.5 UM

EXPERIMENT #: 03

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/07/29

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	COLD	HOT (C)	PLATE TEMP	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	ORS N	EST K	REMARKS
26	26	HINDMAN		6	0843	0931	22.60	22.60		0.00	1.5	.12	23	15		F=0.80, 0.7 UM
26	26	HINDMAN		6	0843	0931	22.60	22.60		0.00	1.5	.059	15	15		E=0.58, 1.4 UM
26	26	HINDMAN		6	0843	0931	22.60	22.60		0.00	1.5	.027	12	12		E=0.010, 3.0 UM
26	26	HINDMAN		6	0843	0931	22.60	22.60		0.00	1.5	.016	.013	.013		E=0.0030, 5.0 UM

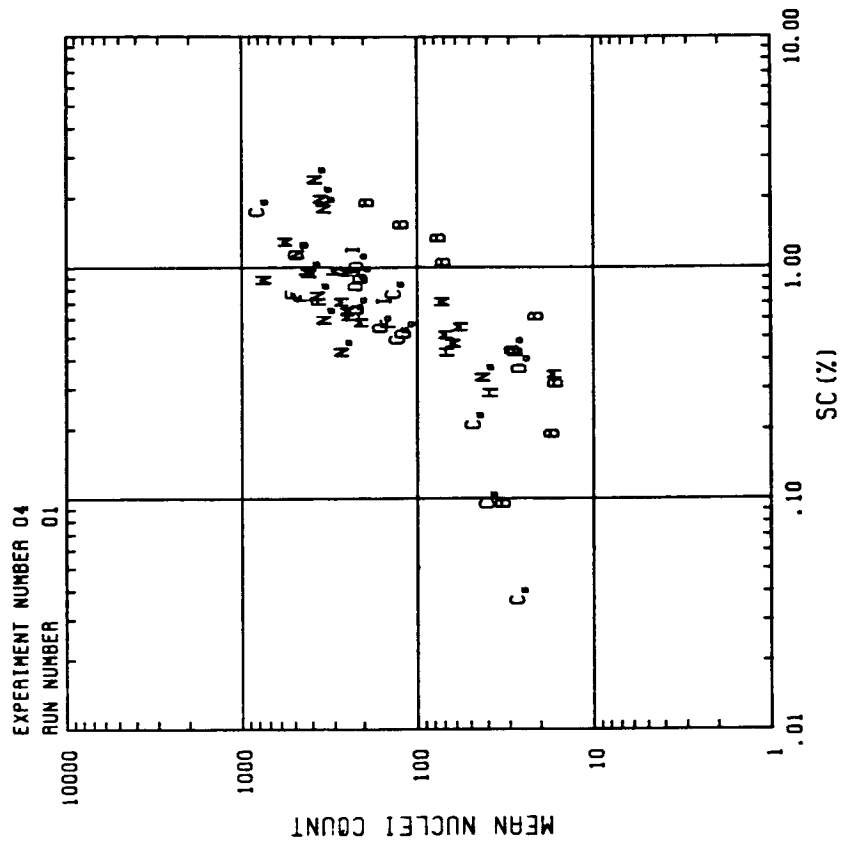
RUN: 1

27 27 OMTAKE 19 0840 0840

3.2

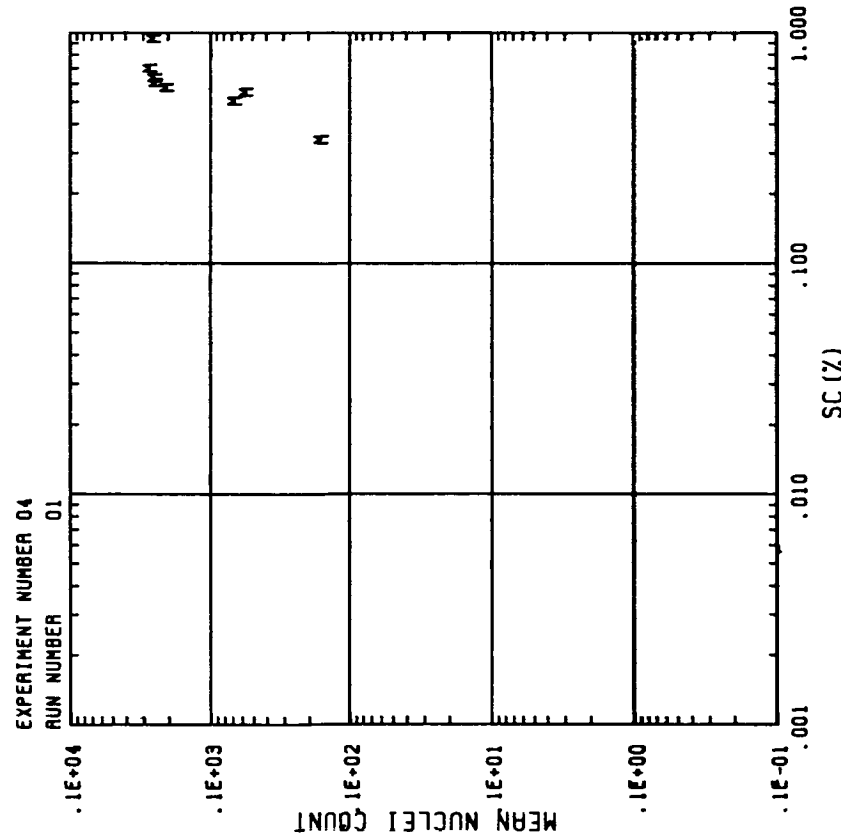
PICTURE IS NOT AVAIL

INSTRUMENT COMPARISON



SDC/CFD RESULTS

INSTRUMENT COMPARISON



HAZE CHAMBER RESULTS

EXPERIMENT # 04

PURPOSE

DETERMINE SC AGREEMENT THROUGH USE OF VERY LARGE K (SHARPLY MONODISPERSE AEROSOL).

DESCRIPTION OF EXPERIMENT

DATE DATE TIME START END
 08 OCT, 1980 0945 1338

NUCLEI TYPE

NACL, MONODISPERSE

GENERATION METHOD

DRI ATOMIZER, NO PRE-DILUTION

SIZE DISTRIBUTION SHAPING

EC; 486V, NOM. 0.036 UM DIA.
 3.3 L/MIN. SAMPLE FLOW

REMARKS

I 73 (F), ID 50 (F), DIL. FLOW 300 L/MIN.
 NOTE DISTURBANCE BETWEEN 1035 AND 1040

WEATHER SYNOPSIS

NA

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS		OUTSIDE CONDITIONS					
AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG WS (M/S)	AVG WD (DEG)
22.0	20.0	852	NA	NA	NA	NA	NA

EXPERIMENT #: 04
 PROCESSING DATE: 81/07/29

NUCLEI MEASUREMENTS

MACH NO.	DUCT NO.	OBSERVER	NO.	SAMPLE START	SAMPLE TIME END	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLFI COUNT	ORS	EST K	REMARKS
						COLD	HOT (C)							
RUN: 1														
5	5	KIICHCEN	2	1127	19:40	26:50	7:20	1.6	1.27	127				
5	5	KIICHCEN	2	1129	20:10	26:50	6:50	1.4	1.79	79				
5	5	KIICHCEN	1	1130	20:40	26:70	5:30	1.6	1.74	74				
5	5	KIICHCEN	2	1131	22:20	26:70	4:10	1.4	2.29	229				
5	5	KIICHCEN	2	1132	23:00	26:70	3:50	1.3	1.17	17				
5	5	KIICHCEN	2	1133	23:30	26:70	2:30	1.2	1.18	18				
5	5	KIICHCEN	2	1134	24:00	26:90	2:30	1.1	1.34	34				
5	5	KIICHCEN	2	1135	25:20	26:90	1:70	1.2	1.27	27				
RUN: 2														
5	5	KIICHCEN	2	1144	19:00	27:00	7:10	1.4	1.27	27				
5	5	KIICHCEN	2	1145	20:40	27:00	6:00	1.4	1.88	88				
5	5	KIICHCEN	2	1146	21:10	27:00	5:30	1.6	1.69	69				
5	5	KIICHCEN	2	1147	21:40	27:10	5:20	1.5	1.57	57				
5	5	KIICHCEN	2	1148	22:40	27:10	4:10	1.5	1.44	44				
5	5	KIICHCEN	2	1149	23:20	27:10	2:90	1.3	1.40	40				
5	5	KIICHCEN	2	1150	24:20	27:10	2:90	1.2	1.40	40				
RUN: 1														
9	9	GAGIN	5	1017	20:20	25:30	5:10	1.0	1.40	40				
9	9	GAGIN	5	1021	22:40	25:30	2:90	1.3	1.70	70				
9	9	GAGIN	5	1023	23:50	25:40	1:90	1.2	1.40	40				
RUN: 2														
9	9	GAGIN	5	1027	20:10	25:30	5:20	1.2	1.08	108				
9	9	GAGIN	5	1029	21:50	25:10	3:00	1.2	1.70	70				
9	9	GAGIN	5	1032	22:30	25:30	3:00	1.2	1.33	33				
RUN: 3														
9	9	GAGIN	5	1037	20:10	25:10	5:00	1.2	1.02	102				
9	9	GAGIN	5	1040	23:40	24:90	1:50	1.2	1.75	75				
RUN: 4														
9	9	GAGIN	5	1052	19:00	25:30	5:50	1.2	1.24	124				
9	9	GAGIN	5	1057	22:50	25:10	2:60	1.2	1.49	49				
RUN: 5														
9	9	GAGIN	5	1103	20:10	25:20	5:00	1.2	1.01	101				
9	9	GAGIN	5	1106	22:50	25:40	2:80	1.2	1.53	53				
9	9	GAGIN	5	1108	23:50	25:30	1:80	1.2	1.17	117				
RUN: 6														
9	9	GAGIN	5	1138	20:10	25:40	5:10	1.2	1.01	101				
9	9	GAGIN	5	1143	21:00	25:30	4:30	1.2	1.66	66				
9	9	GAGIN	5	1145	23:40	25:40	2:00	1.2	1.13	113				
RUN: 1														
13	13	AYERS	2	1013	25:40	26:00	3:40	1.2	1.25	125				SC VALUES ARE PRESET ON INSTR. AND ARE APPROXIMATE ONLY
13	13	AYERS	2	1015	22:00	26:10	4:10	1.2	1.5	150				
13	13	AYERS	2	1018	21:00	26:10	5:10	1.2	1.25	125				
13	13	AYERS	2	1025	20:00	26:10	4:10	1.2	1.25	125				

NUCLEI MEASUREMENTS

EXPERIMENT #: 04

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	SAMPLE TIME END	COLD	HOT (C)	PLATE TEMP	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	ORS N	EST K	REMARKS
RUN: 2																
13	13	AYERS		2	1029	1031	20.61	26.10	5.49			1.25	233			
13	13	AYERS		2	1031	1032	21.80	26.10	4.30			.75	210			
13	13	AYERS		2	1032	1033	22.10	26.10	4.51			.25	0			
13	13	AYERS		2	1033	1034	23.59	26.10	3.51				0			
13	13	AYERS		2	1034	1035	25.74	26.10	2.46				0			
RUN: 3																
13	13	AYERS		2	1035	1036	27.88	26.10	3.49			.25	0			
13	13	AYERS		2	1036	1037	29.52	26.10	3.58			.75	158			
13	13	AYERS		2	1037	1038	31.88	26.10	4.62			1	195	145		
13	13	AYERS		2	1038	1039	34.20	26.10	5.50			1.25	249			
13	13	AYERS		2	1039	1040	36.80	26.10					233			
RUN: 1																
14	14	HUDSON		9	1150	1150				.033				0		
14	14	HUDSON		9	1150	1150				.033				0		
14	14	HUDSON		9	1150	1150				.033				0		
RUN: 1																
17	17	WOJCIK		27	1138	1138						.25	0			
17	17	WOJCIK		27	1138	1138						.75	44			
17	17	WOJCIK		27	1138	1138						.98	281			
17	17	WOJCIK		27	1138	1138						1.19	384			
17	17	WOJCIK		27	1138	1138						1.87	540			
17	17	WOJCIK		27	1138	1138						2.05	820			
17	17	WOJCIK		27	1138	1138							355			
17	17	WOJCIK		27	1138	1138							420			
17	17	WOJCIK		27	1138	1138							500			
17	17	WOJCIK		27	1138	1138							370			
17	17	WOJCIK		27	1138	1138							399			
RUN: 1																
18	18	HUDSON		9	1037	1037	1.50	30.50	4.80			.94		218		
18	18	HUDSON		9	1037	1037	2.43	30.50	4.30			.68		276		
18	18	HUDSON		9	1037	1037	3.35	30.50	3.70			.40		301		
18	18	HUDSON		9	1037	1037	4.27	30.50	3.20			.27		123		
18	18	HUDSON		9	1037	1037	5.20	30.50	2.60			.18		60		
18	18	HUDSON		9	1037	1037	6.12	30.50	2.10			.09		23		
18	18	HUDSON		9	1037	1037	7.05	30.50	1.60			.06		10		
18	18	HUDSON		9	1037	1037	8.00	30.50	1.10			.04		5		
18	18	HUDSON		9	1037	1037	9.00	30.50	.60			.03		2		
18	18	HUDSON		9	1037	1037	10.00	30.50	.10			.01		0		
RUN: 1																
20	20	RADKE		11	1108	1108						.49	59			
20	20	RADKE		11	1108	1108						.33	77			
20	20	RADKE		11	1108	1108						.14	77			
20	20	RADKE		11	1108	1108						.74	82			
20	20	RADKE		11	1108	1108						1.01	307			
20	20	RADKE		11	1108	1108						1.35	590			
RUN: 2																
20	20	RADKE		11	1200	1205						.36	62			
20	20	RADKE		11	1200	1205						.54	50			
20	20	RADKE		11	1200	1205						.78	620			
20	20	RADKE		11	1200	1205						1.04	704			
20	20	RADKE		11	1200	1205						.4	48			

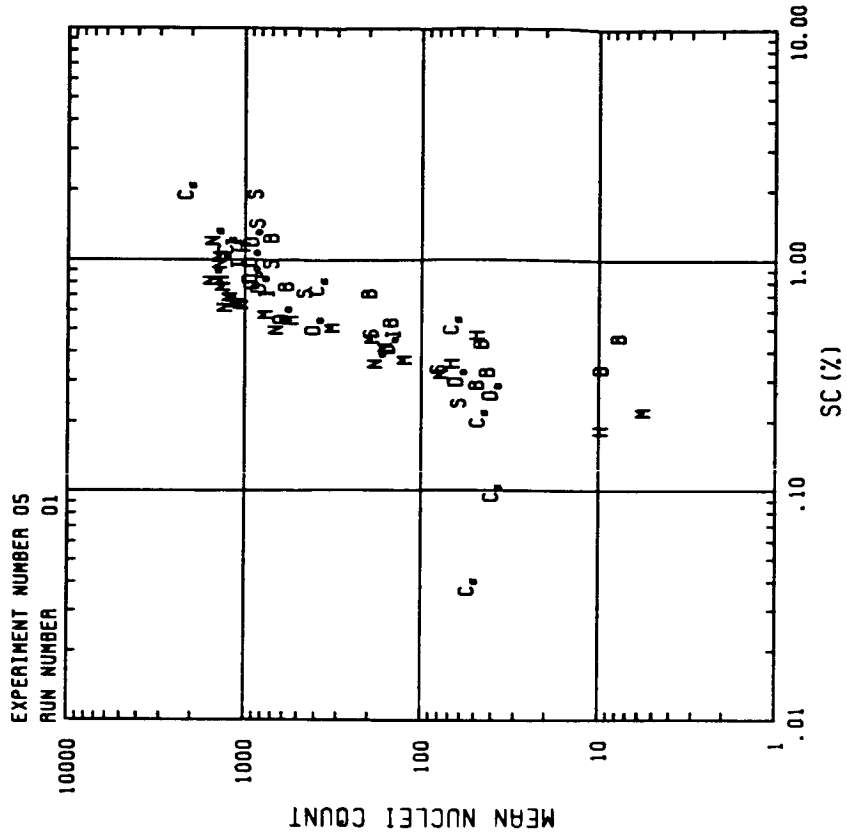
PROCESSING DATE: 81/07/29

NUCLEI MEASUREMENTS

EXPERIMENT #: 04

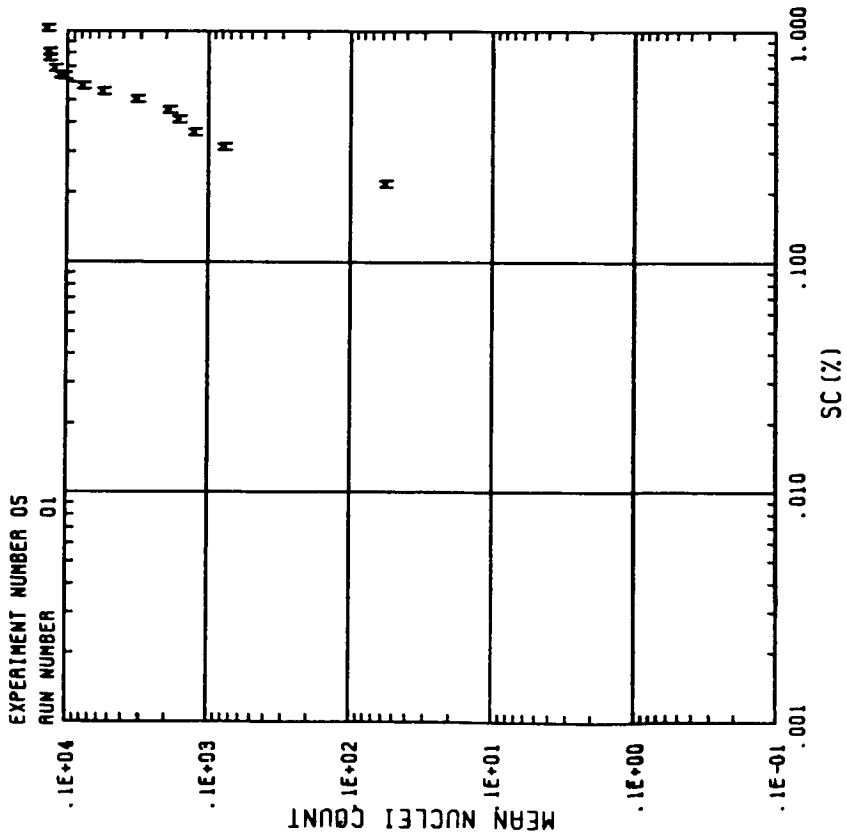
MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	SAMPLE END	COLD	HOT	PLATE TEMP (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OPS	EST K	REMARKS
20	20	RADVE		21	1205							.65	20			
20	20	RADVE		21	1205							.85	593			
20	20	RADVE		21	1205							1.04	601			
RUN: 3																
20	20	DOMONKOS		4	1304	1310						.39	41			
20	20	DOMONKOS		4	1304							.57	28			
20	20	DOMONKOS		4	1304							1.06	638			
20	20	DOMONKOS		4	1304								774			
RUN: 1																
21	21	ALOSS		1	1010	1014	20.90	25.00	25.00	5.00		.1	265.9			
21	21	ALOSS		1	1027	1030	20.90	25.00	25.00	4.70		.72	290.4			
21	21	ALOSS		1	1028	1030	20.90	25.00	25.00	3.80		.67	258.7			
21	21	ALOSS		1	1032	1036	21.20	25.00	25.00	4.00		.64	257.4			
21	21	ALOSS		1	1037	1037	21.10	25.00	25.00	3.90		.68	214.6			
21	21	ALOSS		1	1036	1041	21.35	25.00	25.00	3.65		.533	171.2			
21	21	ALOSS		1	1041	1042	22.00	25.00	25.00	3.00		.36	17.2			
RUN: 1																
24	24	SERPOLAY		25	1045	1055	23.70	27.60	27.60	3.90	.77	.6	150			
24	24	SERPOLAY		25	1115	1115	24.50	28.90	28.90	4.40	.77	.5	380			
24	24	SERPOLAY		25	1120	1135	25.00	30.00	30.00	4.60	.77	.78	570			
RUN: 1																
25	25	BORSS		3	1020	1028	21.60	25.60	25.60	1.00	.51	.38	28		.91	VOL SAMP X 10**5
25	25	BORSS		3	1022	1029	21.00	25.60	25.60	1.00	.51	.09	42			LEAK PROBLEMS
25	25	BORSS		3	1025	1043	21.00	25.60	25.60	1.00	.51	.21	50			
25	25	BORSS		3	1029	1107	21.00	25.60	25.60	1.00	.51	.25	134			
25	25	BORSS		3	1034	1107	21.00	25.60	25.60	1.00	.51	.49	141			
25	25	BORSS		3	1035	1115	21.00	25.60	25.60	1.00	.51	1.182	509			
25	25	BORSS		3	1115	1115	21.00	25.60	25.60	1.00	.51	1.182	509			
RUN: 1																
26	26	HINDMAN		6	1018	1145	22.60	25.60	25.60	0.00	1.1	.16	0			0.5 UH
26	26	HINDMAN		6	1018	1145	22.60	25.60	25.60	0.00	1.1	.12	0			0.7 UH
26	26	HINDMAN		6	1018	1145	22.60	25.60	25.60	0.00	1.1	.057	0			91.4 UH
26	26	HINDMAN		6	1018	1145	22.60	25.60	25.60	0.00	1.1	.027	0			3.0 UH
26	26	HINDMAN		6	1018	1145	22.60	25.60	25.60	0.00	1.1	.016	0			5.0 UH
RUN: 1																
27	27	OHTAKE		19	1016	1016	20.00	25.00	25.00	3.00	.32		405			94% R.H.
27	27	OHTAKE		19	1016	1016	20.00	25.00	25.00	3.00	.32		505			100% R.H.
27	27	OHTAKE		19	1016	1016	20.00	25.00	25.00	3.00	.32		505			101% R.H.
27	27	OHTAKE		19	1016	1016	20.00	25.00	25.00	3.00	.32		505			102% R.H.
27	27	OHTAKE		19	1016	1016	20.00	25.00	25.00	3.00	.32		505			NOMINAL

INSTRUMENT COMPARISON



SDC/CFD RESULTS

INSTRUMENT COMPARISON



HAZE CHAMBER RESULTS

EXPERIMENT # 05

PURPOSE

AS # 4, BUT HIGHER AEROSOL CONCENTRATION

DESCRIPTION OF EXPERIMENT

DATE DATE TIME START END
 08 OCT, 1980 1410 1606

NUCLEI TYPE

NACL, MONODISPERSE

GENERATION METHOD

DRI ATOMIZER, NO PRE-DILUTION, 70 PSI.

SIZE DISTRIBUTION SHAPING

EC, 486V (NOM. 0.036 UM DIA.)
 6.7 L/MIN. SAMPLE FLOW, 250 L/MIN DIL. FLOW.

REMARKS

T 74(F), TD 50(F)

WEATHER SYNOPSIS

NA

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS			OUTSIDE CONDITIONS				
AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG WS (M/S)	AVG WD (DEG)
23.0	21.0	852	NA	NA	NA	NA	NA

EXPERIMENT #: 05

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/07/29

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	SAMPLE END	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLFI COUNT	ORS N	EST	REMARKS
							COLD	HOT (C)							
RUN: 1															
5	5	KITTCHE	N	12	1434	21:40	24:50	3:50	1.2	0.5	48				
5	5	KITTCHE	N	12	1435	22:00	25:00	3:00	1.2	0.35	51				
5	5	KITTCHE	N	12	1436	21:50	25:00	3:10	1.2	0.35	10				
5	5	KITTCHE	N	12	1437	19:40	23:10	3:30	1.2	1.3	743				
5	5	KITTCHE	N	12	1443	20:40	23:10	2:30	1.2	0.75	208				
5	5	KITTCHE	N	12	1445	22:20	25:10	2:50	1.2	0.34	155				
RUN: 2															
5	5	KITTCHE	N	12	1451	21:30	25:60	4:30	1.2	0.68	146				
5	5	KITTCHE	N	12	1452	20:20	25:60	5:40	1.2	1.1	276				
5	5	KITTCHE	N	12	1453	19:10	25:60	6:50	1.2	1.3	540				
5	5	KITTCHE	N	12	1454	17:60	26:20	9:00	1.2	0.8	1446				
5	5	KITTCHE	N	12	1455	22:10	26:90	4:40	1.2	0.88	101				
5	5	KITTCHE	N	12	1510	21:70	26:90	5:20	1.2	1.1	143				
5	5	KITTCHE	N	12	1511	21:00	26:90	5:30	1.2	1.4	255				
5	5	KITTCHE	N	12	1512	21:00	26:90	5:30	1.2	1.4	603				
RUN: 3															
9	9	GAGIN		5	1427	19:60	25:00	5:40	1.2	1.19	1070				
9	9	GAGIN		5	1430	21:50	25:10	3:20	1.2	0.49	50				
9	9	GAGIN		5	1432	23:30	25:00	1:40	1.2	0.37	70				
9	9	GAGIN		5	1435	23:30	25:00	1:40	1.2	0.19	10				
RUN: 4															
9	9	GAGIN		5	1440	19:40	24:70	5:30	1.2	1.17	890				
9	9	GAGIN		5	1442	22:50	25:10	2:20	1.2	0.52	20				
9	9	GAGIN		5	1445	22:50	25:10	2:20	1.2	0.52	20				
RUN: 5															
9	9	GAGIN		5	1451	20:20	25:20	5:00	1.2	1.01	640				
9	9	GAGIN		5	1454	22:10	25:00	2:50	1.2	0.45	80				
9	9	GAGIN		5	1457	22:30	25:30	2:00	1.2	0.15	20				
9	9	GAGIN		5	1459	23:30	25:30	2:00	1.2	0.15	20				
RUN: 6															
9	9	GAGIN		5	1504	19:70	24:80	5:10	1.2	1.04	640				
9	9	GAGIN		5	1507	21:40	25:00	3:20	1.2	0.52	20				
9	9	GAGIN		5	1509	22:40	25:20	2:40	1.2	0.3	40				
9	9	GAGIN		5	1512	23:50	25:40	1:50	1.2	0.15	50				
RUN: 7															
9	9	GAGIN		5	1527	20:00	25:30	5:30	1.2	1.12	500				
9	9	GAGIN		5	1530	21:00	25:30	4:30	1.2	0.53	60				
9	9	GAGIN		5	1531	22:00	25:30	3:30	1.2	0.53	50				
RUN: 8															
9	9	GAGIN		5	1553	20:20	25:00	4:80	1.2	0.94	540				
9	9	GAGIN		5	1555	21:50	25:20	3:20	1.2	0.49	50				
9	9	GAGIN		5	1600	22:50	25:20	2:40	1.2	0.49	50				
RUN: 9															
10	10	LALA		13	1430	22:07	25:46	3:39	0.4	0.25	55			64	
10	10	LALA		13	1430	22:07	25:46	3:39	0.4	0.25	55				
10	10	LALA		13	1430	22:07	25:46	3:39	0.4	0.25	55				

EXPERIMENT #: 05

PROCESSING DATE: 81/07/29

NUCLEI MEASUREMENTS

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	COLD	HOT (C)	PLATE TEMP	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLFI COUNT	ORS N	EST K	REMARKS
10	10	LALA		13	1430		22.61	25.60		2.99	.4	.35	80.8	83.9		
10	10	LALA		13	1430		22.97	25.57		3.56	.4	.5	198	200.5		
10	10	LALA		13	1430		21.15	25.49		4.35	.4	.5	454	477.5		
10	10	LALA		13	1430		21.35	25.45		5.00	.4	.5	728	735		
10	10	LALA		13	1430		20.30	25.30		6.08	.4	1.1	882	884.5		
10	10	LALA		13	1430		19.17	25.25		7.08	.4	1.1	887	907		
10	10	LALA		13	1430	1443	18.08	25.07		6.99	.4	1.2	898	902.5		
RUN: 2																
10	10	LALA		13	1453		22.82	26.36		5.4	.4	.25	54	61.1		
10	10	LALA		13	1453		22.22	26.22		7.0	.4	.25	68.8	80.7		
10	10	LALA		13	1453		22.22	26.22		7.0	.4	.25	76.2	86.2		
10	10	LALA		13	1453		22.22	26.22		7.0	.4	.25	186.2	186.2		
10	10	LALA		13	1453		22.22	26.22		7.0	.4	.25	533	533		
10	10	LALA		13	1453		22.22	26.22		7.0	.4	.25	676.0	688		
10	10	LALA		13	1453		22.22	26.22		7.0	.4	.25	830.9	833		
10	10	LALA		13	1453		18.39	22.40		7.01	.4	1.1	843	864.5		
RUN: 3																
10	10	LALA		13	1553		24.07	26.26		5.5	.4	.25	78.8	66.65		
10	10	LALA		13	1553		23.22	26.22		5.5	.4	.25	107.0	100.45		
10	10	LALA		13	1553		23.22	26.22		5.5	.4	.25	182.2	193.5		
10	10	LALA		13	1553		23.22	26.22		5.5	.4	.25	265.5	265.5		
10	10	LALA		13	1553		23.22	26.22		5.5	.4	.25	428.0	497.2		
10	10	LALA		13	1553		23.22	26.22		5.5	.4	.25	527.0	745.4		
10	10	LALA		13	1553		23.22	26.22		5.5	.4	.25	762.0	877		
10	10	LALA		13	1553	1558	18.74	22.87		7.03	.4	1.1	881.2	877		
10	10	LALA		13	1553		18.74	22.87		7.03	.4	1.1	882.0	877		
RUN: 1																
13	13	AYERS		2	1429		25.52	28.00		4.8	.4	.25	0			35 MM FILM RECORD
13	13	AYERS		2	1431		23.79	27.90		4.37	.4	.25	150			ALSO TAKEN
13	13	AYERS		2	1435		22.05	27.60		5.15	.4	1.25	1200			
RUN: 2																
13	13	AYERS		2	1440		25.03	27.50		7	.4	.25	15			35 MM FILM RECORD
13	13	AYERS		2	1442		23.87	27.40		5.5	.4	.25	270			ALSO TAKEN
13	13	AYERS		2	1447		23.22	27.30		4.5	.4	.25	600			
13	13	AYERS		2	1450		22.22	27.20		4.5	.4	1.25	1095			
RUN: 1																
14	14	HUDSON		0	1610						.019	.21		0		
14	14	HUDSON		0	1610						.019	.13		0		
14	14	HUDSON		0	1610						.019	.02		0		

EXPERIMENT #: 05

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/07/29

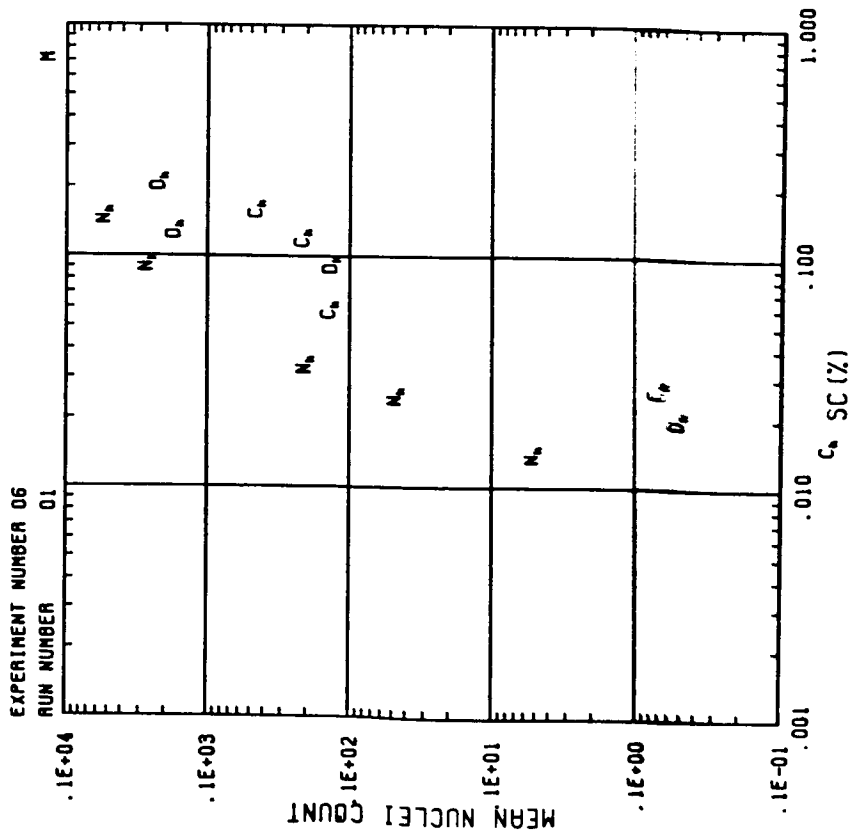
MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	SAMPLE END	COLD	HOT (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLFI COUNT	OPS - N	EST K	REMARKS
RUN: 1															
17	17	HOJCI	CHOWSKI	27	1433	1433	0	0	0	0	0	0	0	0	
17	17	HOJCI	CHOWSKI	27	1433	1433	0	0	0	0	0	0	0	0	
17	17	HOJCI	CHOWSKI	27	1433	1433	0	0	0	0	0	0	0	0	
17	17	HOJCI	CHOWSKI	27	1433	1433	0	0	0	0	0	0	0	0	
17	17	HOJCI	CHOWSKI	27	1433	1433	0	0	0	0	0	0	0	0	
RUN: 2															
17	17	HOJCI	CHOWSKI	27	1506	1506	0	0	0	0	0	0	0	0	
17	17	HOJCI	CHOWSKI	27	1506	1506	0	0	0	0	0	0	0	0	
17	17	HOJCI	CHOWSKI	27	1506	1506	0	0	0	0	0	0	0	0	
RUN: 1															
18	18	HUDSON		0	1427	1427	0	0	0	0	0	0	0	0	
18	18	HUDSON		0	1457	1457	0	0	0	0	0	0	0	0	
18	18	HUDSON		0	1514	1514	0	0	0	0	0	0	0	0	
18	18	HUDSON		0	1529	1529	0	0	0	0	0	0	0	0	
18	18	HUDSON		0	1545	1545	0	0	0	0	0	0	0	0	
18	18	HUDSON		0	1557	1557	0	0	0	0	0	0	0	0	
RUN: 1															
21	21	TRUEBL	OOD	0	1429	1429	0	0	0	0	0	0	0	0	
21	21	TRUEBL	OOD	0	1436	1436	0	0	0	0	0	0	0	0	
21	21	TRUEBL	OOD	0	1440	1440	0	0	0	0	0	0	0	0	
21	21	TRUEBL	OOD	0	1443	1443	0	0	0	0	0	0	0	0	
21	21	TRUEBL	OOD	0	1445	1445	0	0	0	0	0	0	0	0	
21	21	TRUEBL	OOD	0	1449	1449	0	0	0	0	0	0	0	0	
21	21	TRUEBL	OOD	0	1452	1452	0	0	0	0	0	0	0	0	
21	21	TRUEBL	OOD	0	1453	1453	0	0	0	0	0	0	0	0	
21	21	TRUEBL	OOD	0	1454	1454	0	0	0	0	0	0	0	0	
21	21	TRUEBL	OOD	0	1457	1457	0	0	0	0	0	0	0	0	
RUN: 1															
25	25	BORYS		3	1446	1446	0	0	0	0	0	0	0	0	
25	25	BORYS		3	1500	1500	0	0	0	0	0	0	0	0	
25	25	BORYS		3	1515	1515	0	0	0	0	0	0	0	0	
25	25	BORYS		3	1530	1530	0	0	0	0	0	0	0	0	
25	25	BORYS		3	1549	1549	0	0	0	0	0	0	0	0	
RUN: 1															
26	26	HINDMAN		6	1536	1536	0	0	0	0	0	0	0	0	
26	26	HINDMAN		6	1536	1536	0	0	0	0	0	0	0	0	
26	26	HINDMAN		6	1536	1536	0	0	0	0	0	0	0	0	

VOL SAMP. # 10**5
PROBLEM WITH LEAK

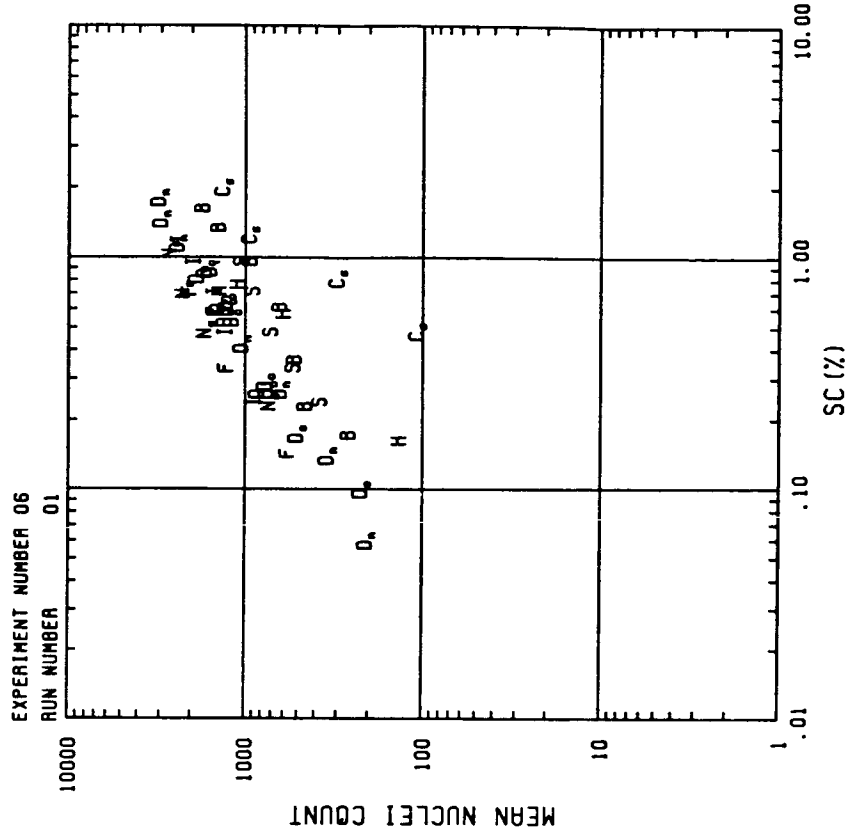
0** 0.5 UM
0** 0.4 UM
0** 1.0 UM
0** 3.0 UM

EXPERIMENT #: 05
 MACH NO. 27
 DUCT NO. 27
 NAME ONTAKA
 OBSERVER NO. 19
 SAMPLE START 1430
 SAMPLE END 1430
 COLD
 PLATE TEMP
 HOT (C)
 DELTA
 VOLUME SAMPLED (L) 3.5
 SUPER SAT (%)
 NUCLEI COUNT 677
 ORS N
 EST K
 REMARKS 118% R.H. NOMINAL
 PROCESSING DATE: 81/07/29

INSTRUMENT COMPARISON



INSTRUMENT COMPARISON



PURPOSE

INSTRUMENT COMPARISON WITH AMBIENT AEROSOL

DESCRIPTION OF EXPERIMENT

DATE 08 OCT 1980 1618 1800
TIME START END

NUCLEI TYPE
AMBIENT

GENERATION METHOD

NA

SIZE DISTRIBUTION SHAPING

NA

REMARKS

I-75(F), ID=54(F)
WIND CALM, W.D. VARIABLE

WEATHER SYNOPSIS

NA

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS			OUTSIDE CONDITIONS				
AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG WS (M/S)	AVG WD (DEG)
25.0	21.0	852	NA	NA	NA	0.0	90

EXPERIMENT #: 06

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/07/29

MACH NO.	DUCT NO.	OBSERVER	NO.	SAMPLE START	SAMPLE TIME END	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLFI COUNT	ORS N	EST K	REMARKS
						COLD	HOT (C)							
RUN: 1														
S	S	KITCHEN	1	1629	1630	26.00	26.00	3.50	2.50	.28	475			
S	S	KITCHEN	1	1630	1631	26.00	26.00	3.90	3.90	.38	522			
S	S	KITCHEN	1	1631	1632	26.00	26.00	4.80	4.80	.64	716			
S	S	KITCHEN	1	1632	1633	26.00	26.00	5.60	5.60	1.47	1056			
S	S	KITCHEN	1	1633	1634	26.00	26.00	6.50	6.50	1.47	1180			
S	S	KITCHEN	1	1634	1635	26.00	26.00	4.80	4.80	1.47	1144			
S	S	KITCHEN	1	1635	1636	26.00	26.00	3.90	3.90	.64	625		1.00	
RUN: 2														
S	S	KITCHEN	1	1637	1638	26.00	26.00	2.20	2.20	.18	273			
S	S	KITCHEN	1	1638	1639	26.00	26.00	3.90	3.90	.64	477			
S	S	KITCHEN	1	1639	1640	26.00	26.00	4.80	4.80	1.70	691			
S	S	KITCHEN	1	1640	1641	26.00	26.00	5.50	5.50	1.70	1053			
S	S	KITCHEN	1	1641	1642	26.00	26.00	4.80	4.80	1.47	1096			
S	S	KITCHEN	1	1642	1643	26.00	26.00	3.90	3.90	.64	571			
S	S	KITCHEN	1	1643	1644	26.00	26.00	3.10	3.10	.38	178		1.00	
RUN: 3														
S	S	GAGIN	5	1634	1635	25.20	25.20	4.70	4.70	.8	150			
S	S	GAGIN	5	1635	1636	25.40	25.40	3.90	3.90	.6	630			
S	S	GAGIN	5	1636	1637	25.60	25.60	1.80	1.80	.17	140			
RUN: 4														
S	S	GAGIN	5	1657	1658	25.20	25.20	5.10	5.10	1.05	1260			
S	S	GAGIN	5	1658	1659	25.40	25.40	3.70	3.70	.55	300			
S	S	GAGIN	5	1659	1660	25.60	25.60	2.20	2.20	.19	200			
RUN: 5														
S	S	GAGIN	5	1712	1713	25.20	25.20	5.00	5.00	1.01	1340			
S	S	GAGIN	5	1713	1714	25.40	25.40	2.10	2.10	.19	330			
RUN: 6														
S	S	LALA	13	1637	1638	26.00	26.00	2.56	2.56	.25	372	393	.73	
S	S	LALA	13	1638	1639	26.00	26.00	3.02	3.02	.35	545	554.5	.73	
S	S	LALA	13	1639	1640	26.00	26.00	3.59	3.59	.55	697	749	.73	
S	S	LALA	13	1640	1641	26.00	26.00	4.68	4.68	.75	803	939.3	.73	
S	S	LALA	13	1641	1642	26.00	26.00	5.46	5.46	.75	975	1099	.73	
S	S	LALA	13	1642	1643	26.00	26.00	5.04	5.04	.1	1090		.73	N=1154 S 0.73
RUN: 7														
S	S	LALA	13	1643	1644	26.00	26.00	5.6	5.6	.25	324	374	.73	
S	S	LALA	13	1644	1645	26.00	26.00	3.2	3.2	.35	456	563.5	.73	
S	S	LALA	13	1645	1646	26.00	26.00	3.54	3.54	.35	574	790.2	.73	
S	S	LALA	13	1646	1647	26.00	26.00	3.0	3.0	.35	495		.73	
S	S	LALA	13	1647	1648	26.00	26.00	4.0	4.0	.75	785		.73	
S	S	LALA	13	1648	1649	26.00	26.00	3.58	3.58	.75	972		.73	

EXPERIMENT # : 06										NUCLEI MEASUREMENTS										PROCESSING DATE: 81/07/29	
MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	SAMPLE END	COLD	HOT	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	ORBS N	EST K	REMARKS						
10	10	LALA		13	1645	1651	22.18	26.57	4.39		.75	1022.9	997.6								
10	10	LALA		13	1645	1651	21.44	26.49	5.05		.75	1095.3	1088.5		N=1202 S 0.77						
10	10	LALA		13	1645	1651	21.37	26.41	5.04		.75	1082.7									
RUN: 1																					
11	11	HOPPEL		7	1632	1644					.15	583.5	583.5		ISOTHERMAL HAZE CHAM						
11	11	HOPPEL		7	1632	1644					.09	290	290								
11	11	HOPPEL		7	1632	1644					.034	22.3	22.3								
11	11	HOPPEL		7	1632	1644					.023	5.07	5.07								
11	11	HOPPEL		7	1632	1644					.014	.55	.55								
RUN: 2																					
11	11	HOPPEL		7	1650	1707					.15	542.7	542.7		ISOTHERMAL HAZE CHAM						
11	11	HOPPEL		7	1650	1707					.09	220	220								
11	11	HOPPEL		7	1650	1707					.034	19.77	19.77								
11	11	HOPPEL		7	1650	1707					.025	2.55	2.55								
11	11	HOPPEL		7	1650	1707					.014	.37	.37								
RUN: 1																					
13	13	AYERS		2	1630	1631	25.71	27.80	2.09		.25	930	930		ALL SC VALUES ARE APPROXIMATE ONLY						
13	13	AYERS		2	1632	1633	24.74	27.70	3.56		.75	1350	1350		35 MM PHOTO TAKEN						
13	13	AYERS		2	1633	1634	23.20	27.60	4.30		.75	1520	1520								
13	13	AYERS		2	1639	1640	21.25	27.60	5.95		1.25	2503	2503								
RUN: 2																					
13	13	AYERS		2	1640	1641	21.86	27.40	5.54		1.25	2513	2513		35 MM PHOTO RECORDS TAKEN						
13	13	AYERS		2	1644	1645	22.48	27.20	4.98		.75	1350	1350								
13	13	AYERS		2	1649	1650	22.69	27.20	4.34		.75	1110	1110								
13	13	AYERS		2	1652	1653	24.70	27.20	2.50		.25	930	930		WARM PLATE DRIED OUT TOWARDS END OF RUN - LAST 3 POINTS QUESTIONARL						
RUN: 3																					
13	13	AYERS		2	1653	1654	24.70	27.20	2.50		.25	930	930		35 MM PHOTO TAKEN						
13	13	AYERS		2	1656	1657	23.05	27.20	4.29		.75	1350	1350								
13	13	AYERS		2	1659	1659	22.13	27.10	4.97		1.25	2250	2250								
13	13	AYERS		2	1700	1701	22.13	27.10	5.54		1.25	2130	2130								
13	13	AYERS		2	1703	1704	21.56	27.10	5.54		1.25	2130	2130								
13	13	AYERS		2	1705	1706	21.56	27.10	5.54		1.25	2130	2130								
RUN: 4																					
13	13	AYERS		2	1717	1718	24.37	26.80	2.43		.25	720	720								
13	13	AYERS		2	1718	1719	24.37	26.80	2.43		.25	720	720								
13	13	AYERS		2	1720	1721	23.43	26.80	3.37		.75	1250	1250								
13	13	AYERS		2	1722	1723	22.08	26.90	4.82		1.25	2273	2273								
13	13	AYERS		2	1725	1726	21.35	26.90	5.55		1.25	2400	2400								
RUN: 1																					
14	14	HUDSON		9	1630	1730					.21	3261	3261		PLATE TEMP APPROX IN ABS VALUE OK IN DELT						
14	14	HUDSON		9	1630	1730	18.30	25.00	6.70	.029	.21	3175	3175								
14	14	HUDSON		9	1630	1730	19.00	25.00	6.00	.029	.033	185	185								
14	14	HUDSON		9	1630	1730	19.70	25.00	5.30	.029	.033	14.4	14.4								
14	14	HUDSON		9	1630	1730	19.70	25.00	5.30	.029	.033	.056	.056								
RUN: 1																					
15	15	KEYSER		11	1640	1641	18.30	25.00	6.70	1.8	1.8	3261	3261								
15	15	KEYSER		11	1644	1645	19.00	25.00	6.00	1.6	1.6	3175	3175								
15	15	KEYSER		11	1648	1649	19.70	25.00	5.30	1.15	1.15	2547	2547								

EXPERIMENT #: 06

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/07/29

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	COLD	HOT	PLATE TEMP (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (g)	NUCLFI COUNT	DRS	EST	REMARKS
15	15	KEYSER		11	1652	1657	21.40	25.00	25.00	4.60	.003	.84	1968			
15	15	KEYSER		11	1706	1711	21.70	25.00	25.00	3.90	.003	.63	1495			
15	15	KEYSER		11	1705	1710	22.25	25.00	25.00	2.20	.003	.427	1103			
15	15	KEYSER		11	1708	1709	23.40	25.00	25.00	1.90	.003	.14	520			
15	15	KEYSER		11	1712	1713	23.80	25.00	25.00	1.20	.003	.06	219			
RUN: 1																
16	16	HUDSON		9	1621	1626	21.40	26.10	26.10	4.70	.003	.9	1663			
16	16	HUDSON		9	1621	1626	21.70	25.40	25.40	3.70	.003	.55	1379			
16	16	HUDSON		9	1621	1626	22.25	24.85	24.85	2.60	.003	.27	897			
RUN: 2																
16	16	HUDSON		9	1637	1641	21.40	26.10	26.10	4.70	.0024	.9	1453			
16	16	HUDSON		9	1637	1641	21.70	25.40	25.40	3.70	.0024	.55	1366			
16	16	HUDSON		9	1637	1641	22.25	24.85	24.85	2.60	.0024	.27	882			
RUN: 3																
16	16	HUDSON		9	1649	1651	21.40	26.10	26.10	4.70	.0018	.9	1875			
16	16	HUDSON		9	1649	1651	21.70	25.40	25.40	3.70	.0018	.55	1394			
16	16	HUDSON		9	1649	1651	22.25	24.85	24.85	2.60	.0018	.27	889			
RUN: 4																
16	16	HUDSON		9	1659	1702	21.40	26.10	26.10	4.70	.0018	.9	1851			
16	16	HUDSON		9	1659	1702	21.70	25.40	25.40	3.70	.0018	.55	1386			
16	16	HUDSON		9	1659	1702	22.25	24.85	24.85	2.60	.0018	.27	886			
RUN: 5																
16	16	HUDSON		9	1708	1710	21.40	26.10	26.10	4.70	.0018	.9	1837			
16	16	HUDSON		9	1708	1710	21.70	25.40	25.40	3.70	.0018	.55	1364			
16	16	HUDSON		9	1708	1710	22.25	24.85	24.85	2.60	.0018	.27	876			
RUN: 6																
16	16	HUDSON		9	1713	1716	21.40	26.10	26.10	4.70	.0018	.9	1851			
16	16	HUDSON		9	1713	1716	21.70	25.40	25.40	3.70	.0018	.55	1376			
16	16	HUDSON		9	1713	1716	22.25	24.85	24.85	2.60	.0018	.27	892			
RUN: 7																
16	16	HUDSON		9	1720	1723	21.40	26.10	26.10	4.70	.0018	.9	1809			
16	16	HUDSON		9	1720	1723	21.70	25.40	25.40	3.70	.0018	.55	1363			
16	16	HUDSON		9	1720	1723	22.25	24.85	24.85	2.60	.0018	.27	872			
RUN: 8																
16	16	HUDSON		9	1730	1732	21.40	26.10	26.10	4.70	.0018	.9	1754			
16	16	HUDSON		9	1730	1732	21.70	25.40	25.40	3.70	.0018	.55	1316			
16	16	HUDSON		9	1730	1732	22.25	24.85	24.85	2.60	.0018	.27	876			
RUN: 1																
17	17	WOJTECHOWSKI		27	1639	1639	21.40	26.10	26.10	4.70	.0018	.9	1789.6			
17	17	WOJTECHOWSKI		27	1639	1639	21.70	25.40	25.40	3.70	.0018	.55	1388.6			
17	17	WOJTECHOWSKI		27	1639	1639	22.25	24.85	24.85	2.60	.0018	.27	880.6			
RUN: 1																
18	18	HUDSON		9	1637	1629	21.40	26.10	26.10	4.70	.03	.64	1357			
18	18	HUDSON		9	1637	1629	21.70	25.40	25.40	3.70	.03	.64	1767			
18	18	HUDSON		9	1637	1629	22.25	24.85	24.85	2.60	.03	.64	1333			
18	18	HUDSON		9	1659	1702	22.60	26.30	26.30	3.70	.024	.55	1246			

EXPERIMENT #: 05

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/07/29

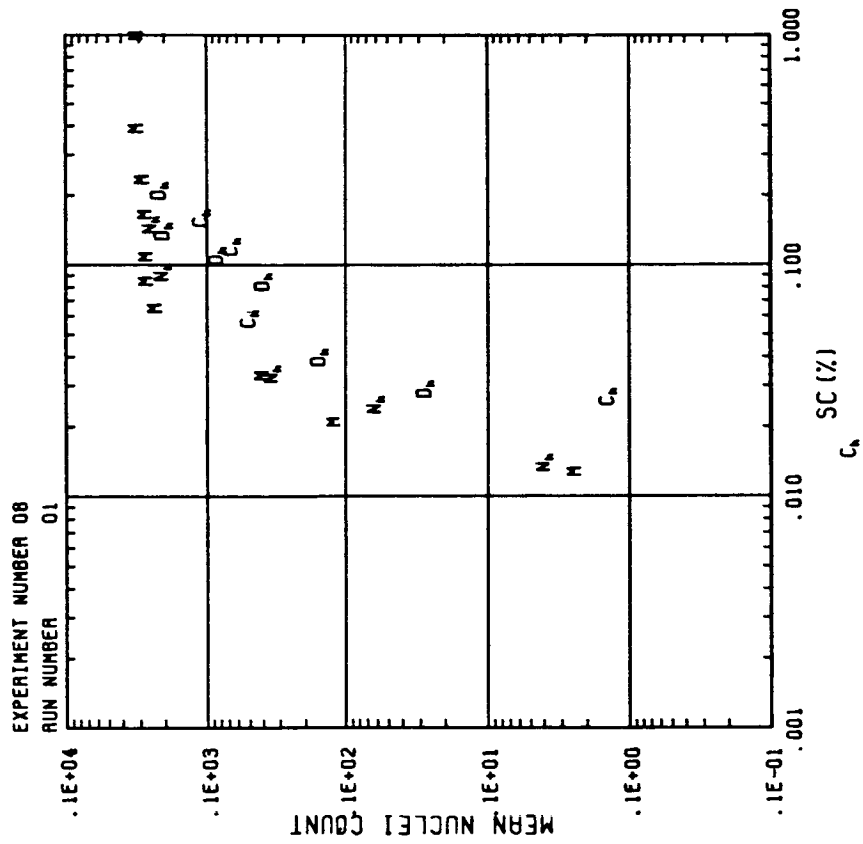
MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	SAMPLE END	COLD	HOT	PLATE TEMP	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLFI COUNT	ORIS - N	EST K	REMARKS
18	18	HUDSON		9	1708	1710	23.10	25.80		2.70	.018	.29	612			
18	18	HUDSON		9	1713	1716	23.10	25.70		2.60	.018	.27	568			
18	18	HUDSON		9	1720	1723	23.35	25.45		2.10	.018	.175	533			
18	18	HUDSON		9	1730	1732	23.60	25.20		1.60	.018	.1	233			
RUN: 1																
21	21	ALOFS		1	1625	1623	20.70	25.00		4.30		.75	1684			
21	21	ALOFS		1	1630	1633	20.70	25.00		4.30		.75	1397			
21	21	ALOFS		1	1635	1637	20.70	25.00		4.30		.75	1353			
21	21	ALOFS		1	1639	1642	20.70	25.00		4.30		.75	1391			
21	21	ALOFS		1	1643	1645	20.70	25.00		4.30		.75	1238			
21	21	ALOFS		1	1647	1649	20.70	25.00		4.30		.75	1547			
RUN: 2																
21	21	ALOFS		1	1701	1702	25.00	25.00		0.00		.068	118			
21	21	ALOFS		1	1701	1702	25.00	25.00		0.00		.089	246			
21	21	ALOFS		1	1701	1702	25.00	25.00		0.00		.114	349			
21	21	ALOFS		1	1701	1702	25.00	25.00		0.00		.173	695			
RUN: 3																
21	21	ALOFS		1	1704	1707	25.00	25.00		0.00		.088	114			
21	21	ALOFS		1	1704	1707	25.00	25.00		0.00		.089	239			
21	21	ALOFS		1	1704	1707	25.00	25.00		0.00		.173	655			
RUN: 4																
21	21	ALOFS		1	1708	1712	25.00	25.00		0.00		.068	119			
21	21	ALOFS		1	1708	1712	25.00	25.00		0.00		.089	254			
21	21	ALOFS		1	1708	1712	25.00	25.00		0.00		.114	358			
21	21	ALOFS		1	1708	1712	25.00	25.00		0.00		.173	720			
RUN: 5																
21	21	ALOFS		1	1713	1714	25.00	25.00		0.00		.068	117			
21	21	ALOFS		1	1713	1714	25.00	25.00		0.00		.089	244			
21	21	ALOFS		1	1713	1714	25.00	25.00		0.00		.173	676			
RUN: 6																
21	21	ALOFS		1	1716	1717	25.00	25.00		0.00		.068	119			
21	21	ALOFS		1	1716	1717	25.00	25.00		0.00		.089	243			
21	21	ALOFS		1	1716	1717	25.00	25.00		0.00		.173	693			
RUN: 1																
24	24	SERPOLAY		25	1625	1635	23.20	25.20		2.00	.77	.15	600			
24	24	SERPOLAY		25	1655	1645	23.70	25.70		3.00	.77	.35	1330			
24	24	SERPOLAY		25	1700	1700	24.60	26.60		4.00	.77	.61	1550			
24	24	SERPOLAY		25	1710	1715	25.20	26.60		4.40	.77	.75	2150			
RUN: 1																
25	25	BORYS		3	1633	1635	15.10	21.90		5.80	2.12	1.25	1415		.91	VOL. SAMP # 10**5
25	25	BORYS		3	1642	1645	16.50	22.90		4.40	3.18	.82	990			
25	25	BORYS		3	1652	1655	17.60	23.00		2.40	4.24	.48	318			
25	25	BORYS		3	1713	1720	19.70	24.00		2.30	8.48	.48	113			
RUN: 1																
26	26	HINDMAN		6	1628	1717	23.00	23.00		0.00	1.2	.16	49			E=1.5, D>= 0.5 UM

EXPERIMENT #: 06
 PROCESSING DATE: 81/07/29

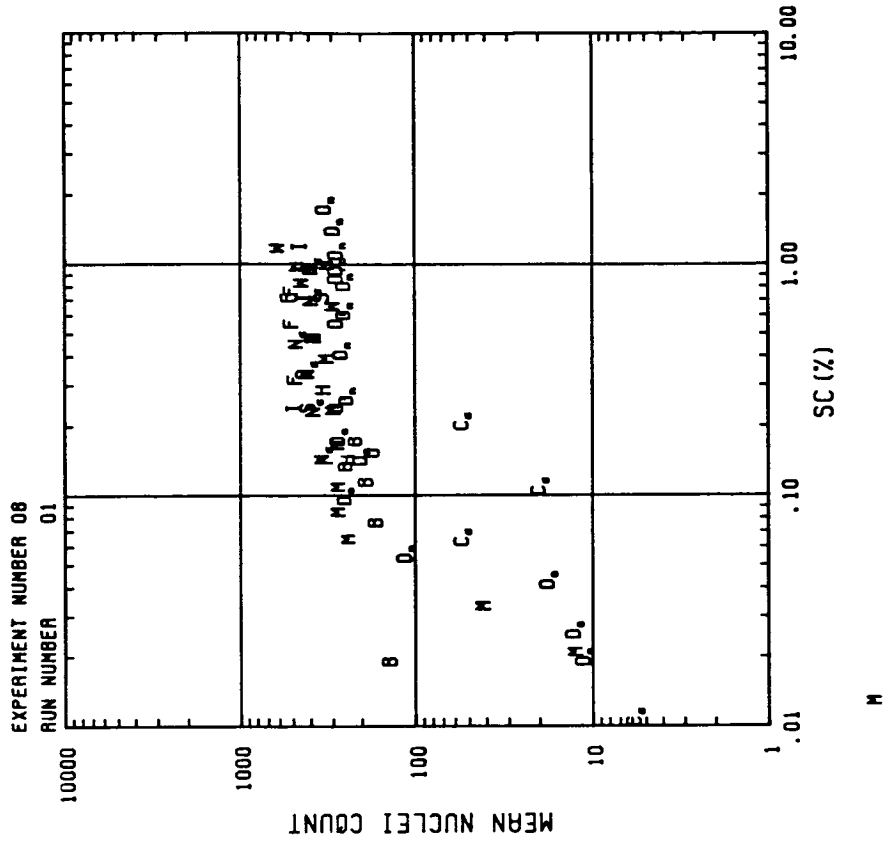
NUCLEI MEASUREMENTS

MACH NO.	DUCT NO.	OBSERVER NAME	OBSERVER NO.	SAMPLE START	SAMPLE TIME END	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLFI COUNT	ORBS N	EST R	REMARKS
						COLD	HOT (C)							
26	26	HINDMAN	6	1628	1717	23.00	23.00	0.00	1.2	.12	23	23	---	E=0.80, D>= 0.7 UM
26	26	HINDMAN	6	1628	1717	23.00	23.00	0.00	1.2	.059	15	15	---	E= 0.57, D>=1.4 UM
26	26	HINDMAN	6	1628	1717	23.00	23.00	0.00	1.2	.027	.075	.075	---	E= 0.12, D>= 3.0 UM
26	26	HINDMAN	6	1628	1717	23.00	23.00	0.00	1.2	.016	.0049	.0049	---	E= 0.0020, D>= 5.0UM
RUN: 1														
27	27	CHTAK	19	1620	1620				.35		2.89			102%
27	27	CHTAK	19	1620	1620				.35		2.35			103%
27	27	CHTAK	19	1620	1620				.35		3.35			115%

INSTRUMENT COMPARISON



INSTRUMENT COMPARISON



EXPERIMENT # 08

PURPOSE

HAZE CHAMBER TEST WITH LARGE PARTICLES

DESCRIPTION OF EXPERIMENT

DATE 09 OCT, 1980 TIME START 0955 END 1152

NUCLEI TYPE

NACL, MONODISPERSE, D=0.18 UM

GENERATION METHOD

DRI ATOMIZER, 40 PSI, 0.025 % SOLUTION.

SIZE DISTRIBUTION SHAPING

EQ. 8550V, MONODISPERSE FLOW 6 L/MIN. DILUTION FLOW 210 L/MIN.

REMARKS

SLIGHT DRIFT OF PARTICLE CONC. TO LOWER VALUES.

WEATHER SYNOPSIS

NONE

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS			OUTSIDE CONDITIONS				
AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG WS (M/S)	AVG WD (DEG)
26.0	22.0	854	NA	NA	NA	NA	NA

PROCESSING DATE: 81/07/29

NUCLEI MEASUREMENTS

EXPERIMENT # : 08		OBSERVER		SAMPLE TIME		PLATE TEMP		VOLUME SAMPLED		SUPER SAI		NUCLEI COUNT		OPS		EST		REMARKS	
MACH NO.	DUCT NO.	NAME	NO.	START	END	COLD	HOT (C)	DELTA	(L)	(%)			N	K					
2	1	ROGERS, DAVE	22	1050	1051	22.40	24.40	2.00	.16				179						
5	1	KITCHEN	12	0959	1004	26.90	27.60	1.70	.02		148								
5	1	KITCHEN	12	1004	1005	26.60	27.70	1.10	.14		208								
5	1	KITCHEN	12	1005	1006	25.40	27.70	2.30	.18		225								
5	1	KITCHEN	12	1006	1007	25.70	27.70	2.00	.14		240								
5	1	KITCHEN	12	1007	1008	25.90	27.70	1.80	.12		146								
5	1	KITCHEN	12	1008	1009	26.30	27.90	1.60	.08		172								
5	2	KITCHEN	12	1038	1039	23.50	28.20	4.70	.9		369								
5	2	KITCHEN	12	1039	1040	22.40	28.30	3.90	.6		347								
5	2	KITCHEN	12	1040	1041	22.40	28.30	3.20	.6		326								
5	2	KITCHEN	12	1041	1042	25.10	28.30	3.20	.3		251								
5	2	KITCHEN	12	1042	1043	25.10	28.30	3.20	.3		271								
5	2	KITCHEN	12	1043	1044	23.50	28.30	3.70	.5		404								
5	2	KITCHEN	12	1044	1045	23.60	28.30	3.70	.5		204								
5	2	KITCHEN	12	1045	1046	24.00	28.30	4.30	.6		331								
5	2	KITCHEN	12	1046	1047	24.00	28.30	4.30	.6		331								
9	1	GAGIN	5	1012	1015	21.80	26.90	5.10	1.02		490								
9	1	GAGIN	5	1015	1017	24.10	26.70	3.60	.3		400								
9	1	GAGIN	5	1017	1020	25.20	26.60	1.40	.15		350								
9	2	GAGIN	5	1024	1026	21.70	26.70	5.00	1.02		460								
9	2	GAGIN	5	1026	1029	23.30	26.90	3.60	.47		450								
9	2	GAGIN	5	1029	1032	24.00	27.90	3.90	.17		310								
9	3	GAGIN	5	1038	1039	21.60	26.70	5.10	1.06		410								
9	3	GAGIN	5	1039	1040	23.20	27.00	3.80	.37		200								
9	3	GAGIN	5	1040	1043	25.10	27.20	2.10	.17		220								
9	4	GAGIN	5	1052	1054	21.90	26.60	4.70	.89		330								
9	4	GAGIN	5	1054	1058	25.10	27.00	3.90	.53		300								
9	4	GAGIN	5	1058	1059	25.10	26.90	1.80	.13		240								
10	1	LALA	13	1006	1007	22.20	25.08	2.88	.25		418							423.5	
10	1	LALA	13	1007	1008	22.20	25.08	2.88	.25		425							445.5	
10	1	LALA	13	1008	1009	22.20	25.08	2.88	.25		425							400.5	
10	1	LALA	13	1009	1010	22.20	25.08	2.88	.25		425							355	
10	1	LALA	13	1010	1011	22.20	25.08	2.88	.25		425							414	
10	1	LALA	13	1011	1012	22.20	25.08	2.88	.25		425								
10	1	LALA	13	1012	1013	22.20	25.08	2.88	.25		425								
10	1	LALA	13	1013	1014	22.20	25.08	2.88	.25		425								
10	1	LALA	13	1014	1015	22.20	25.08	2.88	.25		425								

NORMAL OPERATION ONLY ABOVE 0.15%

EXPERIMENT #: 08

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/07/29

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	SAMPLE END	COLD	HOT (C)	PLATE TEMP	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	ORS N	EST K	REMARKS
RUN: 2																
10	10	LALA		13	1028	1028	20	25	23	53	7.9	25	418	434		
10	10	LALA		13	1023	1023	22	25	21	53	7.9	35	457			
10	10	LALA		13	1023	1023	22	25	19	58	8.8	35	429	6		
10	10	LALA		13	1023	1023	22	25	17	56	9.5	39	392	8		
10	10	LALA		13	1026	1026	22	25	13	53	7.5	36	353	3		
10	10	LALA		13	1027	1027	20	25	10	37	7.5	37	346	5		
10	10	LALA		13	1028	1028	20	25	10	37	7.5	37	353	3		
10	10	LALA		13	1028	1028	19	25	08	39	8.1	40	373	7		
RUN: 3																
10	10	LALA		13	1045	1045	22	26	26	77	3	3	39	5		
10	10	LALA		13	1047	1047	22	26	29	77	3	3	37	5		
10	10	LALA		13	1048	1048	22	26	29	77	3	3	37	5		
10	10	LALA		13	1049	1049	22	26	29	77	3	3	37	5		
10	10	LALA		13	1050	1050	22	26	29	77	3	3	37	5		
10	10	LALA		13	1051	1051	22	26	29	77	3	3	37	5		
10	10	LALA		13	1052	1052	20	26	25	75	3	3	36	4		
RUN: 1																
11	11	HOPPEL		7	1103	1110	22	27	27	15	0.94	15	270	270		
11	11	HOPPEL		7	1103	1110	22	27	27	15	0.94	15	222	222		
11	11	HOPPEL		7	1103	1110	22	27	27	15	0.94	15	365	365		
RUN: 2																
11	11	FITZGERALD		28	1004	1019	24	27	27	9	0.15	9	308	308		
11	11	FITZGERALD		28	1004	1019	22	27	27	9	0.15	9	250	250		
11	11	FITZGERALD		28	1004	1019	22	27	27	9	0.15	9	313	313		
11	11	FITZGERALD		28	1004	1019	22	27	27	9	0.15	9	8	8		
11	11	FITZGERALD		28	1004	1019	22	27	27	9	0.15	9	8	8		
RUN: 1																
13	13	AYERS		2	1028	1019	24	27	27	9	0.25	9	518	518		
13	13	AYERS		2	1022	1023	22	27	27	9	0.25	9	450	450		
13	13	AYERS		2	1027	1028	22	27	27	9	0.25	9	473	473		
RUN: 2																
13	13	AYERS		2	1036	1035	24	26	80	5	0.25	5	233	267.5		
13	13	AYERS		2	1037	1038	24	26	80	5	0.25	5	285	285		
13	13	AYERS		2	1039	1040	22	26	80	5	0.25	5	245	260		
13	13	AYERS		2	1041	1042	22	26	80	5	0.25	5	215	215		
13	13	AYERS		2	1043	1044	22	26	80	5	0.25	5	240	382.5		
13	13	AYERS		2	1045	1046	22	26	80	5	0.25	5	375	375		
13	13	AYERS		2	1047	1048	22	26	70	5	0.25	5	375	337.5		
13	13	AYERS		2	1052	1053	21	26	70	5	0.25	5	300	300		
RUN: 1																
14	14	HUDSON		9	1123	1133	0.065	21	21	0.65	0.65	21	299	299		E48:5: D=0:39
14	14	HUDSON		9	1123	1133	0.065	21	21	0.65	0.65	21	223	223		E48:5: D=0:39
14	14	HUDSON		9	1123	1133	0.065	21	21	0.65	0.65	21	42.8	42.8		E48:5: D=0:39

PHOTO RECORDS TAKEN

E48:5: D=0:39
 F48:5: D=0:39
 F48:5: D=0:39
 F48:5: D=0:39

EXPERIMENT #: 08

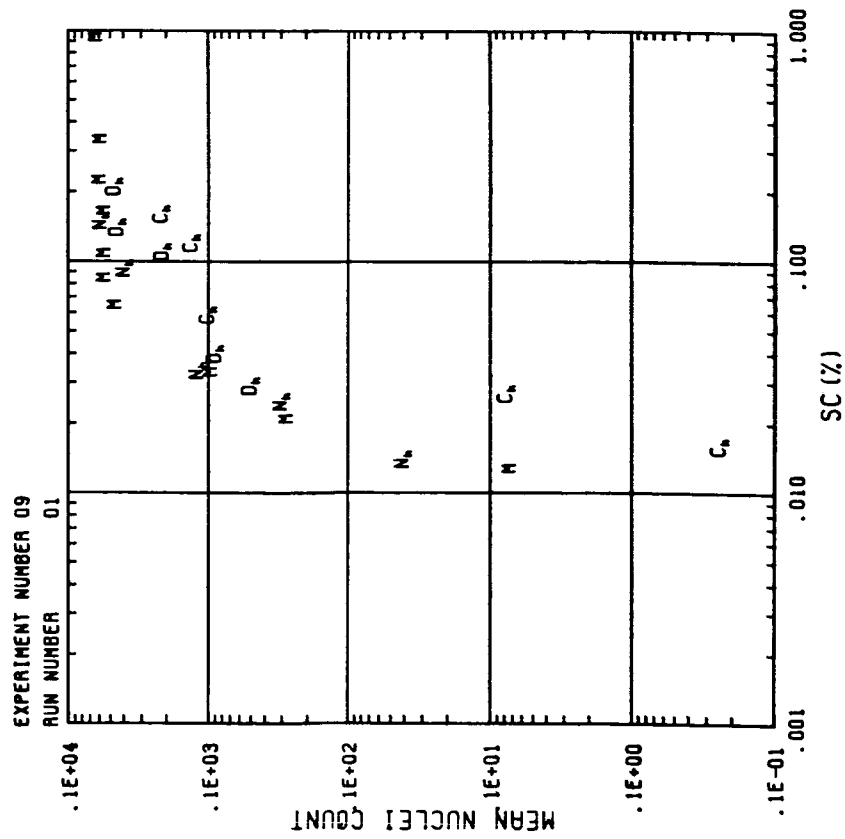
NUCLEI MEASUREMENTS

PROCESSING DATE: 81/07/29

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	COLD	HOT (C)	PLATE TEMP	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OPS N	EST K	REMARKS
14	14	HUDSON		9	1123	1133					.0065	.029	320			F=48.5; D=4.0
14	14	HUDSON		9	1142	1145					.0026	.14	214			F=48.5; D=0.38
14	14	HUDSON		9	1142	1145					.0026	.11	291			F=48.5; D=0.37
14	14	HUDSON		9	1142	1145					.0026	.029	159			F=48.5; D=2.7
14	14	HUDSON		9	1142	1145					.0026	.029	266			F=48.5; D=4.0
RUN: 2																
14	14	HUDSON		9	1152	1155					.0026	.21	170			F=28; D=0.38
14	14	HUDSON		9	1152	1155					.0026	.11	174			F=28; D=0.37
14	14	HUDSON		9	1152	1155					.0026	.029	156			F=28; D=2.7
14	14	HUDSON		9	1152	1155					.0026	.029	309			F=28; D=4.0
RUN: 1																
15	15	ROGERS	FRED	23	1026	1037	18.30	25.05	25.05	6.70		1.8	347			BACKGROUND CNT
15	15	ROGERS	FRED	23	1037	1038	19.12	24.44	24.44	5.32		1.15	307			ABOUT 10 PER CC
15	15	ROGERS	FRED	23	1042	1043	19.87	23.81	23.81	4.04		.84	268			PLATE TEMPS APPROX
15	15	ROGERS	FRED	23	1045	1046	20.23	23.50	23.50	3.94		.83	277			IN ABSOLUTE. OK IN
15	15	ROGERS	FRED	23	1052	1054	20.63	23.19	23.19	2.56		.27	256			DELTA
15	15	ROGERS	FRED	23	1053	1054	21.00	22.89	22.89	1.80		.27	256			READING ROYCO CH. 2
15	15	ROGERS	FRED	23	1057	1058	21.45	22.25	22.25	1.15		.002	119			LEAST BACKGROUND
15	15	ROGERS	FRED	23	1130	1131	22.10	22.25	22.25				140			LAST PT. NEAR ISOTH
RUN: 1																
16	16	HUDSON		9	1005	1008	0	25.10	25.10	4.70	.0018	.9	311			
16	16	HUDSON		9	1005	1008	4.30	25.80	25.80	3.20	.0018	.58	319			
16	16	HUDSON		9	1019	1022	21.30	26.50	26.50	4.20	.0018	.29	298			
16	16	HUDSON		9	1019	1022	21.70	25.80	25.80	3.20	.0018	.58	240			
16	16	HUDSON		9	1035	1037	21.70	26.10	26.10	4.20	.0018	.29	240			
16	16	HUDSON		9	1035	1037	22.30	25.80	25.80	3.20	.0018	.58	240			
RUN: 2																
16	16	HUDSON		9	1057	1058	0	26.10	26.10	4.70	.0024	.9	266			
16	16	HUDSON		9	1057	1058	11.70	25.80	25.80	3.20	.0024	.29	240			
16	16	HUDSON		9	1114	1116	21.70	26.50	26.50	4.20	.0018	.29	240			
16	16	HUDSON		9	1114	1116	21.70	25.80	25.80	3.20	.0018	.58	240			
16	16	HUDSON		9	1122	1124	21.70	26.10	26.10	4.20	.0018	.29	240			
16	16	HUDSON		9	1122	1124	22.30	25.80	25.80	3.20	.0018	.58	240			
RUN: 1																
17	17	WOJTECHOWSKI		7	1018	1019	0	27.00	27.00	2.50		.15	359	26		
17	17	WOJTECHOWSKI		7	1021	1022	4.50	27.00	27.00	3.50		.25	407	29		
17	17	WOJTECHOWSKI		7	1027	1028	22.70	27.00	27.00	3.50		.25	407	29		
17	17	WOJTECHOWSKI		7	1030	1031	22.70	27.00	27.00	4.50		.25	407	29		
RUN: 1																
18	18	HUDSON		9	1059	1062	0	26.90	26.90	4.90	.0018	.98	288			
18	18	HUDSON		9	1059	1062	4.00	25.50	25.50	2.10	.0018	.18	298			
18	18	HUDSON		9	1057	1058	23.65	25.50	25.50	1.60	.0024	.20	118			

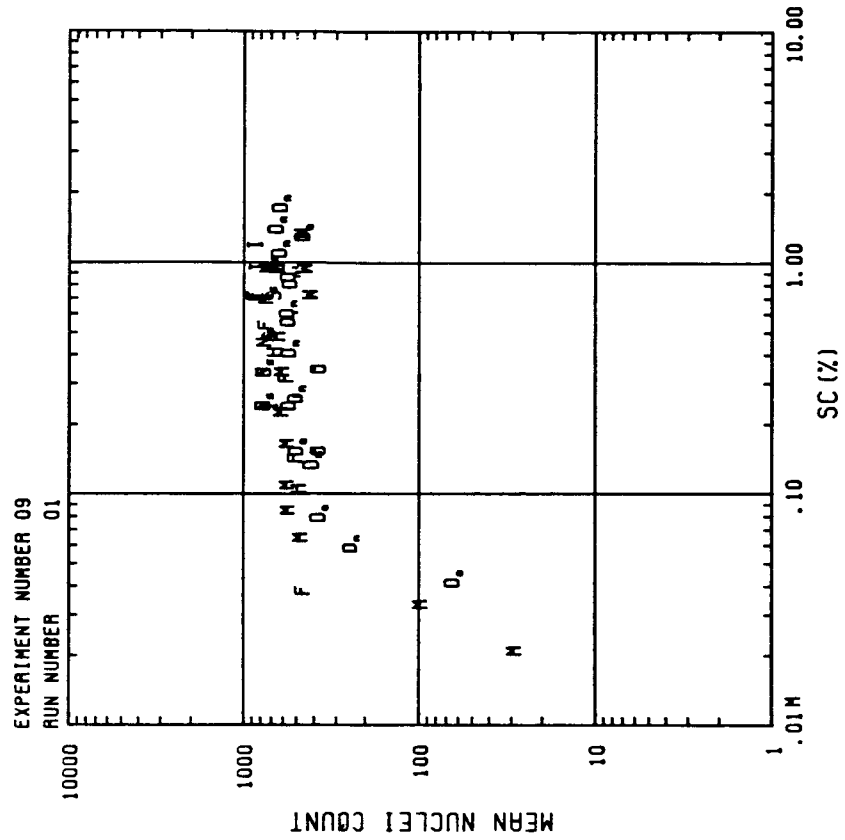
EXPERIMENT #:		NUCLEI MEASUREMENTS										PROCESSING DATE:			
#:												81/07/29			
MACH NO.	DUCT NO.	OBSERVER	NO.	SAMPLE START	TIME END	PLATE TEMP		COLD	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLFI COUNT	ORS	EST K	REMARKS
						HOT (C)	HOT (C)								
18	18	HUDSON	9	1114	1116	24.10	24.00	1.85	0.018	0.043	18.2				
RUN: 1															
20	20	DOMONKOS	4	1120	1130	25.05	25.00	0.69	0.018	0.043	310				
20	20	DOMONKOS	4	1120	1130	25.05	25.00	0.87	0.018	0.043	460				
20	20	DOMONKOS	4	1120	1130	25.05	25.00	1.23	0.018	0.043	630				
RUN: 1															
21	21	TRUEBLOOD	26	1007	1009	19.89	19.89	5.11	0.018	0.043	333				
21	21	TRUEBLOOD	26	1014	1015	21.80	21.80	3.20	0.018	0.043	337				
21	21	TRUEBLOOD	26	1026	1027	22.50	22.50	2.48	0.018	0.043	305				
21	21	TRUEBLOOD	26	1035	1037	22.50	22.50	0.00	0.018	0.043	291				
21	21	TRUEBLOOD	26	1035	1037	22.50	22.50	0.00	0.018	0.043	289				
21	21	TRUEBLOOD	26	1035	1037	22.50	22.50	0.00	0.018	0.043	289				
21	21	TRUEBLOOD	26	1042	1044	22.50	22.50	0.00	0.018	0.043	43				
21	21	TRUEBLOOD	26	1042	1044	22.50	22.50	0.00	0.018	0.043	13				
21	21	TRUEBLOOD	26	1042	1044	22.50	22.50	0.00	0.018	0.043	26				
RUN: 1															
24	24	SERPOLAY	23	1009	1015	26.40	26.40	3.00	0.018	0.043	340				
24	24	SERPOLAY	23	1015	1020	29.40	29.40	3.00	0.018	0.043	500				
24	24	SERPOLAY	23	1030	1035	30.70	30.70	4.00	0.018	0.043	510				
24	24	SERPOLAY	23	1045	1050	31.50	31.50	4.60	0.018	0.043	530				
RUN: 1															
25	25	BORYS	3	1010	1020	23.90	23.90	0.70	0.018	0.043	6				VOL. SAMP. X10**5
25	25	BORYS	3	1024	1030	23.70	23.70	0.00	0.018	0.043	0				CHAMBER PROBLEM.
25	25	BORYS	3	1050	1055	23.50	23.50	0.00	0.018	0.043	0				
25	25	BORYS	3	1103	1115	22.50	22.50	1.70	0.018	0.043	57				
25	25	BORYS	3	1120	1125	22.50	22.50	2.70	0.018	0.043	57				
RUN: 1															
26	26	HINDMAN	6	1007	1125	24.00	24.00	0.00	0.018	0.043	117				E=3.0, D>0.5 UM
26	26	HINDMAN	6	1007	1125	24.00	24.00	0.00	0.018	0.043	117				E=3.0, D>0.7 UM
26	26	HINDMAN	6	1007	1125	24.00	24.00	0.00	0.018	0.043	53				E=2.0, D>1.4 UM
26	26	HINDMAN	6	1007	1125	24.00	24.00	0.00	0.018	0.043	53				E=0.016, D>3.0 UM
26	26	HINDMAN	6	1007	1125	24.00	24.00	0.00	0.018	0.043	0.029				E=0.0011, D>5.0 UM
RUN: 1															
27	27	OHTAKE	19	1024	1024						1.426				109% R.H. NOMINAL

INSTRUMENT COMPARISON



HAZE CHAMBER RESULTS

INSTRUMENT COMPARISON



SDC/CFD RESULTS

PURPOSE

HAZE CHAMBER TESTS (LARGE PARTICLES)

DESCRIPTION OF EXPERIMENT

DATE	TIME START	TIME END
09 OCT.1960	1152	1319

NUCLEI TYPE

NACL, MONODISPERSE, D=0.18 UM

GENERATION METHOD

DRI ATOMIZER, 40 PSI, 0.5% SOLUTION.

SIZE DISTRIBUTION SHAPING

EG, 8550V, MONODISPERSE FLOW 6 L/MIN. DILUTION FLOW 210 L/MIN.

REMARKS

SLIGHT DRIFT IN CONC TO LOWER VALUES DUE TO CHANGE IN MONODISPERSE FLOW

WEATHER SYNOPSIS

NA

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS		OUTSIDE CONDITIONS					
AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG WS (M/S)	AVG WD (DEG)
26.0	22.0	854	NA	NA	NA	NA	NA

PROCESSING DATE: 81/07/29

NUCLEI MEASUREMENTS

EXPERIMENT #: 09		OBSERVER		NUCLEI MEASUREMENTS										PROCESSING DATE: 81/07/29	
MACH NO.	DUCT NO.	NAME	NO.	SAMPLE START	TIME END	COLD	HOT	PLATE TEMP	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OPS N	EST K	REMARKS
RUN: 1				1231	1232	23.30	25.30	3.00			.36		392	0.00	
2	2	ROGERS, DAVE	22	1231	1232	23.30	25.30	3.00			.36		392	0.00	
RUN: 2				1206		21.80	26.90	5.10		1.2	1.01	720			
9	9	GAGIN	5	1206		23.30	26.90	3.60		1.2	.43	670			
9	9	GAGIN	5	1210		23.70	27.00	3.30		1.2	.41	690			
9	9	GAGIN	5	1213		25.20	27.10	1.90			.11	500			
RUN: 3				1218		22.00	26.80	4.80		1.2	.91	760			
9	9	GAGIN	5	1218		23.20	26.90	3.70		1.2	.53	660			
9	9	GAGIN	5	1223		24.10	26.70	2.60		1.2	.3	610			
9	9	GAGIN	5	1226		25.20	27.10	1.90			.11	620			
RUN: 3				1230		21.80	27.00	5.20		1.2	1.08	600			
9	9	GAGIN	5	1230		23.20	26.90	3.70		1.2	.53	660			
9	9	GAGIN	5	1235		23.50	27.20	3.70			.17	530			
9	9	GAGIN	5	1257		25.00	27.00	2.00				510			
RUN: 1				1207		23.70	26.90	3.20			.25	798.1	806.5		
10	10	LALA	13	1207		24.20	26.90	2.70			.25	810.7	800.9		
10	10	LALA	13	1208		24.70	26.90	2.00			.35	800.5	742.7		
10	10	LALA	13	1209		25.10	26.90	1.40			.25	733.8	698.6		
10	10	LALA	13	1210		25.10	26.90	0.00			.75	700.3	655.4		
10	10	LALA	13	1211		25.30	26.90	0.40			.75	690.9	655.4		
10	10	LALA	13	1212		19.70	26.90	4.40			.1	688.9	672.7		
10	10	LALA	13	1213		19.70	26.90	4.40			.1	688.9	672.7		
RUN: 2				1207		21.40	26.90	2.50			.2	812.5	806.5		
10	10	LALA	13	1207		21.96	26.90	2.50			.2	725.2	742.7		
10	10	LALA	13	1208		21.52	26.90	3.50			.4	759	742.7		
10	10	LALA	13	1209		21.52	26.90	3.50			.6	669.6	698.6		
10	10	LALA	13	1210		20.60	26.90	4.10			.8	710.3	698.6		
10	10	LALA	13	1211		20.00	26.90	4.40			.9	626.6	672.7		
10	10	LALA	13	1212		19.70	26.90	4.40			.1	626.6	672.7		
RUN: 3				1221		22.30	26.90	2.50			.25	822.6	852.6		
10	10	LALA	13	1221		22.36	26.90	2.50			.25	880.6	852.6		
10	10	LALA	13	1221		21.99	26.90	2.96			.35	762.8	750.2		
10	10	LALA	13	1221		21.99	26.90	2.96			.35	762.8	750.2		
10	10	LALA	13	1221		21.42	26.90	3.50			.5	670.1	695.8		
10	10	LALA	13	1221		20.60	26.90	3.50			.7	710.3	698.6		
10	10	LALA	13	1221		20.60	26.90	3.50			.7	710.3	698.6		
10	10	LALA	13	1221		19.90	26.90	4.40			.1	632.1	647.6		
10	10	LALA	13	1231		19.90	26.90	4.40			.1	632.1	647.6		
RUN: 1				1251	1307						.15	636	636		ISOTHERMAL HAZE CHAM
11	11	HOPPEL	7	1251	1307						.15	636	636		

PROCESSING DATE: 81/07/29

NUCLEI MEASUREMENTS

EXPERIMENT #: 09

MACH NO.	DUCT NO.	OBSERVER	NAME	NO.	SAMPLE START	TIME END	COLD	HOT (C)	PLATE TEMP	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLFI COUNT	ORS N	EST K	REMARKS
11	11	HOPPEL		7	1251	1307	26.5	26.5	0.0	3.5	0.04	430	430			
11	11	HOPPEL		7	1251	1307	26.5	26.5	0.0	3.5	0.04	128	128			
11	11	HOPPEL		7	1251	1307	26.5	26.5	0.0	3.5	0.04	33	33			
11	11	HOPPEL		7	1251	1307	26.5	26.5	0.0	3.5	0.04	4.4	4.4			
RUN: 1																
13	13	AYERS		2	206	222	26.5	26.5	0.0	5.0	0.04	595	595			POSS. ZERO RESET EBR
13	13	AYERS		2	206	222	26.5	26.5	0.0	5.0	0.04	740	740			
13	13	AYERS		2	206	222	26.5	26.5	0.0	5.0	0.04	840	840			
13	13	AYERS		2	206	222	26.5	26.5	0.0	5.0	0.04	900	900			
13	13	AYERS		2	206	222	26.5	26.5	0.0	5.0	0.04	780	780			
13	13	AYERS		2	206	222	26.5	26.5	0.0	5.0	0.04	840	840			
13	13	AYERS		2	206	222	26.5	26.5	0.0	5.0	0.04	720	720			
13	13	AYERS		2	206	222	26.5	26.5	0.0	5.0	0.04	825	825			
13	13	AYERS		2	206	222	26.5	26.5	0.0	5.0	0.04	877	877			
13	13	AYERS		2	206	222	26.5	26.5	0.0	5.0	0.04	960	960			
RUN: 1																
14	14	HUDSON		9	1211	1222	18.3	18.3	0.5	0.12	0.1	509	509			F=28. D=0.38
14	14	HUDSON		9	1211	1222	18.3	18.3	0.5	0.12	0.1	490	490			F=28. D=0.67
14	14	HUDSON		9	1211	1222	18.3	18.3	0.5	0.12	0.1	128	128			F=28. D=1.3
14	14	HUDSON		9	1211	1222	18.3	18.3	0.5	0.12	0.1	183	183			F=28. D=2.7
14	14	HUDSON		9	1211	1222	18.3	18.3	0.5	0.12	0.1	498	498			F=28. D=4.0
14	14	HUDSON		9	1211	1222	18.3	18.3	0.5	0.12	0.1	476	476			F=42.3. D=0.38
14	14	HUDSON		9	1211	1222	18.3	18.3	0.5	0.12	0.1	309	309			F=42.3. D=1.7
14	14	HUDSON		9	1211	1222	18.3	18.3	0.5	0.12	0.1	244	244			F=42.3. D=2.7
14	14	HUDSON		9	1211	1222	18.3	18.3	0.5	0.12	0.1	395	395			F=25. D=0.38
14	14	HUDSON		9	1211	1222	18.3	18.3	0.5	0.12	0.1	351	351			F=25. D=0.67
14	14	HUDSON		9	1211	1222	18.3	18.3	0.5	0.12	0.1	181	181			F=25. D=1.3
14	14	HUDSON		9	1211	1222	18.3	18.3	0.5	0.12	0.1	181	181			F=25. D=2.7
RUN: 2																
15	15	ROGERS		3	201	225	18.3	18.3	0.5	0.09	0.02	653	653			BACKGROUND ABOUT 10 PER CC.
15	15	ROGERS		3	201	225	18.3	18.3	0.5	0.09	0.02	659	659			TEMPS APPROX IN ABSOLUTE, OK IN DELT
15	15	ROGERS		3	201	225	18.3	18.3	0.5	0.09	0.02	574	574			READING ROYCO CH. 2 LESS BACKGROUND NEAR ISOTHERMAL ON LAST POINT
15	15	ROGERS		3	201	225	18.3	18.3	0.5	0.09	0.02	591	591			
15	15	ROGERS		3	201	225	18.3	18.3	0.5	0.09	0.02	431	431			
15	15	ROGERS		3	201	225	18.3	18.3	0.5	0.09	0.02	288	288			
15	15	ROGERS		3	201	225	18.3	18.3	0.5	0.09	0.02	250	250			
15	15	ROGERS		3	201	225	18.3	18.3	0.5	0.09	0.02	70	70			
15	15	ROGERS		3	201	225	18.3	18.3	0.5	0.09	0.02	50	50			
15	15	ROGERS		3	201	225	18.3	18.3	0.5	0.09	0.02	50	50			
15	15	ROGERS		3	201	225	18.3	18.3	0.5	0.09	0.02	50	50			
15	15	ROGERS		3	201	225	18.3	18.3	0.5	0.09	0.02	50	50			
15	15	ROGERS		3	201	225	18.3	18.3	0.5	0.09	0.02	50	50			
RUN: 1																
16	16	HUDSON		9	1203	1222	22.2	22.2	0.0	0.03	0.02	616	616			
16	16	HUDSON		9	1203	1222	22.2	22.2	0.0	0.03	0.02	616	616			
16	16	HUDSON		9	1203	1222	22.2	22.2	0.0	0.03	0.02	591	591			
16	16	HUDSON		9	1203	1222	22.2	22.2	0.0	0.03	0.02	591	591			
16	16	HUDSON		9	1203	1222	22.2	22.2	0.0	0.03	0.02	591	591			
16	16	HUDSON		9	1203	1222	22.2	22.2	0.0	0.03	0.02	591	591			
16	16	HUDSON		9	1203	1222	22.2	22.2	0.0	0.03	0.02	591	591			
16	16	HUDSON		9	1203	1222	22.2	22.2	0.0	0.03	0.02	591	591			
16	16	HUDSON		9	1203	1222	22.2	22.2	0.0	0.03	0.02	591	591			
16	16	HUDSON		9	1203	1222	22.2	22.2	0.0	0.03	0.02	591	591			

PROCESSING DATE: 81/07/29

NUCLEI MEASUREMENTS

EXPERIMENT #: 09

MACH NO.	DUCT NO.	OBSERVER	NO.	SAMPLE START	TIME END	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPEK SAT (%)	NUCLEI COUNT	ORS N	EST K	REMARKS
						COLD	HOT (C)							
16	16	HUDSON	9	1247	1250	21.70	25.50	3.80	.0018	.58	552			
***	INCORRECT EXPER.	NO.												
16	16	HUDSON	9	1247	1250	22.30	24.80	2.50	.0018	.25	551			
***	INCORRECT EXPER.	NO.												
RUN: 2														
16	16	HUDSON	9	1259	1302	21.40	26.10	4.70	.0024	.9	508			
***	INCORRECT EXPER.	NO.												
16	16	HUDSON	9	1259	1302	21.70	25.50	3.80	.0024	.58	507			
16	16	HUDSON	9	1259	1302	22.30	24.80	2.50	.0024	.25	507			
RUN: 1														
17	17	DOJICIECHOWSKI	27	1209	1207					.25	809			
17	17	DOJICIECHOWSKI	27	1209	1208					.35	809			
17	17	DOJICIECHOWSKI	27	1209	1210					.47	799.2			
17	17	DOJICIECHOWSKI	27	1210	1211					.72	779.4			
17	17	DOJICIECHOWSKI	27	1211	1212					.98	769.6			
RUN: 1														
18	18	HUDSON	9	1203	1208	24.00	25.05	1.05	.003	.043	67			
***	INCORRECT EXPER.	NO.												
18	18	HUDSON	9	1228	1230	23.75	25.20	1.45	.0018	.082	296			
18	18	HUDSON	9	1247	1250	23.50	25.50	2.00	.0018	.136	295			
18	18	HUDSON	9	1259	1302	21.65	27.40	5.75	.0024	1.35	483			
RUN: 1														
20	20	DONKOS	4	1245	1250					.76	436			
20	20	DONKOS	4	1245	1250					.98	463			
20	20	DONKOS	4	1245	1250					1.4	490			
RUN: 2														
20	20	DONKOS	4	1305	1310					.64	405			
20	20	DONKOS	4	1305	1310					.74	440			
20	20	DONKOS	4	1305	1310					1.04	554			
RUN: 1														
21	21	TRUEBLOOD	26	1207	1209	20.04	25.00	4.96		.984	673.5			
21	21	TRUEBLOOD	26	1210	1216	23.04	25.00	2.00		.335	64.8			
21	21	TRUEBLOOD	26	1228	1230	23.57	25.00	1.43		.775	59.2			
21	21	TRUEBLOOD	26	1228	1230	23.00	25.00	2.00		.114	59.0			
21	21	TRUEBLOOD	26	1228	1230	23.00	25.00	2.00		.089	59.2			
21	21	TRUEBLOOD	26	1232	1230	23.00	25.00	2.00		.035	102.6			
21	21	TRUEBLOOD	26	1237	1240	23.00	25.00	2.00		.022	29.4			
21	21	TRUEBLOOD	26	1237	1240	23.00	25.00	2.00		.0133	29.4			
RUN: 2														
21	21	TRUEBLOOD	26	1245	1247	25.00	25.00	0.00		.173	545.4			
21	21	TRUEBLOOD	26	1245	1247	25.00	25.00	0.00		.114	542.8			
21	21	TRUEBLOOD	26	1245	1247	25.00	25.00	0.00		.089	542.8			
21	21	TRUEBLOOD	26	1245	1247	25.00	25.00	0.00		.068	459.6			
RUN: 1														
24	24	SEAPOLAY	25	1205	1210	26.70	27.70	1.00	.77	.04	440			
24	24	SEAPOLAY	25	1233	1240	26.80	29.80	3.00	.77	.15	530			
24	24	SEAPOLAY	25	1247	1255	27.00	30.90	3.90	.77	.33	600			
24	24	SEAPOLAY	25	1305	1310	27.30	31.90	4.60	.77	.56	740			
RUN: 1														
26	26	HINDMAN	6	1213	1315	25.00	25.00	0.00	1.1	.16	233			

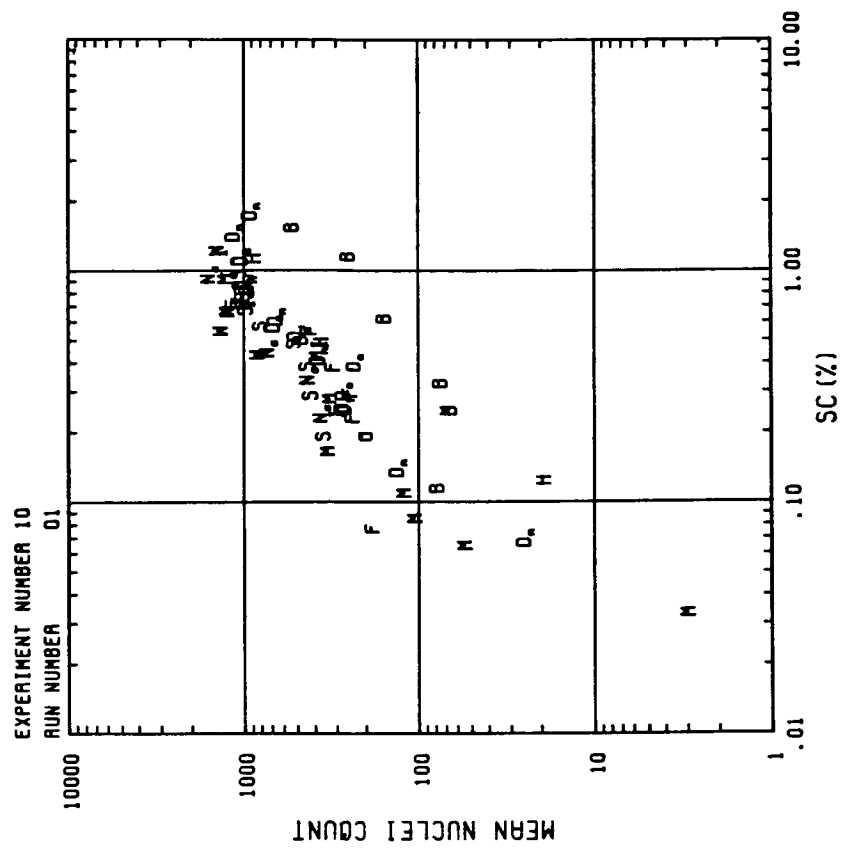
E=6.6 D>=0.5 UM

EXPERIMENT #: 09
 PROCESSING DATE: 81/07/29

NUCLEI MEASUREMENTS

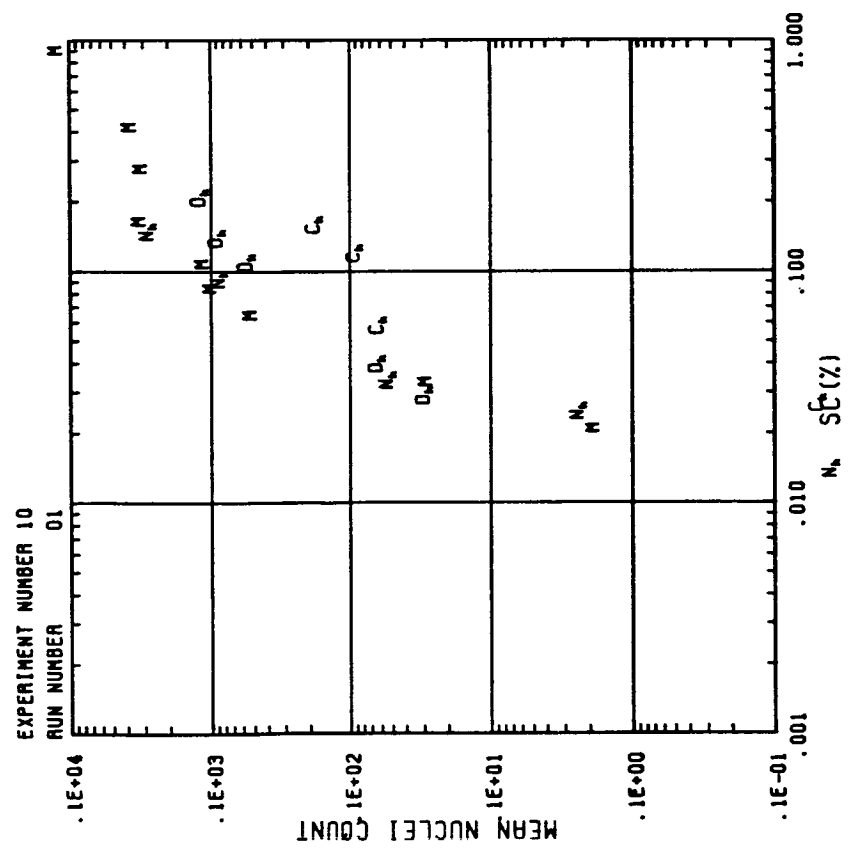
MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLFI COUNT	ORS N	EST K	REMARKS
							COLD	HOT (C)							
26	26	HINDMAN		6	1213	1315	25.00	25.00	0.00	1.1	.12	141	141		E=4.8; D>=0.7 UM
26	26	HINDMAN		6	1213	1315	25.00	25.00	0.00	1.1	.059	108	108		E=3.8; D>=1.4 UM
26	26	HINDMAN		6	1213	1315	25.00	25.00	0.00	1.1	.027	.83	.83		E=0.058; D>=3.0 UM
26	26	HINDMAN		6	1213	1315	25.00	25.00	0.00	1.1	.016	.026	.026		E=0.0068; D>=5.0 UM
RUN: 1															
27	27	ORTAKE		19	1205	1205				.1225		42.318			99% R.H.
27	27	ORTAKE		19	1205	1205				.1225		70.531			108% R.H. NOMINAL

INSTRUMENT COMPARISON



SDC/CFD RESULTS

INSTRUMENT COMPARISON



HAZE CHAMBER RESULTS

EXPERIMENT # 10

PURPOSE

BIMODAL NACL TO GET K-PLATEAU

DESCRIPTION OF EXPERIMENT

DATE	TIME START	END
09 OCT, 1980	1417	1620

NUCLEI TYPE

NACL, 0.1% SOLUTION FOR SMALL, 0.5% FOR LARGE SIZE.

GENERATION METHOD

ATOMIZERS: DRI, 40 PSI, NANOMIST 19PSI.

SIZE DISTRIBUTION SHAPING

EC1, 0.1UM DIA., EC2 0.03 UM DIA., EACH EC APPROX. 5.5 L/MIN, DIL AIR FLOW 410 L/MIN.

REMARKS

TOTAL N STEADY, BUT SMALL SIZE DRIFT UP, LARGE ONE DOWN.

WEATHER SYNOPSIS

NA

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS			OUTSIDE CONDITIONS			
AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG WD (DEG)
26.0	22.0	851	NA	NA	NA	NA

EXPERIMENT #: 10

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/07/29

MACH NO.	DUCT NO.	OBSERVER	NO.	SAMPLE START	SAMPLE END	COLD	HOT	PLATE TEMP	DELTA	VOLUME SAMPLED (L)	SUPER SAT (g)	NUCLEI COUNT	ORS N	EST K	REMARKS
RUN: 1															
2	2	POLITOVICH	20	1506	1507	23.10	25.40	2.30		.2			206		
RUN: 1															
5	5	KITCHEN	12	1503	1504	23.90	26.50	2.60		.26		68			
5	5	KITCHEN	12	1504	1505	23.90	26.50	2.60		1.2		325			
5	5	KITCHEN	12	1505	1506	23.90	26.50	2.60		1.2		554			
5	5	KITCHEN	12	1506	1507	23.90	26.50	2.60		1.2		250			
5	5	KITCHEN	12	1507	1508	23.90	26.50	2.60		.34		176			
5	5	KITCHEN	12	1508	1509	23.90	26.50	2.60		.12		91			
5	5	KITCHEN	12	1509	1510	23.90	26.50	2.60		.12		79			
5	5	KITCHEN	12	1510	1511	23.90	26.50	2.60		1.2		150			
5	5	KITCHEN	12	1511	1512	23.90	26.50	2.60		1.2		276			
RUN: 2															
5	5	KITCHEN	12	1513	1514	23.90	26.50	2.60		1.2		273			
5	5	KITCHEN	12	1514	1515	23.90	26.50	2.60		1.2		266			
5	5	KITCHEN	12	1515	1516	23.90	26.50	2.60		.65		90			
5	5	KITCHEN	12	1516	1517	23.90	26.50	2.60		.34		47			
5	5	KITCHEN	12	1517	1518	23.90	26.50	2.60		.12		47			
RUN: 1															
9	9	GAGIN	5	1506	1508	21.60	27.00	5.40		1.18		920			
9	9	GAGIN	5	1508	1510	21.60	27.00	5.40		1.18		380			
9	9	GAGIN	5	1510	1513	21.60	27.00	5.40		1.18		70			
RUN: 2															
9	9	GAGIN	5	1517	1520	23.30	27.30	4.00		.95		640			
9	9	GAGIN	5	1520	1522	23.30	27.30	4.00		.53		140			
9	9	GAGIN	5	1522	1525	23.30	27.30	4.00		.1		40			
RUN: 3															
9	9	GAGIN	5	1529	1532	23.40	27.30	3.90		.92		670			
9	9	GAGIN	5	1532	1537	23.40	27.30	3.90		.46		170			
9	9	GAGIN	5	1537	1538	23.40	27.30	3.90		.1		150			
RUN: 4															
9	9	GAGIN	5	1540	1542	23.30	27.10	4.80		.9		670			
9	9	GAGIN	5	1542	1544	23.30	27.10	4.80		.51		150			
9	9	GAGIN	5	1544	1548	23.30	27.10	4.80		.3		10			
RUN: 5															
9	9	GAGIN	5	1556	1558	23.60	27.80	4.20		1.06		670			
9	9	GAGIN	5	1558	1605	23.60	27.80	4.20		.62		320			
RUN: 1															
10	10	LJLA	13	1453	1454	23.16	25.4	2.26		.3		370			
10	10	LJLA	13	1454	1455	23.16	25.4	2.26		.3		43			
10	10	LJLA	13	1455	1457	23.16	25.4	2.26		.5		17			
10	10	LJLA	13	1457	1459	23.16	25.4	2.26		.3		84			

PROCESSING DATE: 81/07/29

NUCLEI MEASUREMENTS

EXPERIMENT #: 10

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	COLD	HOT	PLATE TEMP (C)	DELTA	VOLUME SAMPLED (L)	SUPER SA (%)	NUCLEI COUNT	OBS N	EST K	REMARKS	
10	10	LALA		13	1459		21.25	25.5	25.5	4.28		.7	124				
10	10	LALA		13	1459		20.67	25.4	25.4	4.73		.9	102				
10	10	LALA		13	1501		20.40	25.40	25.40	5.00		.1	114				
RUN: 2																	
10	10	LALA		13	1503		23.22	25.9	25.9	2.77		.2	397				
10	10	LALA		13	1505		22.46	25.66	25.66	3.20		.4	396				
10	10	LALA		13	1506		22.13	25.99	25.99	3.56		.5	424				
10	10	LALA		13	1507		22.80	25.70	25.70	3.90		.6	615				
10	10	LALA		13	1508		22.14	25.69	25.69	4.21		.7	818				
10	10	LALA		13	1509		21.88	25.67	25.67	4.49		.8	1022				
10	10	LALA		13	1510		20.89	25.65	25.65	4.76		.9	1099				
10	10	LALA		13	1511		20.63	25.64	25.64	5.01		.1	1028				
RUN: 3																	
10	10	LALA		13	1513		20.69	25.65	25.65	5.94		.1	128				
10	10	LALA		13	1514		20.24	25.54	25.54	5.45		.7	135				
10	10	LALA		13	1515		21.24	25.54	25.54	3.90		.6	290				
10	10	LALA		13	1517		21.68	25.54	25.54	3.59		.5	408				
10	10	LALA		13	1519		22.33	25.54	25.54	3.77		.4	446				
10	10	LALA		13	1520		22.37	25.54	25.54	2.21		.2	306				
RUN: 1																	
11	11	HOPPEL		7	1507	1524						.014	0042	0042			ISOTHERMAL MAGE CHAM
11	11	HOPPEL		7	1507	1524						.15	308	308			
11	11	HOPPEL		7	1507	1524						.094	95	95			
11	11	HOPPEL		7	1507	1524						.034	58	58			
11	11	HOPPEL		7	1507	1524						.025	26	26			
RUN: 1																	
13	13	AYERS		2	1457	1458	26.91	29.00	29.00	2.49		.25	308				
13	13	AYERS		2	1459	1500	25.76	29.00	29.00	2.64		.75	390				
13	13	AYERS		2	1500	1501	21.49	29.00	29.00	3.90		.75	290				
13	13	AYERS		2	1502	1505	23.49	29.00	29.00	5.01		1.25	1425				
RUN: 2																	
13	13	AYERS		2	1508	1509	23.19	28.80	28.80	5.61		1.25	1440				
13	13	AYERS		2	1512	1513	24.35	28.80	28.80	5.05		.75	1750				
13	13	AYERS		2	1514	1515	25.60	28.80	28.80	3.58		.35	447				
13	13	AYERS		2	1517	1520	25.62	28.80	28.80	2.48		.25	223				
13	13	AYERS		2	1519	1520	26.12	28.80	28.80	2.48		.25	223				
RUN: 3																	
13	13	AYERS		2	1526	1527	26.20	28.70	28.70	2.50		.25	255				
13	13	AYERS		2	1529	1530	24.33	28.70	28.70	3.57		.75	390				
13	13	AYERS		2	1531	1532	23.60	28.60	28.60	4.50		1.25	900				
13	13	AYERS		2	1555	1556	23.04	28.60	28.60	5.56		1.25	1145				
RUN: 1																	
14	14	HUDSON		9	1449	1455					.0039	.21	183				F=25.0 D=0.37
14	14	HUDSON		9	1449	1455					.0039	.11	92				F=25.0 D=1.07
14	14	HUDSON		9	1449	1455					.0039	.04	10				F=25.0 D=2.7

PROCESSING DATE: 81/07/79

NUCLEI MEASUREMENTS

EXPERIMENT #: 10

MACH NO.	DUCT NO.	OBSERVER	NAME	NO.	SAMPLE START	TIME END	COLD	HOT (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLFI COUNT	ORS N	EST K	REMARKS
14	14	HUDSON		9	1449	1455				.039	.029	5.53			F=39, D=4.0
14	14	HUDSON		9	1505	1519				.085	.14	143.2			F=39,
14	14	HUDSON		9	1505	1519				.085	.11	104.6			F=39,
14	14	HUDSON		9	1505	1519				.085	.04	51.2			F=39,
14	14	HUDSON		9	1505	1519				.085	.029	3.46			F=39,
RUN: 2															
14	14	HUDSON		9	1520	1549				.017	.21	12.9			F=39
14	14	HUDSON		9	1520	1549				.017	.14	98.2			F=39
14	14	HUDSON		9	1520	1549				.017	.11	38.5			F=39
14	14	HUDSON		9	1520	1549				.017	.029	3.6			F=39
14	14	HUDSON		9	1520	1549				.017	.15	4.2			F=39
14	14	HUDSON		9	1520	1549				.017	.11	40.5			F=39
14	14	HUDSON		9	1520	1549				.017	.11	2.6			F=39
14	14	HUDSON		9	1520	1549				.017	.029	2.6			F=39
RUN: 1															
15	15	ROGERS		2	1448	1450	1837	205	6.70		1.46	9.2			BACKGROUND NOW 10
15	15	ROGERS		2	1457	1459	1912	224	5.32		1.15	121			TO 16 PER CC
15	15	ROGERS		2	1505	1506	2052	223	4.94		1.17	111			READING ROYCD CH.
15	15	ROGERS		2	1509	1510	2052	223	3.27		.63	107			2 LESS 16 PER CC.
15	15	ROGERS		2	1517	1518	2052	223	2.59		.265	401			TEMPSS ARE APPROX.
15	15	ROGERS		2	1517	1518	2052	223	1.22		.17	126			IN ABSOLUTE, OK IN
15	15	ROGERS		2	1517	1518	2052	223	1.22		.17	126			DELTA
15	15	ROGERS		2	1517	1518	2052	223	1.22		.17	126			LAST PT. IS CH 3 CNT
RUN: 1															
16	16	HUDSON		9	1511	1514	2140	265	4.75	.001	.54	193			
16	16	HUDSON		9	1511	1514	2140	265	3.07	.001	.26	44			
16	16	HUDSON		9	1523	1524	2140	265	3.60	.001	.54	25			
16	16	HUDSON		9	1523	1524	2140	265	3.60	.001	.26	55			
16	16	HUDSON		9	1523	1524	2140	265	3.60	.001	.26	22			
16	16	HUDSON		9	1523	1524	2140	265	3.60	.001	.26	22			
16	16	HUDSON		9	1523	1524	2140	265	3.60	.001	.26	45			
RUN: 2															
16	16	HUDSON		9	1545	1548	2140	265	4.70	.001	.9	112			
16	16	HUDSON		9	1545	1549	2140	265	3.65	.001	.54	151			
16	16	HUDSON		9	1600	1602	2140	265	3.65	.001	.54	105			
16	16	HUDSON		9	1600	1602	2140	265	3.65	.001	.54	247			
16	16	HUDSON		9	1607	1610	2140	265	3.65	.001	.54	100			
16	16	HUDSON		9	1607	1610	2140	265	3.65	.001	.54	282			
RUN: 3															
16	16	HUDSON		9	1618	1619	2140	265	4.70	.001	.9	116			
16	16	HUDSON		9	1618	1619	2140	265	3.65	.001	.54	25			
16	16	HUDSON		9	1618	1619	2140	265	3.65	.001	.54	25			
RUN: 1															
17	17	MOJICE		2	1449	1457					.24	385			
17	17	MOJICE		2	1453	1455					.35	454			
17	17	MOJICE		2	1453	1455					.47	770			
17	17	MOJICE		2	1453	1455					.7	1307			

PROCESSING DATE: 81/07/79

NUCLEI MEASUREMENTS

EXPERIMENT #: 10

MACH NO.	DUCT NO.	OBSERVER	NO.	SAMPLE START	SAMPLE END	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
						COLD	HOT (C)							
17	17	WOJCIECHOWSKI	27	1453	1455					.96	1657	1657		
RUN: 1														
18	18	HUDSON	9	1511	1514	25.85	25.75	3.75	.0018	.2	287			
18	18	HUDSON	9	1529	1542	25.75	25.60	3.60	.0022	.95	1055			
18	18	HUDSON	9	1545	1548	25.60	25.30	5.30	.0018	1.75	1082			
18	18	HUDSON	9	1607	1610	25.60	25.30	4.30	.0018	.59	1047			
18	18	HUDSON	9	1618	1619	25.85	25.15	3.15	.0012	.4	1717			
RUN: 1														
20	20	DOMONKOS	4	1530	1540					.45	870			
20	20	DOMONKOS	4	1530	1540					.57	1290			
20	20	DOMONKOS	4	1530	1540					1.27	1410			
RUN: 1														
21	21	TRUEBLOOD	26	1443	1445	25.00	25.00	4.90	.0013	.96	1334			
21	21	TRUEBLOOD	26	1451	1452	25.00	25.00	3.33	.0013	.44	404			
21	21	TRUEBLOOD	26	1456	1458	25.00	25.00	2.70	.0013	.292	340			
21	21	TRUEBLOOD	26	1506	1508	25.00	25.00	0.00	.0013	.173	350			
21	21	TRUEBLOOD	26	1506	1508	25.00	25.00	0.00	.0013	.14	124			
21	21	TRUEBLOOD	26	1506	1508	25.00	25.00	0.00	.0013	.089	109			
21	21	TRUEBLOOD	26	1512	1514	25.00	25.00	0.00	.0013	.058	156			
21	21	TRUEBLOOD	26	1512	1518	25.00	25.00	0.00	.0013	.035	3			
21	21	TRUEBLOOD	26	1512	1518	25.00	25.00	0.00	.0013	.013	.0			
RUN: 2														
21	21	TRUEBLOOD	26	1534	1536	25.00	25.00	4.05	.0013	.66	888			
21	21	TRUEBLOOD	26	1544	1546	25.00	25.00	3.87	.0013	.59	601			
21	21	TRUEBLOOD	26	1549	1550	25.00	25.00	3.38	.0013	.457	401			
RUN: 3														
21	21	TRUEBLOOD	26	1601	1601	25.00	25.00	5.05	.0013	1.02	1428			
21	21	TRUEBLOOD	26	1605	1605	25.00	25.00	4.85	.0013	.97	1428			
21	21	TRUEBLOOD	26	1608	1608	25.00	25.00	4.32	.0013	.76	119			
21	21	TRUEBLOOD	26	1610	1611	25.00	25.00	4.25	.0013	.66	91			
21	21	TRUEBLOOD	26	1613	1614	25.00	25.00	3.90	.0013	.58	621			
21	21	TRUEBLOOD	26	1616	1617	25.00	25.00	3.60	.0013	.57	342			
21	21	TRUEBLOOD	26	1619	1620	25.00	25.00	3.40	.0013	.4	284			
RUN: 1														
24	24	SERPOLAY	25	1445	1500	26.90	1.30	1.30	.77	.08	190			
24	24	SERPOLAY	25	1510	1520	29.60	3.40	3.40	.77	.24	250			
24	24	SERPOLAY	25	1540	1552	31.30	4.60	4.60	.77	.4	330			
24	24	SERPOLAY	25	1600	1605	30.50	2.90	2.90	.77	.57	450			
24	24	SERPOLAY	25	1617	1622	27.00	0.00	0.00	.77	.31	260			
RUN: 1														
26	26	HINDMAN	6	1453	1609	27.00	0.00	0.00	1.4	.16	20			E=0.78, D>=0.5 UM
26	26	HINDMAN	6	1453	1609	27.00	0.00	0.00	1.4	.12	10			E=0.45, D>=0.7 UM
26	26	HINDMAN	6	1453	1609	27.00	0.00	0.00	1.4	.052	6.8			E=0.45, D>=1.4 UM
26	26	HINDMAN	6	1453	1609	27.00	0.00	0.00	1.4	.027	.0052			E=0.0022, D>=3 UM
26	26	HINDMAN	6	1453	1609	27.00	0.00	0.00	1.4	.016	0			0, D>=5 UM

EXPERIMENT #: 10
 MACH NO. 27
 DUCT NO. 27
 NAME OHTAKE
 OBSERVER
 NO. 19
 RUN: 1
 OHTAKE
 OHTAKE

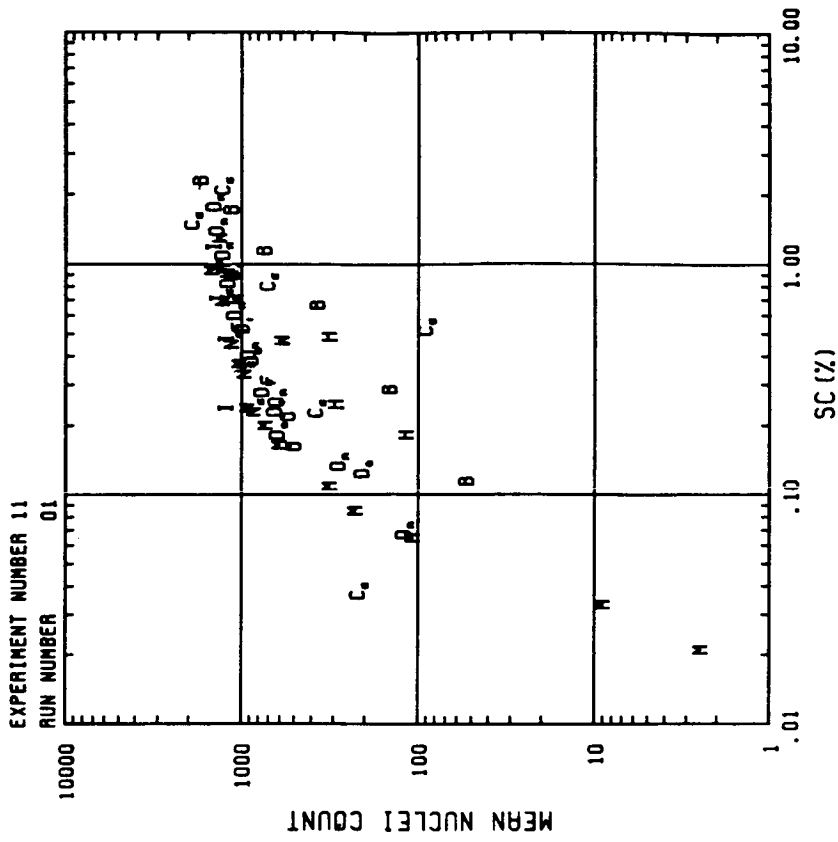
NUCLEI MEASUREMENTS

PROCESSING DATE: 81/07/29
 EST K
 REMARKS
 107% R.M. NOMINAL
 123% R.M. NOMINAL

PLATE TEMP	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OAS
COLD	HOT (C)	DELTA		N
	3.5		1.481	
	3.5		1.975	

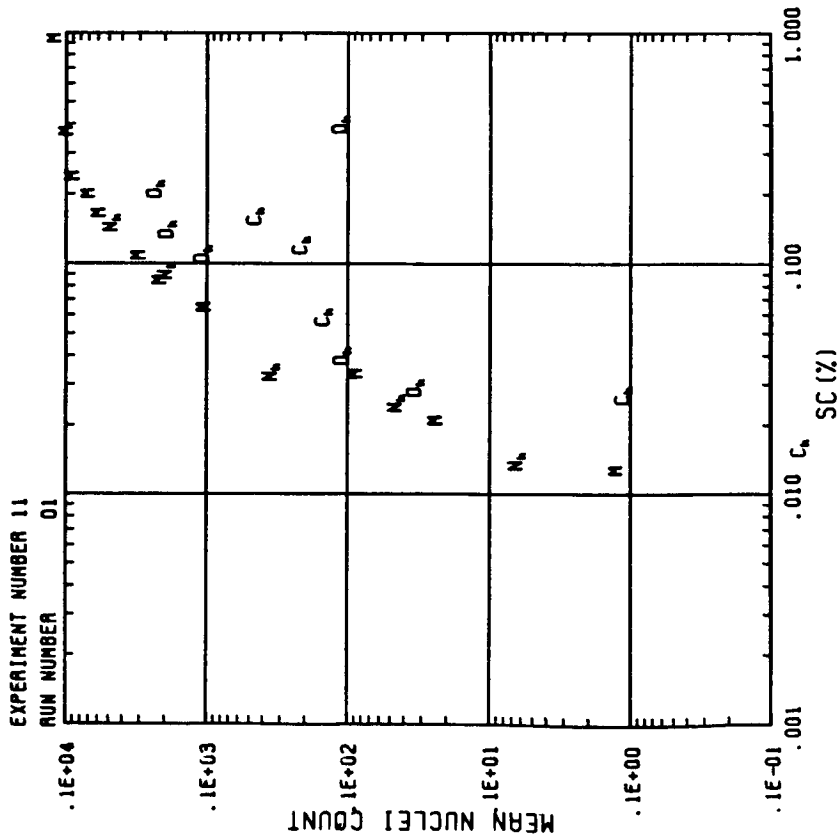
SAMPLE TIME
START
END
1444
1601

INSTRUMENT COMPARISON



M SDC/CFD RESULTS

INSTRUMENT COMPARISON



M HAZE CHAMBER RESULTS

EXPERIMENT # 11

PURPOSE

INSTRUMENT COMPARISON WITH AMBIENT AIR.

DESCRIPTION OF EXPERIMENT

DATE DATE TIME START END

09 OCT 1980 1636 1725

NUCLEI TYPE

AMBIENT

GENERATION METHOD

NA

SIZE DISTRIBUTION SHAPING

NA

REMARKS

FLOW RATE 700 L/MIN.

WEATHER SYNOPSIS

NONE

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS		OUTSIDE CONDITIONS					
AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG WS (M/S)	AVG WD (DEG)
29.0	21.0	851	NA	NA	851	1.0	120

EXPERIMENT #: 11
 PROCESSING DATE: 81/07/29

NUCLEI MEASUREMENTS

MACH NO.	DUCT NO.	OBSERVER	NAME	NO.	SAMPLE START	SAMPLE TIME END	COLD	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	ORS - N	EST K	REMARKS
								HOT (C)	TEMP (C)							
RUN: 1																
2	2	POLITOVICH		20	1702	1703	23.30	25.40	25.40	2.40	1.23	567				
RUN: 1																
5	5	KITTCHEM		12	1640	1640	22.00	27.40	5.40	1.2	935					
5	5	KITTCHEM		12	1641	1641	23.20	27.40	4.20	1.2	512					
5	5	KITTCHEM		12	1642	1642	24.50	27.40	2.90	1.2	94					
5	5	KITTCHEM		12	1643	1643	24.70	27.40	2.70	1.2	55					
5	5	KITTCHEM		12	1644	1644	24.40	27.40	3.00	1.2	207					
5	5	KITTCHEM		12	1645	1645	24.20	27.40	4.20	1.2	257					
5	5	KITTCHEM		12	1646	1646	23.00	27.40	4.40	1.2	1745					
5	5	KITTCHEM		12	1647	1647	24.70	27.40	2.70	1.2	1781					
5	5	KITTCHEM		12	1648	1648	24.70	27.40	2.70	1.2	186					
5	5	KITTCHEM		12	1649	1649	21.90	27.30	5.40	1.2	638					
RUN: 1																
9	9	GAGIN		5	1639	1639	22.90	28.20	5.30	1.2	1330					
9	9	GAGIN		5	1642	1642	24.00	27.60	3.60	1.2	330					
9	9	GAGIN		5	1644	1644	25.10	27.70	2.60	1.2	300					
9	9	GAGIN		5	1646	1646	25.80	28.10	2.30	1.2	120					
RUN: 2																
9	9	GAGIN		5	1649	1649	22.90	27.80	4.90	1.2	1350					
9	9	GAGIN		5	1652	1652	24.40	28.20	3.80	1.2	770					
9	9	GAGIN		5	1655	1655	25.30	28.00	2.70	1.2	390					
9	9	GAGIN		5	1658	1658	26.20	28.10	1.90	1.2	210					
RUN: 1																
11	11	FITZGERALD		28	1644	1700				1.5	525					ISOTHERMAL HOZE CHAM
11	11	FITZGERALD		28	1644	1700				0.94	122					
11	11	FITZGERALD		28	1644	1700				0.24	38					
11	11	FITZGERALD		28	1644	1700				0.14	4					
RUN: 2																
11	11	FITZGERALD		28	1719	1725				1.5	309					ISOTHERMAL HOZE CHAM
11	11	FITZGERALD		28	1719	1725				0.94	140					
11	11	FITZGERALD		28	1719	1725				0.24	38					
11	11	FITZGERALD		28	1719	1725				0.14	4					
RUN: 1																
13	13	AYESS		2	1647	1648	28.33	30.90	2.57	1.25	1275					PHOTO. RECORDS TAKEN
13	13	AYESS		2	1650	1651	27.20	30.80	3.60	1.25	1275					
13	13	AYESS		2	1651	1652	26.39	30.80	4.41	1.25	1425					
13	13	AYESS		2	1653	1654	25.53	30.50	5.07	1.25	1455					
13	13	AYESS		2	1655	1656	24.79	30.50	5.71	1.25	1500					
13	13	AYESS		2	1657	1658	24.69	30.40	5.71	1.25	1515					
RUN: 2																
13	13	AYESS		2	1704	1705	27.84	30.40	2.56	1.25	795					PHOTO. RECORDS TAKEN
13	13	AYESS		2	1706	1707	26.77	30.40	3.63	1.25	1200					
13	13	AYESS		2	1708	1709	22.98	30.40	7.42	1.25	1650					
13	13	AYESS		2	1710	1711	25.32	30.40	5.08	1.25	2100					
13	13	AYESS		2	1713	1714	23.72	30.40	3.68	1.25	1500					CHANGING AMBIENT AEROSOL?
13	13	AYESS		2	1715	1716	24.72	30.40	5.68	1.25	1500					
13	13	AYESS		2	1717	1718	24.72	30.40	5.68	1.25	1425					

EXPERIMENT #: 11

PROCESSING DATE: 81/07/29

NUCLEI MEASUREMENTS

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	COLD	PLATE TEMP HOT (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	ORS	EST K	REMARKS
RUN: 1															
14	14	HUDSON		9	1651	1711	18.37	25.06	6.69	.012	.21	262			F=99
14	14	HUDSON		9	1651	1711	18.74	24.75	6.01	.012	.14	213			F=99
14	14	HUDSON		9	1651	1711	19.12	24.44	5.32	.012	.11	118			F=99
14	14	HUDSON		9	1651	1711	19.49	24.13	4.65	.013	.04	11			F=99
14	14	HUDSON		9	1651	1711	19.86	23.81	3.95	.013	.02	3			F=99
14	14	HUDSON		9	1651	1711	20.22	23.50	3.25	.013	.01	1			F=99
14	14	HUDSON		9	1651	1711	20.59	23.18	2.57	.013	.01	1			F=99
14	14	HUDSON		9	1651	1711	20.99	22.86	1.82	.013	.02	3			F=99
RUN: 1															
15	15	ROGERS	FRED	23	1651	1655	18.37	25.06	6.69		1.86	1506			BACKGROUND 10 PER CC
15	15	ROGERS	FRED	23	1659	1660	19.12	24.44	5.32		1.46	148			AVERAGING LAST 4
15	15	ROGERS	FRED	23	1659	1660	19.49	24.13	4.65		1.87	1345			COUNTS CH. 2
15	15	ROGERS	FRED	23	1659	1660	19.86	23.81	3.95		.63	11958			
15	15	ROGERS	FRED	23	1659	1660	20.22	23.50	3.25		.265	11954			
15	15	ROGERS	FRED	23	1659	1660	20.59	23.18	2.57		.17	660			LAST PT IS CH 3 CNT
15	15	ROGERS	FRED	23	1659	1660	20.99	22.86	1.82		.07	125			
RUN: 1															
16	16	HUDSON		9	1659	1641	0	26.50	4.80	.018	.95	136			
16	16	HUDSON		9	1659	1641	0.50	26.25	4.24	.018	.54	102			
16	16	HUDSON		9	1659	1641	1.00	25.99	3.68	.024	.95	124			
16	16	HUDSON		9	1659	1641	1.50	25.73	3.12	.024	.55	108			
16	16	HUDSON		9	1659	1641	2.00	25.47	2.56	.024	.24	109			
16	16	HUDSON		9	1659	1641	2.50	25.21	1.99	.024	.55	105			
16	16	HUDSON		9	1659	1641	3.00	24.95	1.42	.024	.24	666			
RUN: 2															
16	16	HUDSON		9	1713	1714	1.40	26.50	4.80	.012	.94	1012			
16	16	HUDSON		9	1713	1714	1.80	26.25	4.24	.012	.54	900			
16	16	HUDSON		9	1713	1714	2.20	25.99	3.68	.012	.24	584			
16	16	HUDSON		9	1713	1714	2.60	25.73	3.12	.024	.55	682			
16	16	HUDSON		9	1713	1714	3.00	25.47	2.56	.024	.24	705			
RUN: 1															
17	17	WUJIC	HOWE	27	1641	1642	0	26.50	4.80	.012	.35	893			
17	17	WUJIC	HOWE	27	1641	1642	0.50	26.25	4.24	.012	.27	104			
17	17	WUJIC	HOWE	27	1641	1642	1.00	25.99	3.68	.012	.72	104			
17	17	WUJIC	HOWE	27	1641	1642	1.50	25.73	3.12	.024	.98	154			
17	17	WUJIC	HOWE	27	1641	1642	2.00	25.47	2.56	.024	.24	154			
RUN: 1															
18	18	HUDSON		9	1659	1649	23.85	26.50	3.15	.002	.4	921			
18	18	HUDSON		9	1659	1649	23.40	26.25	2.70	.002	.29	875			
18	18	HUDSON		9	1659	1649	23.00	25.99	2.24	.002	.19	646			
18	18	HUDSON		9	1659	1649	22.60	25.73	1.78	.002	.13	621			
18	18	HUDSON		9	1659	1649	22.20	25.47	1.32	.002	.12	621			
RUN: 1															
20	20	DOMONKOS		4	1640	1650					.49	610			

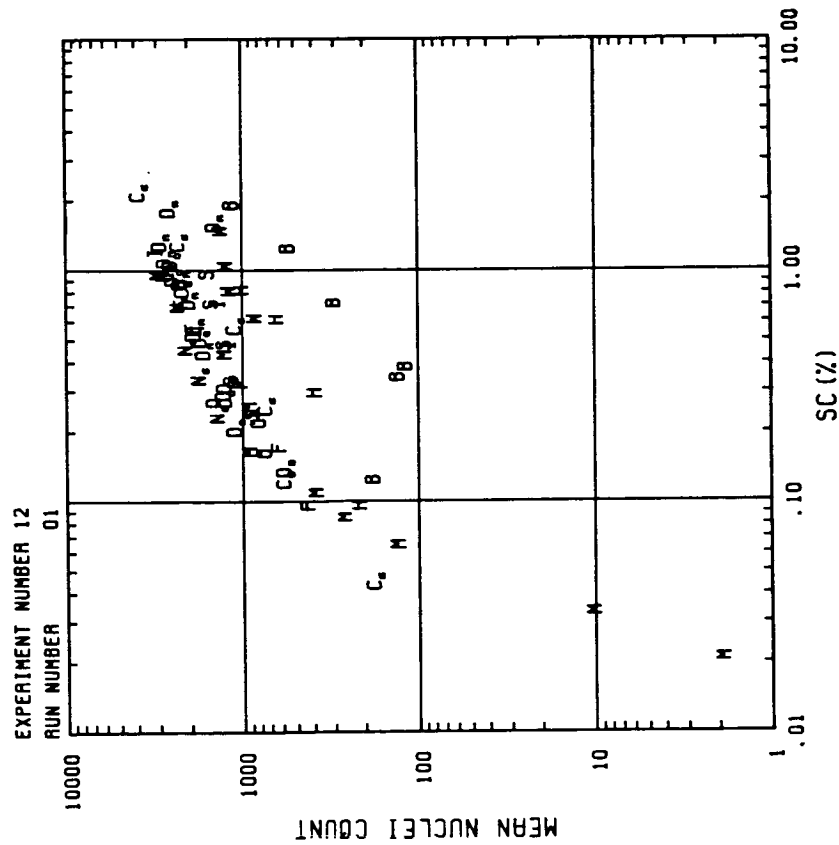
PROCESSING DATE: 81/07/29

NUCLEI MEASUREMENTS

EXPERIMENT #: 11

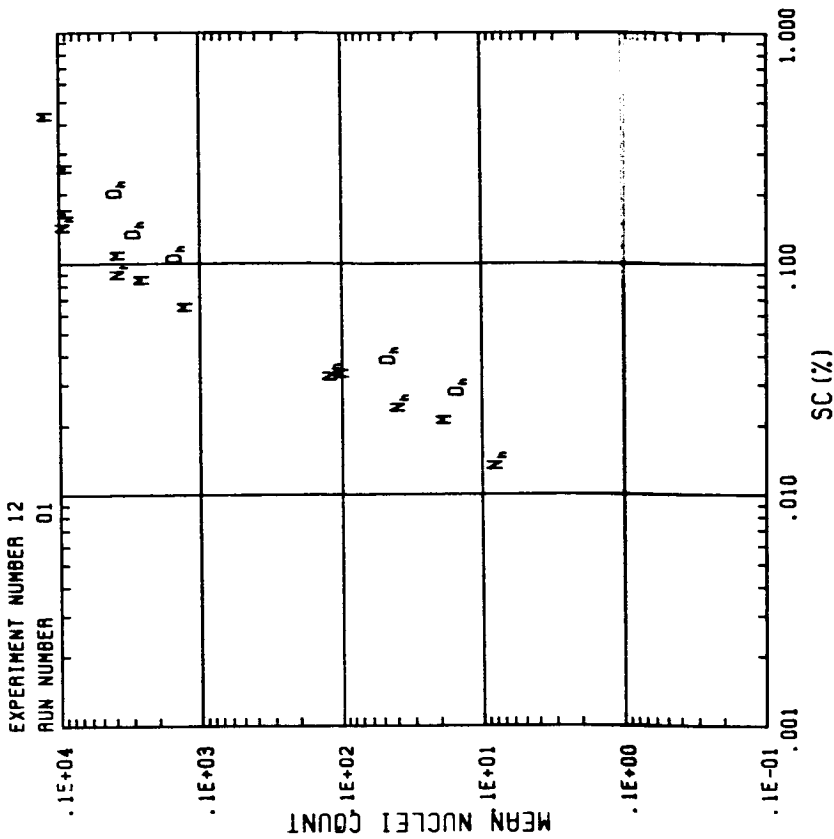
MACH NO.	DUCT NO.	OBSERVER	NO.	SAMPLE START	TIME END	COLD	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	ORS	EST K	REMARKS
							HOT (C)	NUCLEI COUNT							
20	20	DOMONKOS	4	1640	1650						.74	1120			
20	20	DOMONKOS	4	1640	1650						.96	1260			
20	20	DOMONKOS	4	1640	1650						1.37	1360			
RUN: 1															
21	21	TRUEBLOOD	26	1647	1642	19.95	25.00	5.05			.39	1313			
21	21	TRUEBLOOD	26	1650	1644	22.50	25.00	2.50			.25	1970			
21	21	TRUEBLOOD	26	1657	1658	22.50	25.00	2.50			.21	747			
21	21	TRUEBLOOD	26	1710	1711	25.00	25.00	0.00			1.73	637			
21	21	TRUEBLOOD	26	1710	1711	25.00	25.00	0.00			.114	327			
21	21	TRUEBLOOD	26	1710	1711	25.00	25.00	0.00			.089	214			
21	21	TRUEBLOOD	26	1710	1711	25.00	25.00	0.00			.068	112			
21	21	TRUEBLOOD	26	1717	1718	25.00	25.00	0.00			.335	911			
21	21	TRUEBLOOD	26	1717	1718	25.00	25.00	0.00			.022	216			
21	21	TRUEBLOOD	26	1717	1718	25.00	25.00	0.00			.0133	113			
RUN: 1															
24	24	SERPOLAY	25	1645	1650	28.10	31.10	3.00		.77	.33	730			
24	24	SERPOLAY	25	1700	1705	28.50	32.50	4.00		.77	.56	1080			
24	24	SERPOLAY	25	1720	1725	29.00	33.60	4.60		.77	.74	1230			
RUN: 1															
25	25	BORYS	3	1638	1642	17.60	24.80	7.20		2.12	2.18	1344			
25	25	BORYS	3	1655	1652	19.50	25.40	5.90		3.18	1.75	1920			
25	25	BORYS	3	1705	1706	21.50	25.00	3.50		4.03	.84	143			
25	25	BORYS	3	1710	1720	22.30	26.20	3.90		9.36	.33	364			
25	25	BORYS	3	1722	1729	25.60	26.90	1.30		10.8	.038	226			
RUN: 1															
26	26	HINDMAN	6	1643	1719	27.00	27.00	0.00		1.2	.12	43			E=1.4; D>=0.5 UM
26	26	HINDMAN	6	1643	1719	27.00	27.00	0.00		1.2	.12	23			E=0.76; D>=1.7 UM
26	26	HINDMAN	6	1643	1719	27.00	27.00	0.00		1.2	.059	16			E=0.78; D>=1.4 UM
26	26	HINDMAN	6	1643	1719	27.00	27.00	0.00		1.2	.027	12			E=0.02; D>=3 UM
26	26	HINDMAN	6	1643	1719	27.00	27.00	0.00		1.2	.016	12			E=0.0029; D>=.5 UM
RUN: 1															
27	27	OH TAKE	19	1645	1645					3.5	.31	311			101% R.H. NOMINAL
27	27	OH TAKE	19	1645	1645					3.5	.49	49			114% R.H. NOMINAL

INSTRUMENT COMPARISON



SDC/CFD RESULTS

INSTRUMENT COMPARISON



HAZE CHAMBER RESULTS

EXPERIMENT # 12

PURPOSE

INSTRUMENT COMPARISON WITH AMBIENT AEROSOL.

DESCRIPTION OF EXPERIMENT

DATE _____ TIME START _____ END _____
 10 OCT, 1980 0830 1002

NUCLEI TYPE

AMBIENT

GENERATION METHOD

NA

SIZE DISTRIBUTION SHAPING

NA

REMARKS

T 23.5 (C), TD 12 (C)

WEATHER SYNOPSIS

NONE

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS		OUTSIDE CONDITIONS					
AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG WS (M/S)	AVG WD (DEG)
25.0	21.0	852	NA	NA	NA	<1.0	360

PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

EXPERIMENT #: 12

MACH NO.	DUCT NO.	OBSERVER	SAMPLE START	SAMPLE TIME END	COLD	PLATE TEMP		VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
						HOT (C)	DELTA						
RUN: 1													
2	2	POLITOVICH	0926	0931	21.90	24.00	2.10		.17		785		
			0930	0931	21.80	24.20	2.40		.13		853		
RUN: 1													
5	5	KITCHEN	0830	0830	21.20	22.90	1.70		.13	187			
5	5	KITCHEN	0831	0831	19.90	22.90	3.00		.14	122			
5	5	KITCHEN	0832	0832	18.80	23.00	4.20		.17	321			
5	5	KITCHEN	0833	0833	17.60	23.00	5.60		.17	1184			
5	5	KITCHEN	0834	0834	16.40	23.00	6.40		.17	608			
5	5	KITCHEN	0835	0835	17.60	23.00	5.40		.17	300			
5	5	KITCHEN	0836	0836	18.90	23.10	4.20		.16	136		1.20	
5	5	KITCHEN	0837	0837	20.30	23.10	2.80		.16				
RUN: 2													
5	5	KITCHEN	0950	0950	22.50	24.70	2.20		.2	229			
5	5	KITCHEN	0951	0951	21.70	24.70	3.00		.36	445			
5	5	KITCHEN	0952	0952	21.00	24.70	3.70		.71	712			
5	5	KITCHEN	0953	0953	20.10	24.70	4.60		.85	1080			
5	5	KITCHEN	0954	0954	19.10	24.70	5.60		1.4	1650			
RUN: 1													
9	9	GAGIN	0828	0828	23.00	27.70	4.70		.86	1050			
9	9	GAGIN	0829	0829	23.50	27.60	4.10		.91	670			
9	9	GAGIN	0830	0830	25.00	27.80	2.80		.91	410			
9	9	GAGIN	0831	0831	25.90	27.70	1.80		.91	220			
RUN: 2													
9	9	GAGIN	0912	0912	23.10	28.10	5.10		.97	1670			
9	9	GAGIN	0914	0914	24.30	28.00	3.70		.53	1020			
9	9	GAGIN	0916	0916	25.80	27.70	2.70		.29	700			
9	9	GAGIN	0918	0918	26.30	26.00	1.70		.12	270			
RUN: 3													
9	9	GAGIN	0225	0225	22.90	28.00	5.10		1.01	2010			
9	9	GAGIN	0227	0227	24.20	27.90	3.70		.53	1100			
9	9	GAGIN	0230	0230	25.10	28.00	2.90		.31	650			
9	9	GAGIN	0231	0231	26.30	26.50	2.20		.19	320			
RUN: 4													
9	9	GAGIN	0235	0235	22.70	27.80	5.10		.92	2210			
9	9	GAGIN	0237	0237	25.00	27.70	4.10		.92	1000			
9	9	GAGIN	0242	0242	25.20	28.20	2.00		.25	390			
9	9	GAGIN	0242	0242	26.20	28.20	2.00		.16	310			
RUN: 1													
10	10	LALA	0836	0836	21.71	22.23	2.51		.25	950.5			
10	10	LALA	0838	0838	21.73	22.23	2.51		.25	880.8			
10	10	LALA	0839	0839	21.26	22.23	2.51		.25	1220.7			
10	10	LALA	0840	0840	21.26	22.23	2.51		.25	1220.3			
10	10	LALA	0841	0841	20.68	22.23	2.51		.25	1220.3			
10	10	LALA	0842	0842	19.90	22.23	2.51		.25	1336.9			
10	10	LALA	0843	0843	19.89	22.23	2.51		.25	1536.2			
10	10	LALA	0844	0844	19.19	22.23	2.51		.25	1536.2			
10	10	LALA	0845	0845	19.17	22.23	2.51		.25	1621.7			
RUN: 2													
10	10	LALA	0850	0850	22.07	24.58	2.51		.25	957.6			

N=1720.9 S=0.41

EXPERIMENT #: 12

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/09/26

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	COLD	HOT (C)	BELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	ORS N	EST K	REMARKS
RUN: 1															
15	15	ROGERS	FRED	23	0923	0924	18.36	25.05	6.70	6.21	1.86	2716			BACKGROUND 10 PER CC
15	15	ROGERS	FRED	23	0929	0930	18.90	22.44	6.25	5.23	1.32	1570			AEROSOL WILLIY CHANG
15	15	ROGERS	FRED	23	0933	0934	19.16	22.44	5.55	5.27	1.12	3705			NO AVENTRAGING
15	15	ROGERS	FRED	23	0937	0937	19.42	22.44	4.79	4.79	.92	2885			ACCI DENTALLY OMITTED
15	15	ROGERS	FRED	23	0942	0943	19.69	23.75	4.44	4.29	.74	2007			NEW CFD SERVO
15	15	ROGERS	FRED	23	0946	0947	19.95	23.75	3.32	3.32	.57	1943			PARAME TERS
15	15	ROGERS	FRED	23	0951	0951	20.47	23.54	3.32	3.32	.57	1777			N VS. SC SPECTRUM
15	15	ROGERS	FRED	23	0955	0956	20.47	23.31	2.84	2.84	.325	1317			WORTHLESS DUE TO
RUN: 1															
16	16	HUDSON		9	0849	0853	21.45	26.15	4.70	.0024	.54	1848			AEROSOL FLUCTUATIONS
16	16	HUDSON		9	0849	0853	22.45	25.44	3.65	.0024	.54	1648			
16	16	HUDSON		9	0857	0859	22.45	25.44	3.75	.0018	.54	1371			
16	16	HUDSON		9	0857	0859	22.45	25.44	3.65	.0018	.54	2339			
16	16	HUDSON		9	0903	0905	22.45	25.44	4.70	.0018	.54	1625			
16	16	HUDSON		9	0903	0905	22.45	25.44	3.65	.0018	.54	1320			
RUN: 2															
16	16	HUDSON		9	0912	0916	21.45	26.15	4.70	.0024	.56	195			
16	16	HUDSON		9	0912	0916	22.30	25.44	3.65	.0024	.56	2133			
16	16	HUDSON		9	0926	0929	21.70	26.40	4.70	.0018	.56	1236			
16	16	HUDSON		9	0926	0936	22.35	26.40	3.65	.0024	.56	1123			
16	16	HUDSON		9	0932	0936	22.70	26.40	4.70	.0024	.56	1123			
RUN: 3															
16	16	HUDSON		9	0945	0947	21.35	26.05	4.70	.0018	.56	194			
16	16	HUDSON		9	0945	0947	22.70	26.05	3.65	.0018	.56	1257			
16	16	HUDSON		9	0953	0959	22.35	26.05	4.70	.0036	.56	1123			
16	16	HUDSON		9	0953	0959	22.70	26.05	3.65	.0036	.56	1123			
RUN: 1															
17	17	FITZGERALD		28	0906	0907	21.45	25.70	4.70	.0018	.56	1445	1445		
17	17	FITZGERALD		28	0910	0910	22.30	25.70	3.65	.0018	.56	1855	1855		
17	17	FITZGERALD		28	0917	0917	22.35	26.05	4.70	.0018	.56	2105	2105		
17	17	FITZGERALD		28	0919	0920	22.70	26.05	3.65	.0036	.56	2286	2286		
RUN: 1															
18	18	HUDSON		9	0857	0859	23.05	25.70	4.70	.0018	.56	197			
18	18	HUDSON		9	0903	0905	23.05	25.70	3.65	.0018	.56	1154			
18	18	HUDSON		9	0912	0916	23.05	26.05	4.70	.0024	.56	1154			
18	18	HUDSON		9	0926	0929	23.05	26.05	3.65	.0024	.56	2240			
18	18	HUDSON		9	0932	0936	23.40	26.05	4.70	.0024	.56	1168			
18	18	HUDSON		9	0945	0949	23.40	26.05	3.65	.0036	.56	1168			
RUN: 1															
20	20	DOMONKOS		4	0940	0950					.65	890			

PROCESSING DATE: 01/09/26

NUCLEI MEASUREMENTS

EXPERIMENT #: 12

MACH NO. DUCT NO. OBSERVER NO. SAMPLE TIME START END COLD HOT (C) DELTA VOLUME SAMPLED (L) SUPER SAT (%) NUCLEI COUNT OBS N EST K REMARKS

20 20 DOMONKOS 4 0940 0940 25.00 25.00 0.00 10.6 0.85 1230
 20 20 DOMONKOS 4 0940 0940 25.00 25.00 0.00 10.6 1.54 1290
 20 20 DOMONKOS 4 0940 0940 25.00 25.00 0.00 10.6 1.54 1370

RUN: 1
 21 21 TRUEBLOOD 26 0832 0834 20.00 25.00 5.00 10.6 1.15 1152
 21 21 TRUEBLOOD 26 0842 0847 25.00 25.00 0.00 10.6 0.42 1258
 21 21 TRUEBLOOD 26 0900 0901 25.00 25.00 0.00 10.6 0.08 1326
 21 21 TRUEBLOOD 26 0900 0901 25.00 25.00 0.00 10.6 0.09 1402
 21 21 TRUEBLOOD 26 0900 0901 25.00 25.00 0.00 10.6 0.17 1462
 21 21 TRUEBLOOD 26 0912 0913 25.00 25.00 0.00 10.6 0.25 1497
 21 21 TRUEBLOOD 26 0912 0913 25.00 25.00 0.00 10.6 0.35 1658

RUN: 2
 21 21 TRUEBLOOD 26 0924 0925 25.00 25.00 0.00 10.6 0.68 1253
 21 21 TRUEBLOOD 26 0924 0925 25.00 25.00 0.00 10.6 0.89 1372
 21 21 TRUEBLOOD 26 0924 0925 25.00 25.00 0.00 10.6 1.13 1453

RUN: 1
 24 24 SERPOLAY 25 0840 0847 25.30 29.30 4.00 10.6 0.34 1080
 24 24 SERPOLAY 25 0855 0905 25.00 29.80 4.80 10.6 0.59 1960
 24 24 SERPOLAY 25 0910 0915 26.10 30.70 4.60 10.6 0.75 2290
 24 24 SERPOLAY 25 0945 0955 26.20 27.80 1.60 10.6 0.18 450

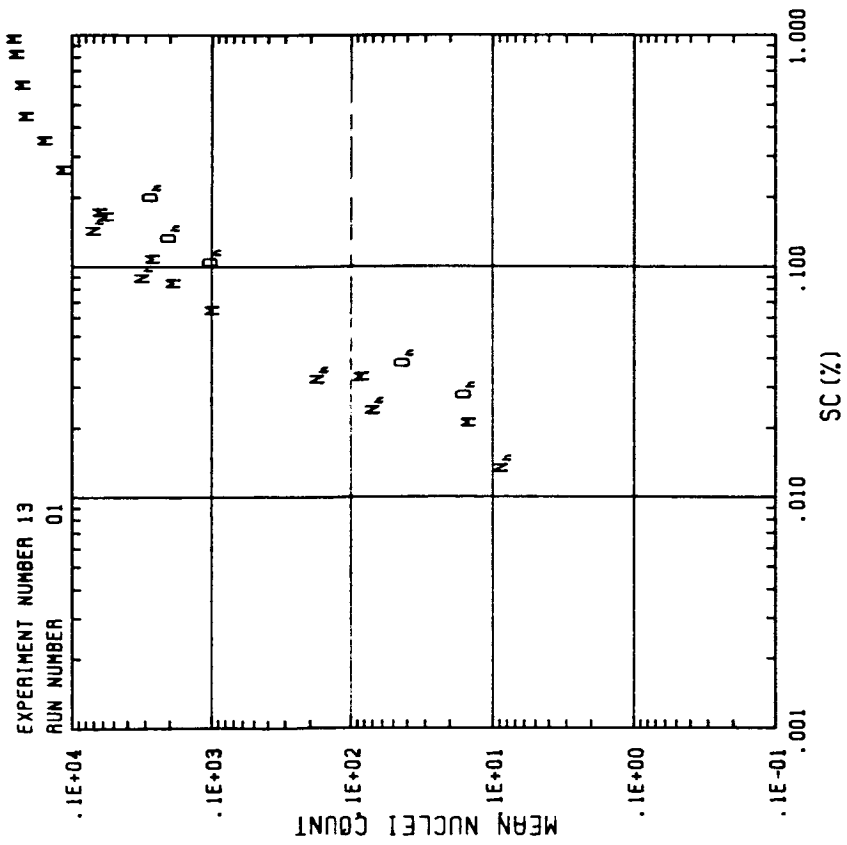
RUN: 1
 25 25 BORYS 33 0840 0846 22.20 23.40 1.20 10.6 0.45 184
 25 25 BORYS 33 0850 0856 21.50 23.70 2.20 10.6 0.125 608
 25 25 BORYS 33 0906 0915 20.80 23.80 3.00 10.6 0.26 778
 25 25 BORYS 33 0929 0935 19.10 23.80 4.70 10.6 0.58 1160
 25 25 BORYS 33 0943 0949 18.20 23.90 5.70 10.6 0.93 2373
 25 25 BORYS 33 0948 0955 16.20 23.70 7.50 10.6 1.32 4032
 25 25 BORYS 33 0948 0955 16.20 23.70 7.50 10.6 1.19 4032

RUN: 1
 26 26 HINDMAN 6 0843 0932 09.00 09.32 0.32 10.6 0.00 184
 26 26 HINDMAN 6 0843 0932 09.00 09.32 0.32 10.6 0.00 608
 26 26 HINDMAN 6 0843 0932 09.00 09.32 0.32 10.6 0.00 778
 26 26 HINDMAN 6 0843 0932 09.00 09.32 0.32 10.6 0.00 1160
 26 26 HINDMAN 6 0843 0932 09.00 09.32 0.32 10.6 0.00 2373
 26 26 HINDMAN 6 0843 0932 09.00 09.32 0.32 10.6 0.00 4032

RUN: 1
 27 27 OHTAKE 19 0833 0833 10.5 125% R.H. NOMINAL 1.737

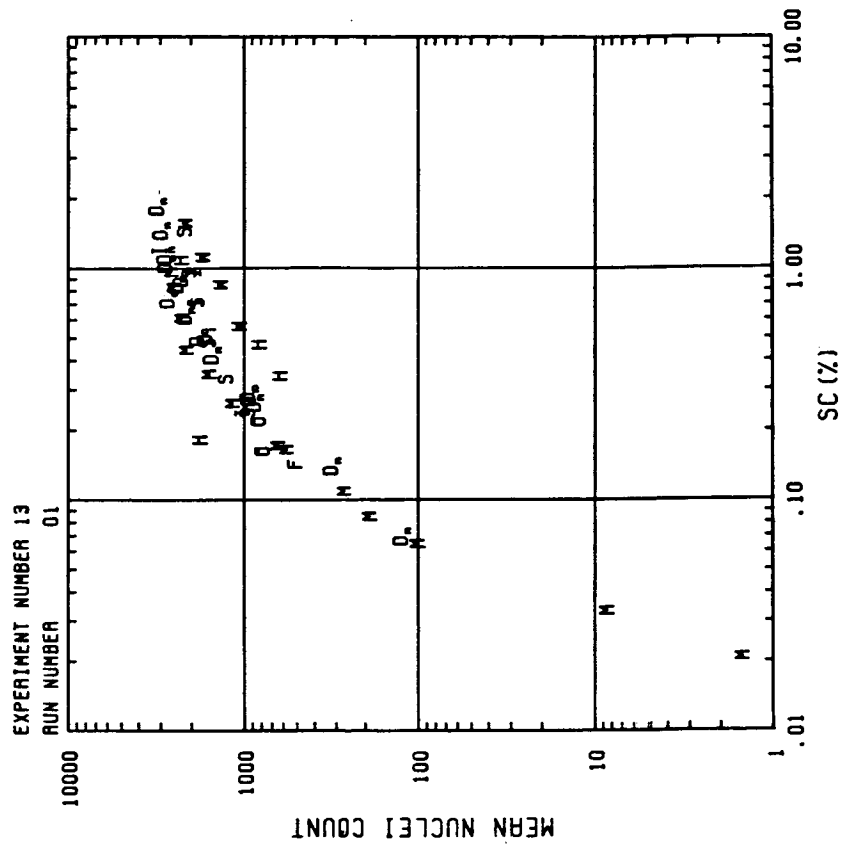
INSTRUMENT DOWN.
 ALL DATA FROM ALL PROBLEMS OFF. USE 1/8" SPHERE CALIBR. AREA. A LOT OF COUNTS. AIR DRY. TO BE SUBSATURATED HAZE CHAMBER.

INSTRUMENT COMPARISON



HAZE CHAMBER RESULTS

INSTRUMENT COMPARISON



SDC/CFD RESULTS

EXPERIMENT # 13

PURPOSE

POLYDISPERSE FOR MODERATE K.

DESCRIPTION OF EXPERIMENT

DATE 10 OCT 1980 TIME START 1017 END 1300

NUCLEI TYPE (NH4)2SO4

GENERATION METHOD

NANOMIST, 5.5 PSI, PREDILUTION 150 BR., 1.77 G/L SOLUTION.

SIZE DISTRIBUTION SHAPING

NONE

REMARKS

DILUTION FLOW RATE 590 L/MIN., UNSTABLE, SEE STRIPCHART FOR REFERENCE.

WEATHER SYNOPSIS

NONE

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS		OUTSIDE CONDITIONS					
AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG WS (M/S)	AVG WD (DEG)
27.0	21.0	850	NA	NA	NA	NA	NA

PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

EXPERIMENT #: 13

MACH NO. DUCT NO. OBSERVER NO. SAMPLE TIME START END COLD HOT PLATE TEMP DELTA VOLUME SAMPLED (L) SUPER SAT (%) NUCLEI COUNT OBS N EST R REMARKS

MACH NO.	DUCT NO.	OBSERVER	NO.	SAMPLE TIME START	END	COLD	HOT	PLATE TEMP	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST R	REMARKS
RUN: 1															
2	2	ROGERS: DAVE	22	1205	1207	23.80	26.00	26.00	2.10		.17		827		
2	2	ROGERS: DAVE	22	1208	1209	23.80	26.20	26.20	2.40		.23		867		
RUN: 1															
9	9	GAGIN	5	1144		22.70	28.10		5.40	1.2	1.14	2340			
9	9	GAGIN	5	1147		24.60	28.30		3.70	1.2	.49	850			
9	9	GAGIN	5	1152		25.50	28.50		3.00	1.2	.36	650			
9	9	GAGIN	5	1154		26.10	28.30		2.20	1.2	.19	1840			
RUN: 2															
9	9	GAGIN	5	1159		25.60	28.10		2.50	1.2	.44	2450			
9	9	GAGIN	5	1201		26.30	28.30		2.00	1.2	.22	1030			
9	9	GAGIN	5	1203		26.30	28.30		2.00	1.2	.16	740			
9	9	GAGIN	5	1206		26.30	28.30		2.00	1.2	.16	240			
RUN: 3															
9	9	GAGIN	5	1211		23.20	27.90		4.70	1.2	.85	1440			
9	9	GAGIN	5	1213		24.60	28.20		3.60	1.2	.49	930			
9	9	GAGIN	5	1216		25.10	28.40		3.30	1.2	.42	640			
9	9	GAGIN	5	1218		26.20	28.40		2.20	1.2	.42	640			
RUN: 1															
10	10	LALA	13	1144		23.00	25.20		2.20	.4	.25	930			
10	10	LALA	13	1145		23.50	25.20		1.70	.4	.17	1380			
10	10	LALA	13	1147		23.50	25.20		1.70	.4	.17	1380			
10	10	LALA	13	1149		23.50	25.20		1.70	.4	.17	1380			
10	10	LALA	13	1151		23.50	25.20		1.70	.4	.17	1380			
10	10	LALA	13	1153		23.50	25.20		1.70	.4	.17	1380			
10	10	LALA	13	1155		23.50	25.20		1.70	.4	.17	1380			
10	10	LALA	13	1157		23.50	25.20		1.70	.4	.17	1380			
10	10	LALA	13	1159		23.50	25.20		1.70	.4	.17	1380			
10	10	LALA	13	1161		23.50	25.20		1.70	.4	.17	1380			
10	10	LALA	13	1163		23.50	25.20		1.70	.4	.17	1380			
RUN: 2															
10	10	LALA	13	1155		23.40	25.60		2.20	.4	.23	891.9			
10	10	LALA	13	1156		23.40	25.60		2.20	.4	.23	891.9			
10	10	LALA	13	1157		23.40	25.60		2.20	.4	.23	891.9			
10	10	LALA	13	1158		23.40	25.60		2.20	.4	.23	891.9			
10	10	LALA	13	1159		23.40	25.60		2.20	.4	.23	891.9			
10	10	LALA	13	1160		23.40	25.60		2.20	.4	.23	891.9			
10	10	LALA	13	1161		23.40	25.60		2.20	.4	.23	891.9			
10	10	LALA	13	1162		23.40	25.60		2.20	.4	.23	891.9			
10	10	LALA	13	1163		23.40	25.60		2.20	.4	.23	891.9			
RUN: 1															
11	11	FITZGERALD	28	1255		26.01	28.50		2.49		.15	740			ISOTHERMAL CHAMBER
11	11	FITZGERALD	28	1255		26.01	28.50		2.49		.15	740			ISOTHERMAL CHAMBER
11	11	FITZGERALD	28	1255		26.01	28.50		2.49		.15	740			ISOTHERMAL CHAMBER
11	11	FITZGERALD	28	1255		26.01	28.50		2.49		.15	740			ISOTHERMAL CHAMBER
RUN: 1															
13	13	AYERS	2	1145		26.01	28.50		2.49		.15	740			ISOTHERMAL CHAMBER
13	13	AYERS	2	1147		26.01	28.50		2.49		.15	740			ISOTHERMAL CHAMBER

AEROSOL FLUCTUATIONS

ISOTHERMAL CHAMBER

EXPERIMENT #: 13
 PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	SAMPLE END	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
							COLD	HOT (C)							
RUN: 1															
13	13	AYERS		2	1150	1150	23.14	28.50	4.36		1965				
13	13	AYERS		2	1150	1150	22.77	28.40	5.03	.75	2025				
13	13	AYERS		2	1151	1152	22.77	28.40	5.03	1.25	3150				
RUN: 2															
13	13	AYERS		2	1153	1153	23.77	28.40	5.63	1.25	3975			PHOTOGRAPHIC RECORDS TAKEN	
13	13	AYERS		2	1154	1154	23.40	28.40	5.00		2025				
13	13	AYERS		2	1155	1155	23.93	28.30	4.37	.75	2025				
13	13	AYERS		2	1156	1157	24.76	28.30	3.54	.25	1760				
13	13	AYERS		2	1158	1159	25.80	28.30	2.50						
RUN: 3															
13	13	AYERS		2	1243	1243	27.03	29.60	2.57	.25	1283			PHOTOGRAPHIC RECORDS TAKEN	
13	13	AYERS		2	1244	1244	26.05	29.60	4.36	.75	2130				
13	13	AYERS		2	1245	1246	25.14	29.50	4.36	.75	2400				
13	13	AYERS		2	1246	1246	24.35	29.40	5.62	1.25	2850				
13	13	AYERS		2	1247	1248	23.78	29.40			3750				
RUN: 1															
14	14	HUDSON		9	1158	1158				.08	283			E=41	
14	14	HUDSON		9	1158	1158				.11	209			E=41	
14	14	HUDSON		9	1158	1158				.08	106			E=41	
14	14	HUDSON		9	1158	1158				.029	1.62			E=41	
14	14	HUDSON		9	1158	1158				.21	306			E=41	
14	14	HUDSON		9	1158	1158				.11	109			E=41	
14	14	HUDSON		9	1158	1158				.029	1.92			E=41	
RUN: 2															
14	14	HUDSON		9	1221	1226				.21	338			E=41	
14	14	HUDSON		9	1221	1226				.11	123			E=41	
14	14	HUDSON		9	1221	1226				.04	5.52			E=41	
14	14	HUDSON		9	1221	1226				.029				E=41	
RUN: 1															
15	15	ROGERS	FRED	23	1159	1159	18.36	25.06	6.70	1.86	3291			BACKGROUND 10 PER CC	
15	15	ROGERS	FRED	23	1200	1200	18.74	24.75	6.01	1.46	3114			AVERAGING LAST 4	
15	15	ROGERS	FRED	23	1201	1201	19.11	24.43	5.44	1.15	2554			COUNTS PER SS	
15	15	ROGERS	FRED	23	1210	1210	19.87	23.81	3.44	.63	2554			MAY NEED TO REDUCE	
15	15	ROGERS	FRED	23	1210	1210	20.25	23.50	2.55	.265	1596			MAIN FLOW FOR	
15	15	ROGERS	FRED	23	1220	1220	20.99	23.19	2.66	.17	1915			LOWEST SS PTS	
15	15	ROGERS	FRED	23	1231	1232	21.35	22.56	1.21	.07	334			LAST PNT CH 3	
RUN: 1															
16	16	HUDSON		9	1148	1148				.08	2120				
16	16	HUDSON		9	1148	1148				.08	1526				
16	16	HUDSON		9	1148	1148				.08	2766				
16	16	HUDSON		9	1148	1148				.033	1802				
16	16	HUDSON		9	1148	1148				.004	1029				
16	16	HUDSON		9	1148	1148				.004	1034				
16	16	HUDSON		9	1148	1148				.002					
RUN: 2															
16	16	HUDSON		9	1229	1233	21.35	26.10	4.75	.0024					

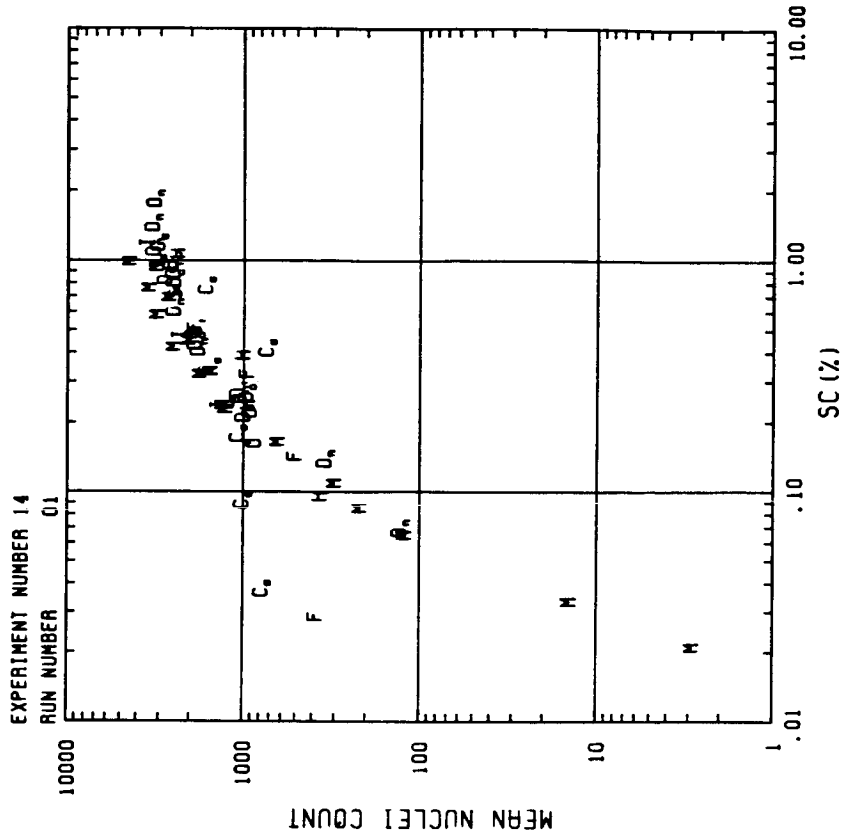
PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

EXPERIMENT #: 13

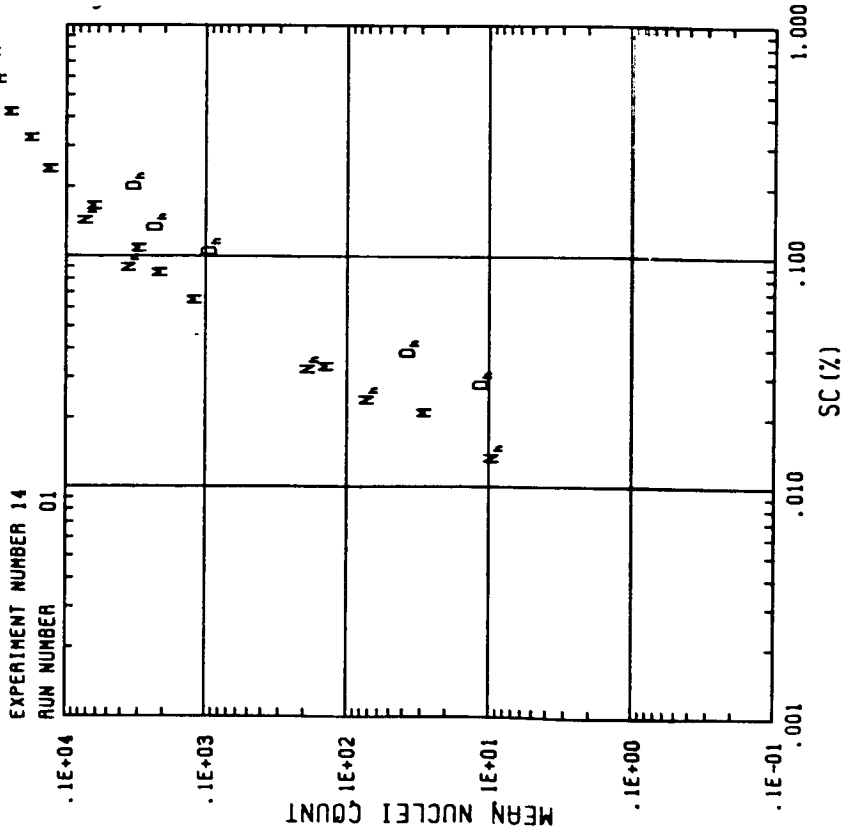
MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS	EST K	REMARKS
							COLD	HOT (C)							
16	16	HUDSON		9	1229	1233	21.70	25.30	3.60	.024	.52	1941			
16	16	HUDSON		9	1229	1233	22.20	24.85	2.65	.024	.78	1069			
16	16	HUDSON		9	1238	1242	21.35	25.10	4.75	.003	.92	2498			
16	16	HUDSON		9	1238	1242	21.70	25.30	3.60	.003	.52	2163			
16	16	HUDSON		9	1238	1242	22.20	24.85	2.65	.003	.52	1146			
RUN: 1															
18	18	HUDSON		9	1146	1148	23.15	26.85	2.70	.0018	.29	990			
18	18	HUDSON		9	1155	1200	22.65	26.35	3.70	.003	.21	1910			
18	18	HUDSON		9	1206	1209	22.20	26.85	4.60	.003	.21	2532			
18	18	HUDSON		9	1229	1233	22.20	26.85	4.60	.003	1.02	2938			
18	18	HUDSON		9	1238	1242	22.40	26.70	4.50	.003	.74	2847			
RUN: 1															
20	20	DOMONKOS		4	1130	1135					.59	1103			
20	20	DOMONKOS		4	1130	1135					.89	1410			
20	20	DOMONKOS		4	1130	1135					1.17	1780			
RUN: 1															
21	21	TRUEBLOOD		26	1145	1147	19.48	25.00	5.52	.02	1.01	2727			
21	21	TRUEBLOOD		26	1159	1150	20.40	25.00	4.60	.02	.67	2610			
21	21	TRUEBLOOD		26	1157	1158	19.59	25.00	5.43	.02	.67	2610			
21	21	TRUEBLOOD		26	1157	1202	21.97	25.00	3.60	.02	.29	2374			
21	21	TRUEBLOOD		26	1159	1206	22.38	25.00	3.60	.02	.73	1212			
21	21	TRUEBLOOD		26	1159	1210	22.00	25.00	3.60	.02	.73	1669			
21	21	TRUEBLOOD		26	1159	1210	22.00	25.00	3.60	.02	.73	5239			
21	21	TRUEBLOOD		26	1159	1210	22.00	25.00	3.60	.02	.73	5239			
21	21	TRUEBLOOD		26	1159	1210	22.00	25.00	3.60	.02	.73	5239			
21	21	TRUEBLOOD		26	1159	1210	22.00	25.00	3.60	.02	.73	5239			
21	21	TRUEBLOOD		26	1159	1210	22.00	25.00	3.60	.02	.73	5239			
21	21	TRUEBLOOD		26	1159	1210	22.00	25.00	3.60	.02	.73	5239			
21	21	TRUEBLOOD		26	1159	1210	22.00	25.00	3.60	.02	.73	5239			
RUN: 1															
24	24	SERPOLAY		25	1155	1200	27.40	29.40	2.00	.77	.15	530			ONLY ONE MEASUREMENT AT S=000.150
24	24	SERPOLAY		25	1155	1200	27.40	29.40	2.00	.77	.15	530			
RUN: 1															
26	26	HINDMAN		6	1148	1237									INSTRUMENT DOWN, REPAIRING CHAMBER.
26	26	HINDMAN		6	1148	1237									

INSTRUMENT COMPARISON



SDC/CFD RESULTS

INSTRUMENT COMPARISON



HAZE CHAMBER RESULTS

EXPERIMENT # 14

PURPOSE

SAME AS #13.

DESCRIPTION OF EXPERIMENT

DATE 10 OCT. 1980 TIME START 1335 END 1444

NUCLEI TYPE (NH4)2SO4, POLYDISPERSE.

GENERATION METHOD

NANOMIST, 5.5 PSI, PREDILUTION 150 BR., 1.77 G/L SOLUTION.

SIZE DISTRIBUTION SHAPING

NONE

REMARKS

STABILITY VARIES, SEE STRIPCHART FOR ADJUSTMENTS.

WEATHER SYNOPSIS

NONE

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS		OUTSIDE CONDITIONS					
AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG WS (M/S)	AVG WD (DEG)
27.0	21.0	850	NA	NA	NA	NA	NA

EXPERIMENT #: 14

PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	SAMPLE END	COLD	HOT	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
RUN: 1															
2	2	POLITOVICH		20	1428	1431	18.90	21.60	2.70	1.2	.17	950	950		
RUN: 2															
9	9	GAGIN		5	1347		22.90	28.30	5.40	1.2	1.12	2480			
9	9	GAGIN		5	1349		24.60	27.90	3.30	1.2	.41	1050			
9	9	GAGIN		5	1352		26.60	28.00	1.40	1.2	.25	980			
RUN: 3															
9	9	GAGIN		5	1356		23.30	28.10	4.80	1.2	.87	1950			
9	9	GAGIN		5	1402		25.60	28.30	2.70	1.2	.22	640			
9	9	GAGIN		5	1408		26.40	28.10	1.70	1.2	.12	220			
RUN: 4															
9	9	GAGIN		5	1412		22.80	28.00	5.20	1.2	1.08	2670			
9	9	GAGIN		5	1414		24.50	28.00	3.50	1.2	.49	940			
9	9	GAGIN		5	1416		25.60	27.90	2.30	1.2	.19	550			
9	9	GAGIN		5	1418		26.40	28.40	2.00	1.2	.16	300			
RUN: 5															
10	10	LALA		13	1348		23.20	25.80	2.60	.4	.55	1318			
10	10	LALA		13	1349		22.70	25.70	3.00	.4	.55	1549			
10	10	LALA		13	1351		22.70	25.70	3.00	.4	.55	1549			
10	10	LALA		13	1352		22.70	25.70	3.00	.4	.55	1549			
10	10	LALA		13	1353		22.70	25.70	3.00	.4	.55	1549			
10	10	LALA		13	1354		22.70	25.70	3.00	.4	.55	1549			
10	10	LALA		13	1355		22.70	25.70	3.00	.4	.55	1549			
10	10	LALA		13	1356		22.70	25.70	3.00	.4	.55	1549			
RUN: 6															
10	10	LALA		13	1357		23.10	25.80	2.70	.4	.37	1016			
10	10	LALA		13	1358		22.70	25.70	3.00	.4	.55	1084			
10	10	LALA		13	1359		22.70	25.70	3.00	.4	.55	1084			
10	10	LALA		13	1401		22.70	25.70	3.00	.4	.55	1084			
10	10	LALA		13	1402		22.70	25.70	3.00	.4	.55	1084			
10	10	LALA		13	1403		22.70	25.70	3.00	.4	.55	1084			
10	10	LALA		13	1404		22.70	25.70	3.00	.4	.55	1084			
10	10	LALA		13	1405		22.70	25.70	3.00	.4	.55	1084			
RUN: 7															
11	11	FITZGERALD		28	1337	1353	20.90	22.50	1.60	.4	.15	773	773		
11	11	FITZGERALD		28	1337	1353	20.90	22.50	1.60	.4	.15	377	377		
11	11	FITZGERALD		28	1337	1353	20.90	22.50	1.60	.4	.15	19.9	19.9		
11	11	FITZGERALD		28	1337	1353	20.90	22.50	1.60	.4	.15	7.6	7.6		
RUN: 8															
11	11	FITZGERALD		28	1359	1416	20.90	22.50	1.60	.4	.15	758	758		
11	11	FITZGERALD		28	1359	1416	20.90	22.50	1.60	.4	.15	358	358		
11	11	FITZGERALD		28	1359	1416	20.90	22.50	1.60	.4	.15	1.6	1.6		
11	11	FITZGERALD		28	1359	1416	20.90	22.50	1.60	.4	.15	1.06	1.06		

PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

EXPERIMENT #: 14

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	SAMPLE END	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	ORBS	EST K	REMARKS
							COLD	HOT (C)							
RUN: 1															
13	13	AYERS		2	1352	1353	27.23	29.80	2.57	.25	1440				
13	13	AYERS		2	1353	1354	26.20	29.80	3.60	.75	2400				
13	13	AYERS		2	1356	1357	25.22	29.80	4.38	.75	2625				
13	13	AYERS		2	1357	1358	24.50	29.80	5.10	1.25	3255				
13	13	AYERS		2	1358	1359	23.87	29.50	5.63		3675				
RUN: 2															
13	13	AYERS		2	1359	1400	23.89	29.50	5.61	1.25	4050				
13	13	AYERS		2	1400	1401	24.35	29.40	5.05	.75	2000				
13	13	AYERS		2	1402	1403	24.32	29.30	4.38	.75	2250				
13	13	AYERS		2	1405	1406	25.75	29.30	3.53	.25	1935				
13	13	AYERS		2	1406	1407	23.27	29.20	3.93	.25	1110				
RUN: 1															
14	14	HUDSON		9	1345	1355	18.36	25.06	6.70	.0056	396			F=41	
14	14	HUDSON		9	1346	1356	19.74	24.75	6.01	.0056	391			F=41	
14	14	HUDSON		9	1347	1357	19.49	24.44	5.33	.0056	425			F=41	
14	14	HUDSON		9	1348	1358	19.87	23.81	4.64	.0056	623			F=41	
14	14	HUDSON		9	1349	1359	20.25	23.50	3.94	.0056	273			F=55	
14	14	HUDSON		9	1407	1407	20.69	23.19	3.59	.0016	1093			F=55	
14	14	HUDSON		9	1407	1407	21.35	22.86	1.81	.0016	116			F=55	
14	14	HUDSON		9	1407	1411	21.35	22.86	1.81	.0016	116			F=55	
RUN: 1															
15	15	ROGERS	FRED	23	1337	1338	18.36	25.06	6.70	1.86	378				
15	15	ROGERS	FRED	23	1345	1346	19.74	24.75	6.01	1.15	348				
15	15	ROGERS	FRED	23	1350	1351	19.87	24.44	5.33	1.15	348				
15	15	ROGERS	FRED	23	1358	1359	20.25	23.81	4.64	.63	292				
15	15	ROGERS	FRED	23	1406	1407	20.69	23.50	3.94	.265	1093				
15	15	ROGERS	FRED	23	1406	1411	21.35	23.19	3.59	.107	116				
15	15	ROGERS	FRED	23	1410	1411	21.35	22.86	1.81	.107	116				
RUN: 1															
16	16	HUDSON		9	1420	1424	35	35	4.75	.0024	379				
16	16	HUDSON		9	1420	1424	35	35	4.75	.0024	379				
16	16	HUDSON		9	1423	1423	35	35	4.75	.0024	379				
16	16	HUDSON		9	1423	1423	35	35	4.75	.0024	379				
16	16	HUDSON		9	1423	1423	35	35	4.75	.0024	379				
16	16	HUDSON		9	1423	1423	35	35	4.75	.0024	379				
16	16	HUDSON		9	1423	1423	35	35	4.75	.0024	379				
16	16	HUDSON		9	1423	1423	35	35	4.75	.0024	379				
RUN: 1															
17	17	WUJICIECHOWSKI		27	1343	1344	14.1	10	3.35	.25	1598	1312			
17	17	WUJICIECHOWSKI		27	1347	1349	14.9	9.5	3.35	.25	1598	1598			
17	17	WUJICIECHOWSKI		27	1352	1353	14.9	9.5	3.35	.25	1598	1598			
17	17	WUJICIECHOWSKI		27	1355	1357	14.9	9.5	3.35	.25	1598	1598			
RUN: 1															
18	18	HUDSON		9	1327	1337	22.80	26.15	3.35	.0054	1965				

BACKGROUND CNT
10 PER CC.
DROPSIZE PLATEAU
BAD AT LOWER SS
LOWER MAIN FLO
FROM 145 TO 1315.
LAST FEW MIN 15 PCT.
AEROSOL LOW 15 PCT.

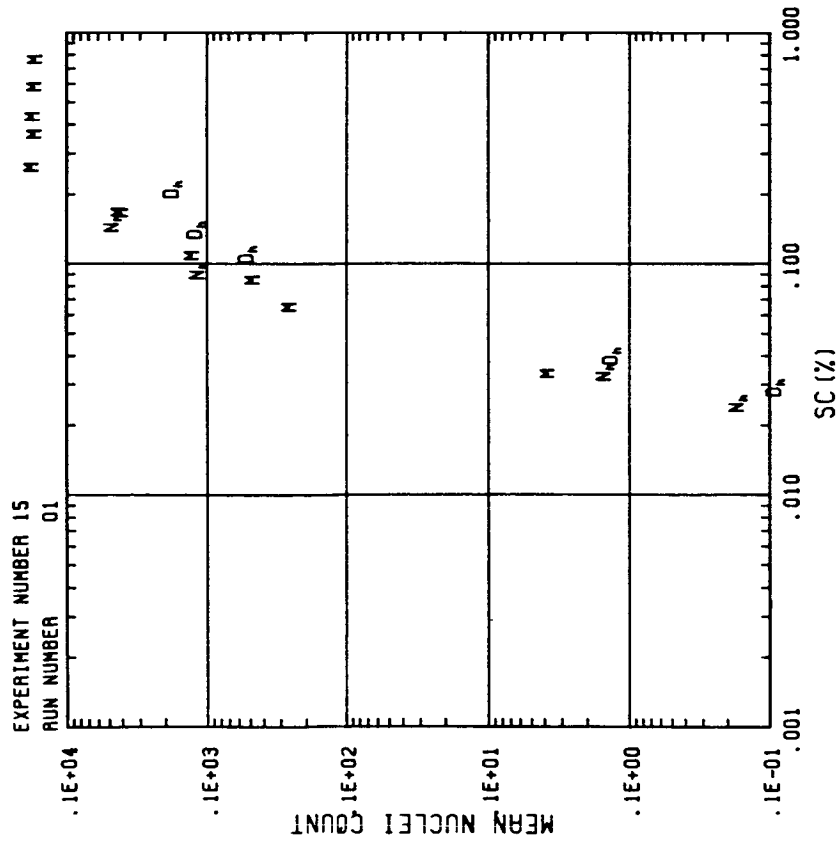
PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

EXPERIMENT #: 14

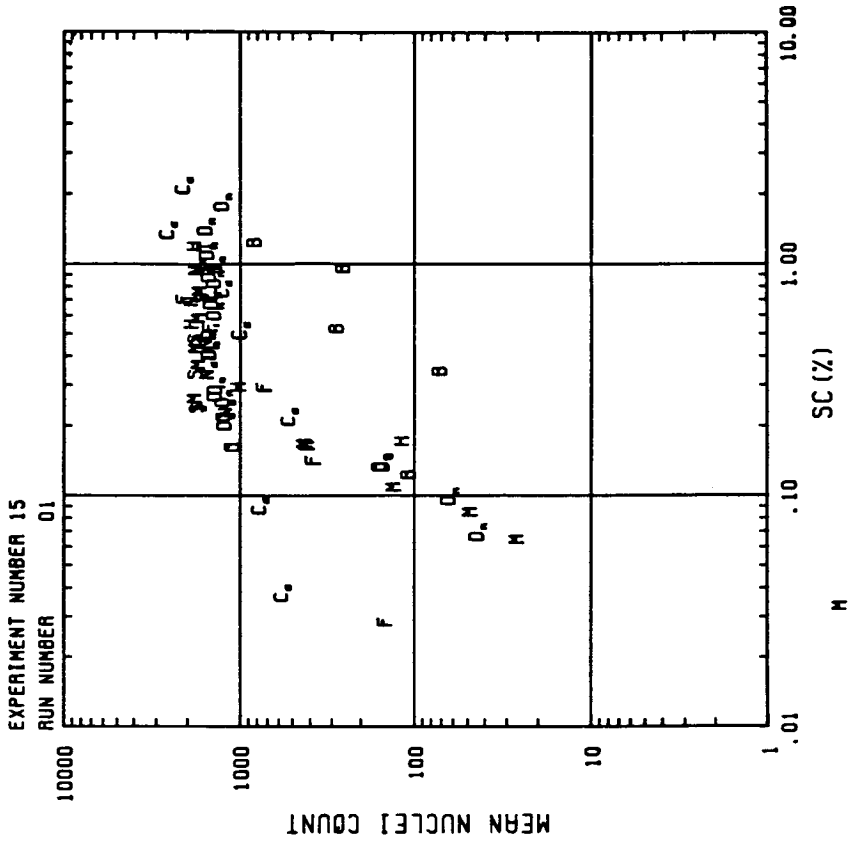
MACH NO.	DUCT NO.	NAME	OBSERVER	NG.	SAMPLE START	TIME END	COLD	HOT	PLATE TEMP	BELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	ORS - N	EST K	REMARKS
18	18	HUDSON		9	1402	1406	23:20	25:05		2.75	.0024	.27	1051			
18	18	HUDSON		9	1420	1425	22:25	25:00		2.55	.0024	.84	1018			
18	18	HUDSON		9	1434	1438	22:00	27:00		5.00	.0024	.1	2713			
RUN: 1																
21	21	TRUERBLOOD		26	1348	1349	19:52	25:00		5.10		1.04	4628.6			
21	21	TRUERBLOOD		26	1352	1357	21:08	25:00		4.92		.803	3530			
21	21	TRUERBLOOD		26	1356	1403	22:08	25:00		3.32		.614	3230			
21	21	TRUERBLOOD		26	1406	1407	22:50	25:00		2.50		.344	1852			
21	21	TRUERBLOOD		26	1410	1412	22:52	25:00		0.08		.173	1371			
21	21	TRUERBLOOD		26	1421	1422	25:00	25:00		0.00		.114	640.7			
21	21	TRUERBLOOD		26	1421	1422	25:00	25:00		0.00		.109	315.25			
21	21	TRUERBLOOD		26	1421	1422	25:00	25:00		0.00		.068	126.1			
21	21	TRUERBLOOD		26	1429	1431	25:00	25:00		0.00		.035	126.8			
21	21	TRUERBLOOD		26	1429	1431	25:00	25:00		0.00		.022	14.97			
21	21	TRUERBLOOD		26	1429	1431	25:00	25:00		0.00		.0133	2.			
RUN: 1																
24	24	SERPOLAY		25	1340	1355	25:70	26:70		1.00	.77	.03	400			
24	24	SERPOLAY		25	1407	1415	26:10	28:70		3.00	.77	.15	530			
24	24	SERPOLAY		25	1445	1447	27:20	31:00		3.80	.77	.54	1000			
RUN: 1																
25	25	BORYS		3	1341	1348	25:50	26:50		1.00	10.6	.038	820		.33	VOL SAMP X 10**5
25	25	BORYS		3	1354	1358	26:00	28:30		1.20	1.6	.076	1051			
25	25	BORYS		3	1406	1412	27:00	28:00		3.20	1.6	.13	1732			
25	25	BORYS		3	1418	1422	27:10	28:00		3.50	1.6	.79	1697			
25	25	BORYS		3	1438	1438	20:00	23:00		5.20	9.18	1.2	3206			
RUN: 1																
26	26	HINDMAN		6	1336	1349										INSTRUMENT DOWN REPAIRING CHAMBER
RUN: 1																
27	27	OMTAK		19	1358	1358					11		.716			103% R.M. NOMINAL
27	27	OMTAK		19	1358	1358					11		.1.1			114% R.M. NOMINAL

INSTRUMENT COMPARISON



HAZE CHAMBER RESULTS

INSTRUMENT COMPARISON



SDC/CFD RESULTS

EXPERIMENT # 15

PURPOSE

MONODISPERSE AEROSOL WITH SIZE USEFUL TO BOTH HAZE AND CCN INSTRUMENTS,
I.E., $D \text{ APPROX. } \approx 0.092 \text{ UM.}$

DESCRIPTION OF EXPERIMENT

DATE DATE TIME START END
10 OCT. 1980 1455 1630

NUCLEI TYPE
(NH₄)₂ SO₄

GENERATION METHOD

DRI ATOMIZER, 50 PSI, PUMP 2.

SIZE DISTRIBUTION SHAPING

EC, 2780 V., MONODISPERSE FLOWS 6 L/MIN.

REMARKS

DILUTION FLOW 390 L/MIN.

WEATHER SYNOPSIS

NONE

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS		OUTSIDE CONDITIONS					
AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG TEMP (C)	AVG RH (%)	AVG PRESS (MR)	AVG WS (M/S)	AVG WD (DEG)
27.0	21.0	850	NA	NA	NA	NA	NA

PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

EXPERIMENT #: 15

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	COLD	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
								HOT (C)	(C)							
RUN: 1																
2	2	ROGERS, DAVE	DAVE	22	1544	1545	21.70	23.80	2.10		.17		1138			
		ROGERS, DAVE	DAVE	22	1547	1549	21.60	24.00	2.40		.23		1279			
RUN: 1																
5	5	KITCHEN		12	1555	1555	20.50	24.20	3.70		.55	294				
5	5	KITCHEN		12	1556	1556	21.20	24.20	3.00		.36	114				
5	5	KITCHEN		12	1557	1557	22.40	24.20	1.80		.13	457				
5	5	KITCHEN		12	1559	1559	18.60	24.20	5.60		1.3	268				
5	5	KITCHEN		12	1600	1600	19.40	24.20	4.80		1					
RUN: 1																
9	9	GAGIN		5	1512		22.70	24.40	5.70		1.25	1900				
9	9	GAGIN		5	1514		24.30	28.20	2.90		.59	1070				
9	9	GAGIN		5	1517		25.30	28.20	2.90		.30	1060				
9	9	GAGIN		5	1520		26.30	28.10	1.80		.18	120				
RUN: 2																
9	9	GAGIN		5	1524		22.60	28.20	5.60		1.23	2100				
9	9	GAGIN		5	1525		24.40	28.30	3.90		.18	2400				
9	9	GAGIN		5	1532		26.40	28.30	1.90		.18	140				
RUN: 3																
9	9	GAGIN		5	1536		22.80	28.00	5.20		1.08	2080				
9	9	GAGIN		5	1538		24.40	28.10	3.70		.52	2070				
9	9	GAGIN		5	1542		25.40	28.30	2.90		.31	1000				
RUN: 1																
10	10	LALA		13	1512		23.84	26.38	2.54		.25	1915	2			
10	10	LALA		13	1513		23.37	26.38	2.99		.35	1725	2			
10	10	LALA		13	1515		23.36	26.37	3.01		.35	1870	2			
10	10	LALA		13	1516		22.72	26.35	3.63		.55	1903	5			
10	10	LALA		13	1517		22.72	26.30	3.38		.55	1856	3			
10	10	LALA		13	1518		22.72	26.30	3.38		.55	1856	3			
10	10	LALA		13	1519		21.95	26.12	4.17		.75	1976	5			
10	10	LALA		13	1520		21.70	26.12	4.37		.75	1712	5			
10	10	LALA		13	1521		20.84	25.86	5.02		1	1616	1			
RUN: 2																
10	10	LALA		13	1526		23.92	26.31	2.38		.25	1606	2			
10	10	LALA		13	1527		23.42	26.31	2.89		.35	1841	1			
10	10	LALA		13	1528		23.71	26.31	2.60		.25	1847	1			
10	10	LALA		13	1529		23.71	26.30	2.59		.25	1678	6			
10	10	LALA		13	1530		23.35	26.30	3.95		.07	1611	4			
10	10	LALA		13	1531		23.35	26.30	3.95		.07	1611	4			
10	10	LALA		13	1532		23.35	26.30	3.95		.07	1611	4			
10	10	LALA		13	1533		23.35	26.30	3.95		.07	1611	4			
10	10	LALA		13	1534		23.35	26.30	3.95		.07	1611	4			
10	10	LALA		13	1535		23.35	26.30	3.95		.07	1611	4			
10	10	LALA		13	1536		23.35	26.30	3.95		.07	1611	4			
10	10	LALA		13	1537		23.35	26.30	3.95		.07	1611	4			
10	10	LALA		13	1538		23.35	26.30	3.95		.07	1611	4			
10	10	LALA		13	1539		23.35	26.30	3.95		.07	1611	4			
10	10	LALA		13	1540		23.35	26.30	3.95		.07	1611	4			
10	10	LALA		13	1541		23.35	26.30	3.95		.07	1611	4			
10	10	LALA		13	1542		23.35	26.30	3.95		.07	1611	4			
RUN: 1																
11	11	FITZGERALD		28	1528						.15	501	501			
11	11	FITZGERALD		28	1528						.094	125	125			
11	11	FITZGERALD		28	1528						.034	.16	.16			
11	11	FITZGERALD		28	1528						.025	.018	.018			
11	11	FITZGERALD		28	1528						.014	.010	.010			
RUN: 1																
13	13	AYERS		2	1512		27.63	30.20	2.57		.25	1770				

PHOTOGRAPHIC RECORDS

PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

EXPERIMENT #: 15

MACH NO.	DUCT NO.	NAME	OBSERVER	SAMPLE START	SAMPLE END	COLD	HOT	PLATE TEMP (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
13	13	AYERS		1513	1514	26.48	30.10	30.10	3.62		5	1755			TAKEN
13	13	AYERS		1515	1516	25.63	30.00	30.00	4.37		.75	1515			ALL S. VALVES ARE
13	13	AYERS		1516	1517	24.94	30.00	30.00	5.06		1.25	1613			APPROXIMATE ONLY.
13	13	AYERS		1518	1519	24.25	29.90	29.90	5.65			1635			
RUN: 2															
13	13	AYERS		1519	1520	24.16	29.80	29.80	5.64		1.25	1815			PHOTOGRAPHIC RECORDS
13	13	AYERS		1520	1521	24.04	29.70	29.70	5.06		.75	1800			TAKEN
13	13	AYERS		1521	1522	25.19	29.60	29.60	4.41			1710			
13	13	AYERS		1522	1523	26.00	29.60	29.60	3.60		.5	1740			
13	13	AYERS		1523	1524	27.97	29.50	29.50	2.53		.25	1493			
13	13	AYERS		1524	1525	27.57	29.50	29.50	1.93		.25	1492			
13	13	AYERS		1525	1526	28.05	29.50	29.50	1.45		.075	121			
RUN: 3															
13	13	AYERS		1608	1609	25.53	30.20	30.20	5.67		1.25	1925			PHOTOGRAPHIC RECORDS
13	13	AYERS		1611	1612	25.28	30.00	30.00	7.72		.75	1525			TAKEN
13	13	AYERS		1614	1615	27.58	29.90	29.90	2.26		.25	1920			
13	13	AYERS		1617	1618	27.95	29.90	29.90	2.50		.25	1440			
13	13	AYERS		1619	1620	28.45	29.90	29.90	1.47		.075	1490			
13	13	AYERS		1620	1621	28.03	30.10	30.10	3.07		1	1850			
RUN: 1															
14	14	HUDSON		1522	1523	15.55	22.00	22.00	18.6	.0186	.21	188			PHOTOGRAPHIC RECORDS
14	14	HUDSON		1522	1523	15.55	22.00	22.00	18.6	.0186	.11	187			TAKEN
14	14	HUDSON		1522	1523	15.55	22.00	22.00	18.6	.0186	.11	187			
14	14	HUDSON		1522	1523	15.55	22.00	22.00	18.6	.0186	.029	52			
14	14	HUDSON		1522	1523	15.55	22.00	22.00	18.6	.0186	.21	188			
14	14	HUDSON		1522	1523	15.55	22.00	22.00	18.6	.0186	.11	187			
14	14	HUDSON		1522	1523	15.55	22.00	22.00	18.6	.0186	.11	187			
14	14	HUDSON		1522	1523	15.55	22.00	22.00	18.6	.0186	.11	187			
14	14	HUDSON		1522	1523	15.55	22.00	22.00	18.6	.0186	.029	52			
14	14	HUDSON		1522	1523	15.55	22.00	22.00	18.6	.0186	.029	52			
RUN: 1															
15	15	ROGERS	FRED	1513	1514	18.36	25.06	25.06	6.70		1.86	1365			REDUCED MAIN FLOW
15	15	ROGERS	FRED	1523	1526	19.19	24.43	24.43	6.01		1.46	1297			TO ABOUT 100 CMTS.
15	15	ROGERS	FRED	1523	1526	19.87	24.81	24.81	5.33		1.15	1458			DROP SIZE PLATEAUS.
15	15	ROGERS	FRED	1534	1539	20.63	23.81	23.81	3.94		.63	1147			BACKGROUNDS TWO PLS.
15	15	ROGERS	FRED	1542	1547	20.99	23.50	23.50	3.25		.25	1150			DROP SIZE PLATEAU
15	15	ROGERS	FRED	1542	1547	21.35	22.88	22.88	1.89		.14	1162			WORKING AGAIN.
15	15	ROGERS	FRED	1548	1549	21.12	22.72	22.72	1.52		.07	1146			AVERAGING LAST 4
15	15	ROGERS	FRED	1548	1549	21.12	22.72	22.72	1.52		.1	1166			COUNTS PER RUN
15	15	ROGERS	FRED	1548	1549	21.12	22.72	22.72	1.52		.1	1166			FOR WHOLE RUN
RUN: 1															
16	16	HUDSON		1510	1511	21.35	26.00	26.00	7.5	.0018	.22	133			
16	16	HUDSON		1510	1511	21.35	26.00	26.00	7.5	.0018	.22	133			
16	16	HUDSON		1510	1511	21.35	26.00	26.00	7.5	.0018	.22	133			
16	16	HUDSON		1510	1511	21.35	26.00	26.00	7.5	.0018	.22	133			
16	16	HUDSON		1510	1511	21.35	26.00	26.00	7.5	.0018	.22	133			
16	16	HUDSON		1510	1511	21.35	26.00	26.00	7.5	.0018	.22	133			
16	16	HUDSON		1510	1511	21.35	26.00	26.00	7.5	.0018	.22	133			
16	16	HUDSON		1510	1511	21.35	26.00	26.00	7.5	.0018	.22	133			
16	16	HUDSON		1510	1511	21.35	26.00	26.00	7.5	.0018	.22	133			
16	16	HUDSON		1510	1511	21.35	26.00	26.00	7.5	.0018	.22	133			
RUN: 2															
16	16	HUDSON		1542	1545	21.35	26.00	26.00	7.5	.0018	.22	133			
16	16	HUDSON		1542	1545	21.35	26.00	26.00	7.5	.0018	.22	133			
16	16	HUDSON		1542	1545	21.35	26.00	26.00	7.5	.0018	.22	133			

EXPERIMENT #: 15

NUCLEI MEASUREMENTS

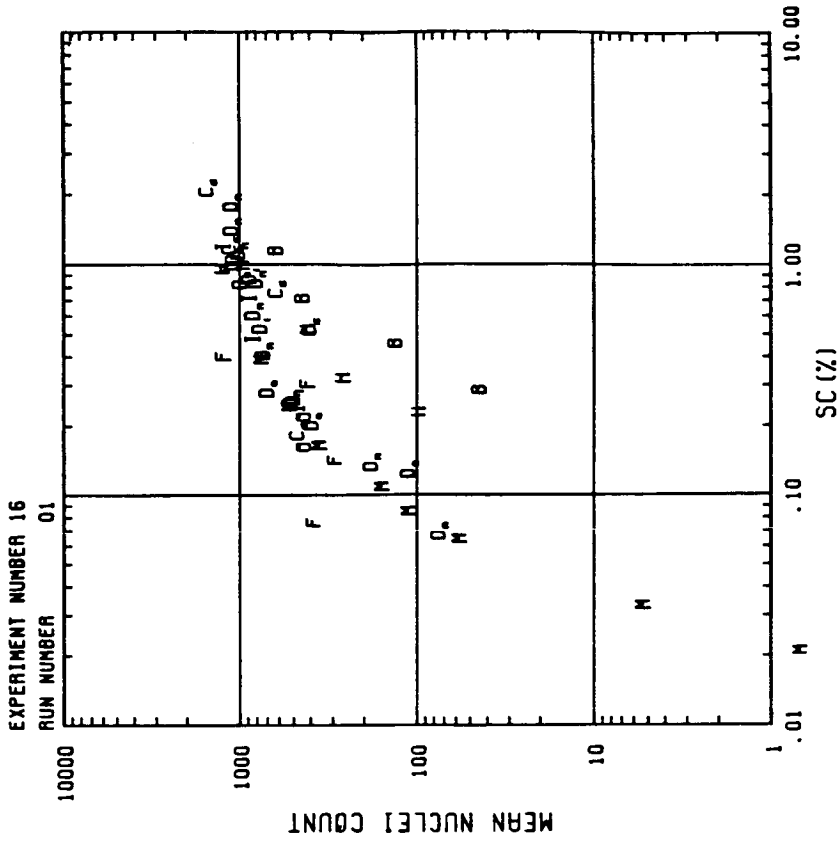
PROCESSING DATE: 81/09/26

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	COLD	HOT (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
16	16	HUDSON		9	1604	1606	21.35	22.10	4.75	.0012	.92	1573			
16	16	HUDSON		9	1604	1606	22.20	22.85	2.65	.0012	.28	1355			
16	16	HUDSON		9	1619	1623	21.70	22.10	4.75	.0024	.92	1575			
16	16	HUDSON		9	1619	1623	22.10	22.85	4.65	.0024	.28	1362			
RUN: 1															
17	17	WOJCIECHOWSKI		27	1515	1516						1253	1253		
17	17	WOJCIECHOWSKI		27	1527	1528						1259	1259		
17	17	WOJCIECHOWSKI		27	1527	1528						1259	1259		
17	17	WOJCIECHOWSKI		27	1527	1528						1259	1259		
RUN: 1															
18	18	HUDSON		9	1518	1519	22.05	22.65	5.00	.0018	.1	1450			
18	18	HUDSON		9	1526	1529	22.40	22.15	2.25	.0018	.43	1567			
18	18	HUDSON		9	1545	1545	23.10	22.90	2.20	.0018	.21	1448			
18	18	HUDSON		9	1602	1623	23.10	22.50	2.10	.0024	.14	155			
RUN: 1															
21	21	TRUEBLOOD		25	1515	1516	20.53	20.00	4.8		.812	183	183		
21	21	TRUEBLOOD		25	1527	1528	21.30	20.00	3.3		.412	182	182		
21	21	TRUEBLOOD		25	1527	1528	21.30	20.00	3.3		.412	182	182		
21	21	TRUEBLOOD		25	1527	1528	21.30	20.00	3.3		.412	182	182		
21	21	TRUEBLOOD		25	1527	1528	21.30	20.00	3.3		.412	182	182		
21	21	TRUEBLOOD		25	1527	1528	21.30	20.00	3.3		.412	182	182		
21	21	TRUEBLOOD		25	1527	1528	21.30	20.00	3.3		.412	182	182		
21	21	TRUEBLOOD		25	1527	1528	21.30	20.00	3.3		.412	182	182		
21	21	TRUEBLOOD		25	1527	1528	21.30	20.00	3.3		.412	182	182		
21	21	TRUEBLOOD		25	1527	1528	21.30	20.00	3.3		.412	182	182		
21	21	TRUEBLOOD		25	1527	1528	21.30	20.00	3.3		.412	182	182		
RUN: 1															
24	24	SERPOLAY		25	1523	1538	27.40	28.40	1.00	.77	.03	150			
24	24	SERPOLAY		25	1556	1602	28.50	30.90	2.90	.77	.15	400			
24	24	SERPOLAY		25	1615	1620	28.50	32.50	4.00	.77	.16	760			
24	24	SERPOLAY		25	1625	1630	28.80	33.40	4.60	.77	.14	1530			
RUN: 1															
25	25	BORYS		33	1520	1528	25.30	26.30	1.00	12.72	.08	601	601		
25	25	BORYS		33	1537	1538	25.00	26.00	1.00	12.72	.08	601	601		
25	25	BORYS		33	1547	1548	25.00	26.00	1.00	12.72	.08	601	601		
25	25	BORYS		33	1547	1548	25.00	26.00	1.00	12.72	.08	601	601		
25	25	BORYS		33	1547	1548	25.00	26.00	1.00	12.72	.08	601	601		
25	25	BORYS		33	1547	1548	25.00	26.00	1.00	12.72	.08	601	601		
25	25	BORYS		33	1547	1548	25.00	26.00	1.00	12.72	.08	601	601		
25	25	BORYS		33	1547	1548	25.00	26.00	1.00	12.72	.08	601	601		
25	25	BORYS		33	1547	1548	25.00	26.00	1.00	12.72	.08	601	601		
RUN: 1															
26	26	HINDMAN		6	1353	1421									

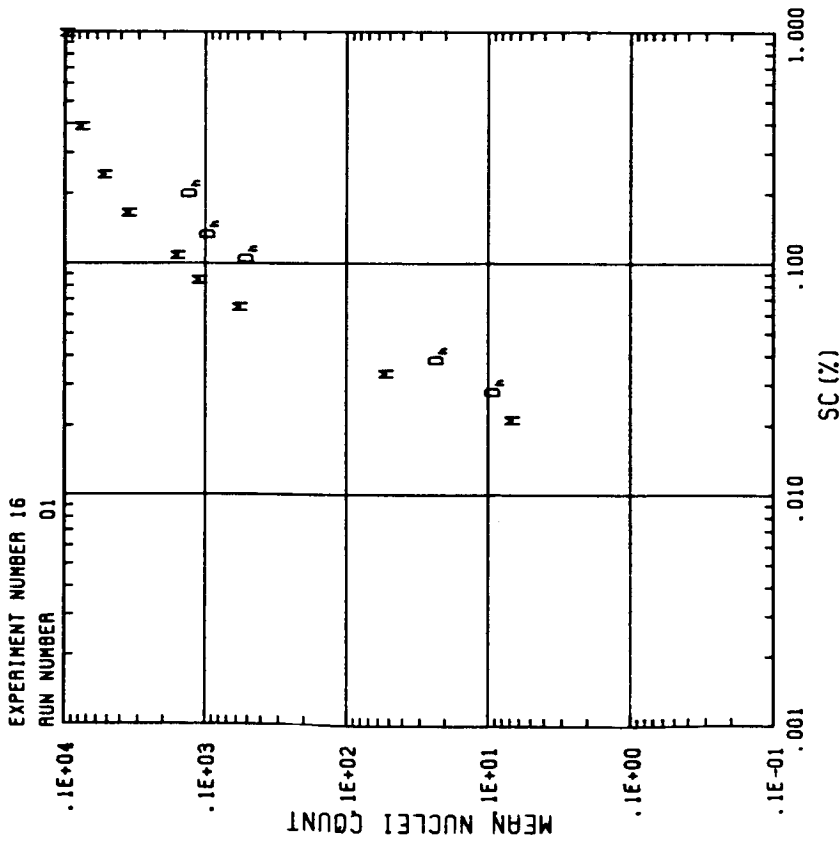
VOL SAMP X 10**5

INSTRUMENT DOWN. REPAIRING CHAMBER.

INSTRUMENT COMPARISON



INSTRUMENT COMPARISON



SDC/CFD RESULTS

HAZE CHAMBER RESULTS

EXPERIMENT # 16

PURPOSE

INSTRUMENT COMPARISON WITH AMBIENT AEROSOL.

DESCRIPTION OF EXPERIMENT

DATE	TIME START	END
10 OCT. 1980	1635	1726

NUCLEI TYPE
 AMBIENT

GENERATION METHOD
 NA

SIZE DISTRIBUTION SHAPING
 NA

REMARKS
 NONE

WEATHER SYNOPSIS
 NONE

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS		OUTSIDE CONDITIONS					
AVG TEMP (C)	AVG RH (%)	AVG PPRESS (MB)	AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG WS (M/S)	AVG WD (DEG)
29.0	20.0	850	NA	NA	NA	4.0	270

EXPERIMENT #: 16

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/09/26

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	SAMPLE END	COLD	HOT (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
2	2	POLITOVICH		20	1656	1657	23.90	26.00	2.10		.17		456		
					1658	1700	23.80	26.20	2.40		.23		456		
5	5	KITCHEN		12	1635	1635	21.20	24.90	3.70		.3	49			
5	5	KITCHEN		12	1637	1637	21.50	24.90	3.40		.48	131			
5	5	KITCHEN		12	1638	1638	20.90	24.90	4.00		.75	456			
5	5	KITCHEN		12	1639	1639	19.80	24.90	5.10		1.2				
5	5	KITCHEN		12	1640	1640	20.60	24.90	4.30		.75	330			
5	5	KITCHEN		12	1641	1641	21.60	24.90	3.30		.48	173			
5	5	KITCHEN		12	1642	1642	22.20	24.90	2.70		.3	34			
5	5	KITCHEN		12	1643	1643	20.30	24.90	4.60		1.25	256			
5	5	KITCHEN		12	1644	1644	19.90	24.90	5.00		1.75	552			
5	5	KITCHEN		12	1645	1645	20.60	24.90	4.30		.48	343			
5	5	KITCHEN		12	1646	1646	21.60	25.00	3.40		.48	125			
9	9	GAGIN		5	1636	1636	23.00	28.10	5.10	1.2	.1	1310			
9	9	GAGIN		5	1638	1638	24.30	28.10	3.80	1.2	.55	430			
9	9	GAGIN		5	1642	1642	25.10	28.10	3.00	1.2	.24	270			
9	9	GAGIN		5	1650	1650	26.00	28.50	2.50			100			
9	9	GAGIN		5	1649	1649	22.70	28.10	5.40	1.2	1.14	1550			
9	9	GAGIN		5	1652	1652	25.00	28.10	3.00	1.2	.64	400			
9	9	GAGIN		5	1654	1654	25.10	28.10	3.00	1.2	.16	200			
9	9	GAGIN		5	1701	1701	22.90	27.90	5.00	1.2	.98	1150			
9	9	GAGIN		5	1704	1704	24.40	28.30	3.90	1.2	.55	300			
9	9	GAGIN		5	1708	1708	26.10	28.30	2.20		.19	150			
13	13	AYERS		2	1635	1636	28.01	30.50	2.49		.25	488			
13	13	AYERS		2	1640	1641	26.82	30.40	3.58		.5	87			
13	13	AYERS		2	1643	1643	25.98	30.40	4.42		.75	915			
13	13	AYERS		2	1644	1645	24.75	30.40	5.65		1.25	1120			
13	13	AYERS		2	1646	1647	24.64	30.30	5.66		1.25	100			
13	13	AYERS		2	1648	1649	25.13	30.30	5.07		.75	1020			
13	13	AYERS		2	1650	1651	25.68	30.20	4.52		.25	800			
13	13	AYERS		2	1652	1653	27.64	30.20	2.54		.25	615			
13	13	AYERS		2	1654	1655	27.66	30.20	2.54		.25	615			
14	14	HUDSON		9	1642	1702				.0148	.21	124			55555555
14	14	HUDSON		9	1643	1702				.0143	.11	973			88888888
14	14	HUDSON		9	1645	1702				.0143	.04	738			33333333
14	14	HUDSON		9	1646	1702				.0143	.029	102			55555555
14	14	HUDSON		9	1703	1720				.0143	.21	124			88888888

EXPERIMENT #: 16

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/09/26

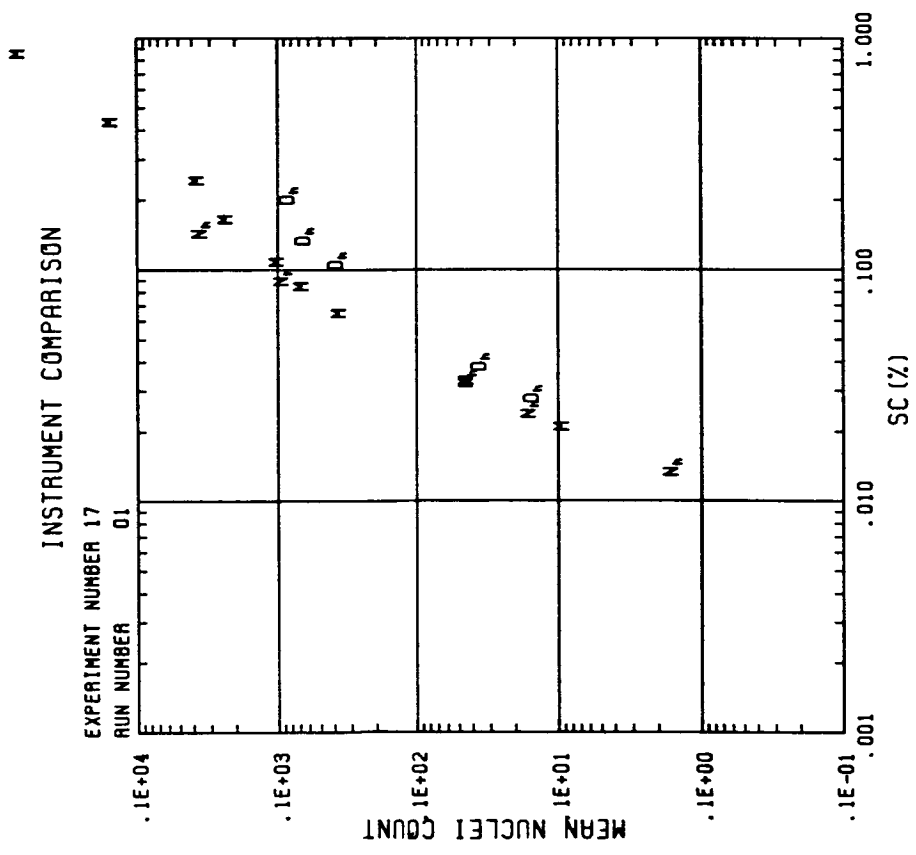
MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	SAMPLE END	COLD	HOT	PLATE TEMP	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
14	14	HUDSON		9	1703	1728					.0143	.11	112			F=38.5
14	14	HUDSON		9	1703	1728					.0143	.04	6187			F=38.5
14	14	HUDSON		9	1703	1728					.0143	.029	1114			F=38.5
RUN: 1																
15	15	ROGERS	FRED	23	1645	1646	18.36	25.06	6.70			1.46	1155			MAIN FLO ABT 1300
15	15	ROGERS	FRED	23	1653	1654	18.74	24.47	6.01			1.46	1155			CNTS. BACKGROUND 10.
15	15	ROGERS	FRED	23	1657	1658	19.11	24.44	5.33			1.15	1059			
15	15	ROGERS	FRED	23	1702	1703	19.87	23.43	4.64			.87	870			AEROSOL CHANGING
15	15	ROGERS	FRED	23	1706	1707	20.25	23.50	3.25			.63	773			AS FCN. TIME ?
15	15	ROGERS	FRED	23	1710	1711	20.63	23.10	2.56			.265	540			
15	15	ROGERS	FRED	23	1714	1715	20.99	22.84	1.89			.14	190			MAIN FLO 1287
15	15	ROGERS	FRED	23	1718	1719	21.35	22.54	1.21			.07	178			
RUN: 1																
16	16	HUDSON		9	1636	1641	21.40	26.10	4.70		.003	.9	936			
16	16	HUDSON		9	1636	1641	21.70	25.80	3.70		.003	.27	729			
16	16	HUDSON		9	1653	1655	22.17	25.10	2.90		.0012	.57	884			
16	16	HUDSON		9	1653	1655	22.50	24.80	2.70		.0012	.27	798			
16	16	HUDSON		9	1710	1713	22.70	24.40	2.90		.0012	.57	1032			
16	16	HUDSON		9	1710	1713	22.80	24.80	2.60		.0018	.27	858			
RUN: 2																
16	16	HUDSON		9	1720	1721	21.40	26.10	4.70		.0012	.9	1072			
16	16	HUDSON		9	1720	1721	21.70	25.80	3.70		.0012	.57	944			
16	16	HUDSON		9	1726	1727	22.17	26.10	2.60		.0012	.27	1155			
16	16	HUDSON		9	1726	1727	22.40	25.40	2.60		.0012	.27	1155			
16	16	HUDSON		9	1726	1727	22.80	24.80	2.60		.0012	.27	657			
RUN: 1																
18	18	HUDSON		9	1636	1641	23.65	25.45	1.80		.003	.13	116			
18	18	HUDSON		9	1653	1655	23.40	25.70	2.30		.0012	.29	471			
18	18	HUDSON		9	1720	1721	22.50	29.90	2.70		.0012	.16	731			
18	18	HUDSON		9	1726	1727	22.80	27.20	5.60		.0012	.11	1037			
RUN: 1																
21	21	TRUEBLOOD		26	1643	1645	19.90	25.00	5.10			1.04	1003			
21	21	TRUEBLOOD		26	1659	1657	22.48	25.00	2.52			.41	1792			
21	21	TRUEBLOOD		26	1708	1710	22.50	25.00	0.00			.54	546			
21	21	TRUEBLOOD		26	1708	1710	22.50	25.00	0.00			.14	366			
21	21	TRUEBLOOD		26	1708	1710	22.50	25.00	0.00			.14	164			
21	21	TRUEBLOOD		26	1718	1710	22.50	25.00	0.00			.08	115			
21	21	TRUEBLOOD		26	1718	1721	22.50	25.00	0.00			.035	5			
21	21	TRUEBLOOD		26	1718	1721	22.50	25.00	0.00			.023	.0			
RUN: 1																
24	24	SERPOLAY		25	1640	1650	29.20	32.20	3.00		.77	.32	430			IMPORTANT FLUCTUATIO
24	24	SERPOLAY		25	1656	1659	29.50	31.60	2.20		.77	.15	300			LOW LEVEL AT ABOUT
24	24	SERPOLAY		25	1705	1710	29.70	33.10	3.40		.77	.42	1280			1647
RUN: 1																
25	25	BORYS		3	1636	1638	18.40	25.50	7.10		4.24	2.15	1591		1.16	VOL SAMP X 10**5

EXPERIMENT #: 16

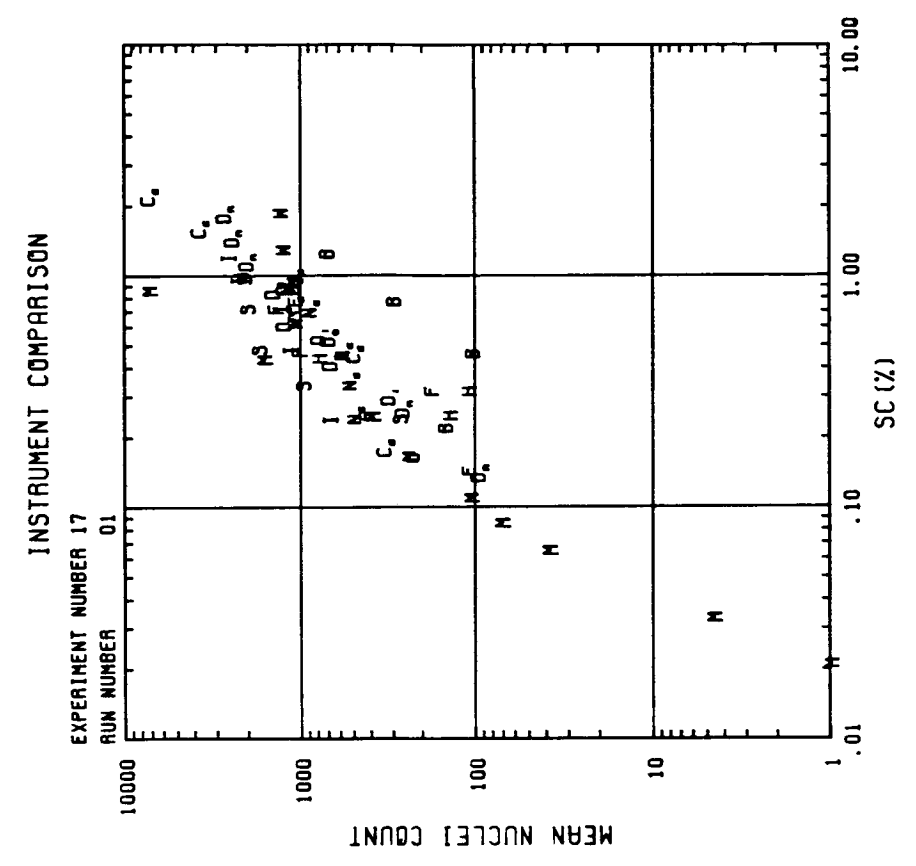
NUCLEI MEASUREMENTS

PROCESSING DATE: 81/09/26

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	COLD	HOT (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST R	REMARKS
25	25	BORYS		3	1643	1645	20.10	25.60	5.50	8.48	1.22	1185	1185		
25	25	BORYS		3	1650	1656	21.20	25.70	4.50	121.2	.54	648	648		
25	25	BORYS		3	1701	1709	22.50	26.00	3.50	21.2	.19	417	417		
25	25	BORYS		3	1717	1725	24.30	26.60	2.30	21.2		495	495		



HAZE CHAMBER RESULTS



SDC/CFD RESULTS

PURPOSE

INSTRUMENT COMPARISON WITH AMBIENT AEROSOL.

DESCRIPTION OF EXPERIMENT

DATE 10 OCT 1980 0823 0957
TIME START END

NUCLEI TYPE
AMBIENT

GENERATION METHOD

NA

SIZE DISTRIBUTION SHAPING

NA

REMARKS

20% R.H.

WEATHER SYNOPSIS

NONE

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS		OUTSIDE CONDITIONS						
AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG WS (M/S)	AVG WD (DEG)	VAR
26.0	21.0					<1.0		

EXPERIMENT #: 17

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/09/26

MACH NO.	DUCT NO.	OBSERVER	NO.	SAMPLE START	SAMPLE END	COLD	HOT	PLATE TEMP	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
RUN: 1															
2	2	ROGERS, DAVE	22	0918	0919	21.70	24.10	24.10	2.40		.17		235		
				0920	0921	21.70	24.10	24.10	2.40		.23		206		
RUN: 1															
5	5	KITCHEN	12	0831	0831	19.20	22.60	22.60	3.40		.48	88			
5	5	KITCHEN	12	0832	0833	18.30	22.60	22.60	4.30		1.3	277			
5	5	KITCHEN	12	0833	0834	17.30	22.60	22.60	5.30		1.3	333			
5	5	KITCHEN	12	0834	0835	18.40	22.70	22.70	4.30		.8	333			
5	5	KITCHEN	12	0835	0836	19.40	22.80	22.80	3.40		.48	125		1.30	
5	5	KITCHEN	12	0836	0836	20.40	22.80	22.80	2.40		.23	194			
RUN: 2															
5	5	KITCHEN	12	0922	0922	22.00	25.40	25.40	3.40		.48	229			
5	5	KITCHEN	12	0923	0923	21.10	25.40	25.40	4.30		1.3	391			
5	5	KITCHEN	12	0924	0925	22.10	25.50	25.50	5.40		.48	188			
RUN: 1															
9	9	GAGIN	5	0837	0837	23.00	27.10	27.10	5.10	1.2	1.03	240			
9	9	GAGIN	5	0840	0840	24.30	27.20	27.20	2.90	1.2	.46	800			
9	9	GAGIN	5	0844	0844	25.00	27.50	27.50	2.50	1.2	.26	110			
RUN: 2															
9	9	GAGIN	5	0847	0847	21.80	27.10	27.10	5.30	1.2	1.1	1530			
9	9	GAGIN	5	0849	0849	23.60	27.30	27.30	3.70	1.2	.52	1160			
9	9	GAGIN	5	0853	0853	24.70	27.30	27.30	2.60	1.2	.26	1190			
RUN: 3															
9	9	GAGIN	5	0901	0901	22.20	27.20	27.20	5.00	1.2	.99	1350			
9	9	GAGIN	5	0907	0907	24.70	27.70	27.70	3.10	1.2	.48	800			
9	9	GAGIN	5	0910	0910	25.80	27.90	27.90	2.10	1.2	.17	120			
RUN: 4															
9	9	GAGIN	5	0917	0917	22.30	27.70	27.70	5.40	1.2	1.13	1270			
9	9	GAGIN	5	0920	0920	24.80	27.90	27.90	3.10	1.2	.57	320			
9	9	GAGIN	5	0924	0924	24.80	27.90	27.90	3.10	1.2	.57	320			
RUN: 1															
10	10	LALA	13	0836	0836	21.09	23.50	23.50	2.49	.4	.25	268.9			
10	10	LALA	13	0838	0838	20.58	23.48	23.48	2.94	.4	.25	2870.9			
10	10	LALA	13	0840	0840	20.02	23.52	23.52	3.94	.4	.25	870.1			
10	10	LALA	13	0841	0841	20.02	23.52	23.52	3.50	.4	.25	1120.1			
10	10	LALA	13	0842	0842	19.29	23.52	23.52	4.28	.4	.25	1738.9			
10	10	LALA	13	0843	0843	19.29	23.51	23.51	4.28	.4	.25	2157.9			
10	10	LALA	13	0844	0844	18.51	23.46	23.46	4.92	.4	.25	1928.9			
10	10	LALA	13	0845	0845	18.51	23.46	23.46	4.92	.4	.25	2329.9			
RUN: 2															
10	10	LALA	13	0846	0846	21.32	23.82	23.82	3.50	.4	.25	365.9			
10	10	LALA	13	0847	0847	20.32	23.81	23.81	4.92	.4	.25	271.8			
10	10	LALA	13	0849	0849	20.87	23.82	23.82	3.51	.4	.25	349.8			
10	10	LALA	13	0850	0850	20.90	23.81	23.81	3.51	.4	.25	94.8			

VERY UNSTABLE AEROSO

EXPERIMENT #: 17

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/09/26

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	COLD	HOT	PLATE TEMP (C)	BELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
10	10	LALA		13	0851		20.31	23.83	3.52	.4	.5	.75	1602.1			
10	10	LALA		13	0852		19.53	23.83	4.29	.4	.4	.75	2084.2			
10	10	LALA		13	0853		19.53	23.82	4.29	.4	.4	.75	1407.7			
10	10	LALA		13	0854		18.84	23.78	4.94	.4	.4	.75	1417.1			
10	10	LALA		13	0855		18.85	23.79	4.94	.4	.4	.75	1395.7			VERY UNSTABLE AEROSO
RUN: 1																
11	11	FITZGERALD		28	0845	0857						.15	378	378		
11	11	FITZGERALD		28	0845	0857						.094	109	109		
11	11	FITZGERALD		28	0845	0857						.026	4.7	4.7		
11	11	FITZGERALD		28	0845	0857						.014	1.7	1.7		
RUN: 2																
11	11	FITZGERALD		28	0910	0927						.15	307	307		
11	11	FITZGERALD		28	0910	0927						.094	96	96		
11	11	FITZGERALD		28	0910	0927						.034	6.8	6.8		
11	11	FITZGERALD		28	0910	0927						.014	2.4	2.4		
RUN: 1																
13	13	AYERS		2	0839	0840	21.89	24.30	2.41		.25		690			
13	13	AYERS		2	0841	0842	20.91	24.40	3.49		.15		1170			
13	13	AYERS		2	0842	0843	20.53	24.40	4.20		.1		1290			
13	13	AYERS		2	0846	0847	18.96	24.40	5.44		1.25		2325			
RUN: 1																
14	14	HUDSON		9	0911	0933					.014	.21	94.1			F=40
14	14	HUDSON		9	0911	0933					.014	.11	73.7			F=40
14	14	HUDSON		9	0911	0933					.014	.11	43.2			F=40
14	14	HUDSON		9	0911	0933					.014	.06	4.22			F=40
14	14	HUDSON		9	0911	0933					.014	.06	1.86			F=40
14	14	HUDSON		9	0914	0937					.014	.11	65.1			F=40
14	14	HUDSON		9	0934	0957					.014	.11	98.2			F=40
14	14	HUDSON		9	0934	0957					.014	.029	3.2			F=40
RUN: 1																
15	15	ROGERS, FRED		23	0925	0926	18.36	25.06	6.70		1.86		2807			FIRST POINT NEEDED
15	15	ROGERS, FRED		23	0928	0929	18.74	24.75	6.01		1.46		2548			MAIN FLO PE ADJUST.
15	15	ROGERS, FRED		23	0931	0936	19.49	24.44	5.33		1.187		2097			BACKGROUND LESS
15	15	ROGERS, FRED		23	0935	0940	19.87	24.81	4.64		.63		1488			THAN 10 PER CC.
15	15	ROGERS, FRED		23	0939	0944	19.87	23.50	3.94		.425		1298			ROYCO CAL OK
15	15	ROGERS, FRED		23	0943	0948	20.25	23.81	3.25		.255		670			SIGNIFICANT AEROSOL
15	15	ROGERS, FRED		23	0947	0952	20.99	22.88	1.89		.214		298			FLUCTUATIONS.
RUN: 1																
16	16	HUDSON		9	0931	0938	21.35	26.05	4.70		.042		1281			
16	16	HUDSON		9	0931	0938	21.70	25.40	3.70		.042		814			
16	16	HUDSON		9	0931	0938	22.20	24.70	2.50		.042		328			
RUN: 1																
17	17	WOJCIECHOWSKI		27	0905	0907					.25		508			
17	17	WOJCIECHOWSKI		27	0905	0907					.25		540			
17	17	WOJCIECHOWSKI		27	0905	0907					.27		592			

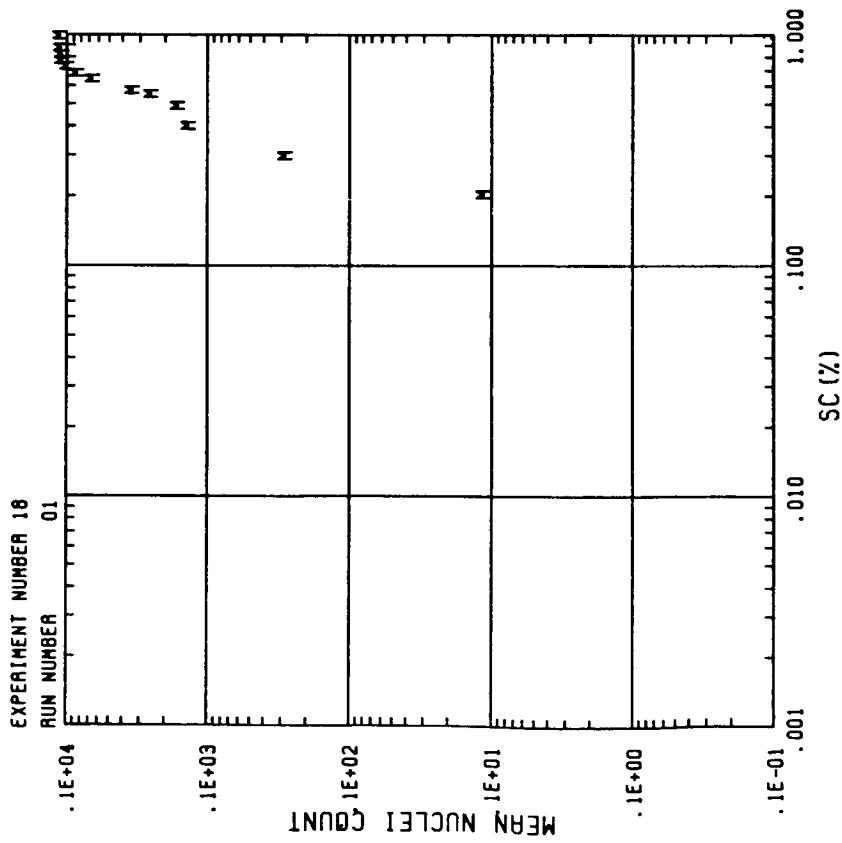
PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

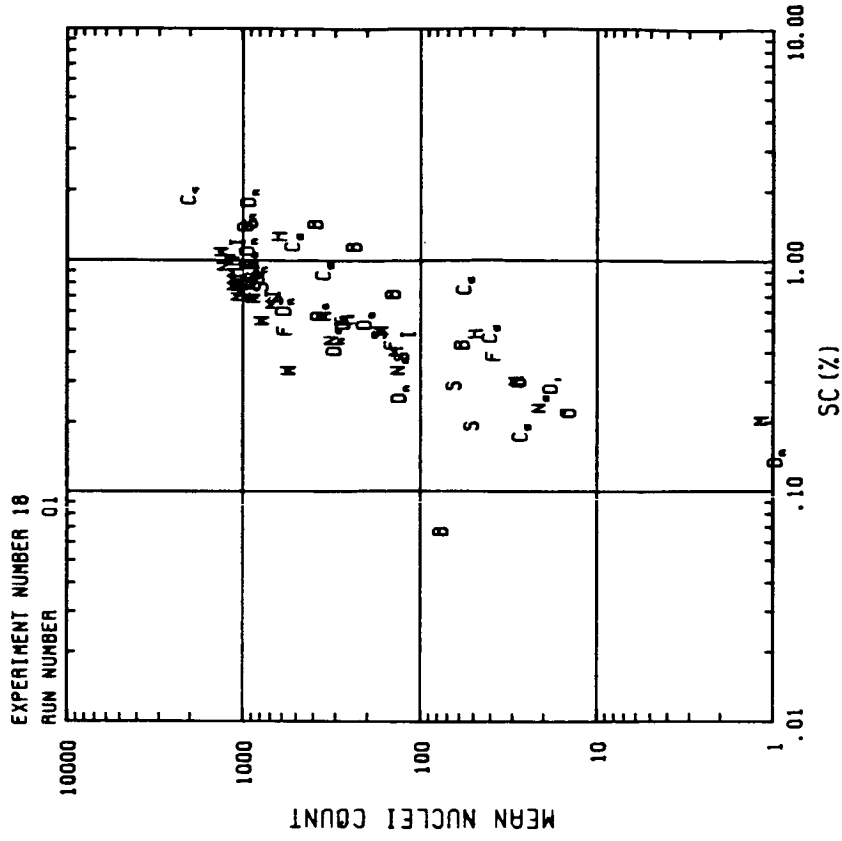
EXPERIMENT #: 17

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE TIME		COLD	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SA (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
					START	END		HOT (C)	HOT (C)							
17	17	WOJCIECHOWSKI	WOJCIECHOWSKI	27	0905	0907					.72	925	925			
17	17	WOJCIECHOWSKI	WOJCIECHOWSKI	27	0905	0907					.98	1132	1132			
RUN: 1																
18	18	HUDSON		9	0921	0924	22.10	26.80	4.70	.0018	.9	1146				
18	18	HUDSON		9	0931	0938	22.65	26.30	3.65	.0042	.54	734				
18	18	HUDSON		9	0951	0955	22.65	26.30	3.65	.0024	.54	715				
RUN: 1																
20	20	DOMONKOS		4	0925	0930					.65	1095				
20	20	DOMONKOS		4	0925	0930					.93	1210				
20	20	DOMONKOS		4	0925	0930					1.35	1293				
20	20	DOMONKOS		4	0925	0930					1.95	1340				
RUN: 1																
21	21	TRUEBLOOD		26	0831	0832	20.25	25.00	4.75		.92	7422				
21	21	TRUEBLOOD		26	0837	0837	20.25	25.00	2.30		.256	1295				
21	21	TRUEBLOOD		26	0849	0849	22.00	25.00	2.30		.273	1295				
21	21	TRUEBLOOD		26	0857	0858	22.50	25.00	0.00		.114	248.8				
21	21	TRUEBLOOD		26	0857	0858	22.50	25.00	0.00		.114	248.8				
21	21	TRUEBLOOD		26	0857	0858	22.50	25.00	0.00		.114	248.8				
21	21	TRUEBLOOD		26	0857	0858	22.50	25.00	0.00		.114	248.8				
21	21	TRUEBLOOD		26	0906	0907	22.50	25.00	0.00		.335	1072.4				
21	21	TRUEBLOOD		26	0906	0907	22.50	25.00	0.00		.335	1072.4				
21	21	TRUEBLOOD		26	0906	0907	22.50	25.00	0.00		.335	1072.4				
21	21	TRUEBLOOD		26	0906	0907	22.50	25.00	0.00		.335	1072.4				
RUN: 1																
24	24	SERPOLAY		25	0830	0831	24.20	26.70	2.50		.77	430				IMPORTANT FLUCTUATIO
24	24	SERPOLAY		25	0845	0852	24.20	26.70	3.60		.77	1050				TILL 0905
24	24	SERPOLAY		25	0901	0920	25.40	29.90	4.50		.77	1430				THEN STABLE.
24	24	SERPOLAY		25	0915	0920	26.10	30.80	4.70		.77	1100				
24	24	SERPOLAY		25	0937	0953	26.50	29.80	3.00		.77	1100				
24	24	SERPOLAY		25	0950	0953	27.10	29.10	2.00		.77	110				
RUN: 1																
25	25	BORIS		3	0830	0831	16.40	23.70	7.30		2.6	7592				SAMP VOL X10**5
25	25	BORIS		3	0835	0837	18.00	23.50	5.50		1.6	3890				MEAS. 1 OFF SCALE
25	25	BORIS		3	0842	0844	19.50	23.90	4.40		1.75	1132				
25	25	BORIS		3	0846	0847	20.70	24.10	3.40		.46	1513				
25	25	BORIS		3	0901	0907	22.40	24.60	2.20		.18	348				
RUN: 1																
27	27	OHTAKE		19	0839	0839					1.1	4.01				101% NOMINAL
27	27	OHTAKE		19	0839	0839					1.1	5.93				105% NOMINAL

INSTRUMENT COMPARISON



INSTRUMENT COMPARISON



80/10/13

EXPERIMENT # 18

PURPOSE

INSTRUMENT COMPARISON WITH SMALL MONODISPERSE CCN.

DESCRIPTION OF EXPERIMENT

DATE 13 OCT 1980 TIME START 1010 END 1130

NUCLEI TYPE (NH4)2 SO4, MONODISPERSE, D=0.04 UM.

GENERATION METHOD

DRI ATOMIZER, 70 PSI, 1.77 G/L SOLUTION.

SIZE DISTRIBUTION SHAPING

EC, 634V, MONODISPERSE, FLOW RATE = 5.8 L/MIN. DILUTION FLOW 570 L/MIN.

REMARKS

NONE

WEATHER SYNOPSIS

NONE

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS		OUTSIDE CONDITIONS					
AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG WS (M/S)	AVG WD (DEG)
26.0	21.0	848					

EXPERIMENT #: 18

NUCLEI MEASUREMENTS

PROCESSING DATE: 01/09/26

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	SAMPLE END	COLD	PLATE TEMP HOT (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	ORBS N	EST K	REMARKS
RUN: 1															
2	2	ROGERS	DAVE	22	1104	1105	23.40	25.50	2.10		.17				
2	2	ROGERS	DAVE	22	1107	1108	23.40	26.90	3.40		.23				
2	2	ROGERS	DAVE	22	1113	1114	23.40	26.20	2.80		.31				LESS THAN 6 PER CC.
RUN: 1															
5	5	KITCHEN		12	1030	1030	25.50	26.80	1.40		.07	79			
5	5	KITCHEN		12	1031	1031	22.50	26.90	4.40		.15	68			
5	5	KITCHEN		12	1032	1033	21.60	26.90	5.30		.15	183			
5	5	KITCHEN		12	1034	1034	20.80	26.90	6.10		.15	243			
RUN: 2															
5	5	KITCHEN		12	1036	1036	25.10	27.00	1.90		.14	81			
5	5	KITCHEN		12	1037	1037	22.00	27.00	4.70		.15	57			
5	5	KITCHEN		12	1038	1039	20.50	27.00	6.10		.15	47			
5	5	KITCHEN		12	1040	1040	19.50	27.00	7.50		.20	71			
5	5	KITCHEN		12	1041	1041	22.80	27.00	4.20		.14	139			
5	5	KITCHEN		12	1042	1042	23.80	27.10	3.30		.14	51			
5	5	KITCHEN		12	1043	1043	23.50	27.10	1.90		.14	139			
RUN: 1															
9	9	GAGIN		5	1027	1027	22.10	27.90	5.80		1.35	640			
9	9	GAGIN		5	1029	1029	24.10	27.90	3.80		.36	50			
9	9	GAGIN		5	1032	1032	25.10	28.10	3.00		.16	0			
9	9	GAGIN		5	1035	1035	26.00	28.10	2.10			0			
RUN: 2															
9	9	GAGIN		5	1039	1039	22.40	27.80	5.40		1.13	470			
9	9	GAGIN		5	1041	1041	24.30	27.90	3.80		.56	60			
9	9	GAGIN		5	1045	1045	25.30	28.00	2.70		.25	0			
9	9	GAGIN		5	1048	1048	25.90	28.00	2.10		.16	0			
RUN: 3															
9	9	GAGIN		5	1052	1052	22.40	27.90	5.50		1.15	570			
9	9	GAGIN		5	1055	1055	24.50	28.00	3.50		.46	50			
9	9	GAGIN		5	1058	1058	25.00	28.00	2.90		.31	0			
9	9	GAGIN		5	1100	1100	26.20	28.00	1.80		.18	0			
RUN: 1															
10	10	LALA		13	1020	1020	24.01	26.90	2.70		.23	52			
10	10	LALA		13	1021	1021	23.15	26.90	3.75		.45	66			
10	10	LALA		13	1022	1022	22.13	26.90	4.76		.67	131			
10	10	LALA		13	1023	1023	21.33	26.90	5.57		.89	382			
10	10	LALA		13	1023	1023	21.93	26.90	4.97		.89	674			
10	10	LALA		13	1023	1023	20.98	26.90	5.91		1.15	819			
10	10	LALA		13	1023	1023	19.86	26.90	6.12		1.15	911			
10	10	LALA		13	1023	1023	20.98	26.90	5.91		1.15	945			
RUN: 2															
10	10	LALA		13	1035	1035	23.57	26.90	3.70		.23	57			
10	10	LALA		13	1036	1036	23.13	26.90	4.77		.45	80			
10	10	LALA		13	1037	1037	22.99	26.90	4.91		.67	237			
10	10	LALA		13	1038	1038	21.91	26.90	5.01		.89	444			
10	10	LALA		13	1040	1040	21.56	26.90	5.21		.89	710			

PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

EXPERIMENT #	MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	SAMPLE END	PLATE TEMP		BETA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
								COLD	HOT (C)							
18	10	10	LALA		13	1041	1041	21.22	25.72	4.50	.4	846.7				
	10	10	LALA		13	1042	1042	20.91	25.67	4.76	.4	891.9				
	10	10	LALA		13	1043	1043	20.61	25.62	5.01	.4	929.5				
	10	10	LALA		13	1044	1044	19.47	25.58	6.11	.4	930.4				
RUN: 3																
10	10	10	LALA		13	1120	1120	21.18	26.46	3.88	.4	46.9				
	10	10	LALA		13	1121	1121	21.66	26.45	3.79	.4	85.2				
	10	10	LALA		13	1122	1122	22.27	26.32	3.21	.4	142.2				
	10	10	LALA		13	1123	1123	22.37	26.32	3.22	.4	279.3				
	10	10	LALA		13	1124	1124	22.30	26.32	4.57	.4	75.8				
	10	10	LALA		13	1125	1125	22.05	26.11	4.78	.4	95.2				
	10	10	LALA		13	1126	1126	21.93	26.03	4.57	.4	85.2				
	10	10	LALA		13	1128	1128	19.78	25.90	5.02	.4	937.2				
RUN: 1																
11	11	11	FITZGERALD		28	1030	1045					0	0			
	11	11	FITZGERALD		28	1030	1045					15	0	0		
	11	11	FITZGERALD		28	1030	1045					.04	0	0		
	11	11	FITZGERALD		28	1030	1045					.01	0	0		
RUN: 1																
13	13	13	AYERS		2	1025	1026	25.39	27.90	3.51	.5	110	0			
	13	13	AYERS		2	1026	1027	22.39	27.80	3.56	.5	110	0			
	13	13	AYERS		2	1027	1028	22.54	27.90	4.36	.5	703	0			
	13	13	AYERS		2	1028	1029	22.77	27.80	5.03	.5	945	0			
RUN: 2																
13	13	13	AYERS		2	1035	1036	25.11	27.60	3.49	.5	0	0			
	13	13	AYERS		2	1036	1037	24.09	27.60	3.71	.5	150	0			
	13	13	AYERS		2	1038	1039	22.29	27.60	4.11	.5	120	0			
	13	13	AYERS		2	1040	1041	22.60	27.60	5.00	.5	1295	0			
RUN: 3																
13	13	13	AYERS		2	1041	1042	21.94	27.50	5.36	1.25	1295	0			
	13	13	AYERS		2	1041	1042	25.40	27.90	3.50	.5	50	0			
	13	13	AYERS		2	1041	1042	24.50	27.90	3.70	.5	155	0			
	13	13	AYERS		2	1041	1042	22.72	27.70	4.58	1.25	1200	0			
RUN: 1																
15	15	15	ROGERS		3	1025	1026	18.36	25.05	6.70	1.86	965				BACKGROUND 10 PER CC.
	15	15	ROGERS		3	1026	1027	19.74	24.44	6.53	1.45	1079				AT THIS PNT. STARTED AVGS. LAST
	15	15	ROGERS		3	1027	1028	19.49	24.44	4.44	1.87	860				4 AT THIS PNT. DROP
	15	15	ROGERS		3	1028	1029	20.87	23.65	3.32	1.99	614				SIZE PLATEAU WOR-
	15	15	ROGERS		3	1029	1030	20.63	22.88	3.25	1.89	1				LAST PNT. EST. CNT.
	15	15	ROGERS		3	1030	1031	20.99	22.88	1.89	1.89	1				
	15	15	ROGERS		3	1031	1032	20.99	22.88	1.89	1.89	1				
	15	15	ROGERS		3	1032	1033	20.99	22.88	1.89	1.89	1				
	15	15	ROGERS		3	1033	1034	20.99	22.88	1.89	1.89	1				
	15	15	ROGERS		3	1034	1035	20.99	22.88	1.89	1.89	1				
	15	15	ROGERS		3	1035	1036	20.99	22.88	1.89	1.89	1				
	15	15	ROGERS		3	1036	1037	20.99	22.88	1.89	1.89	1				
RUN: 1																
16	16	16	HUDSON		9	1030	1032	21.40	26.05	4.65	.88	948				
	16	16	HUDSON		9	1030	1032	21.70	25.40	3.70	.55	292				

EXPERIMENT #: 18

PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	COLD	HOT	PLATE TEMP	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS	EST K	REMARKS
16	16	HUDSON		9	1030	1032	22.15	22.85	2.70	.0018	.29	19.4				
16	16	HUDSON		9	1038	1041	22.15	22.85	2.70	.0018	.29	19.4				
16	16	HUDSON		9	1038	1041	22.15	22.85	2.70	.0018	.29	19.4				
16	16	HUDSON		9	1044	1047	22.15	22.85	2.70	.0018	.29	19.4				
16	16	HUDSON		9	1044	1047	22.15	22.85	2.70	.0018	.29	19.4				
RUN: 2																
16	16	HUDSON		9	1050	1053	21.40	25.40	4.00	.0018	.88	97.4				
16	16	HUDSON		9	1050	1053	22.15	26.05	3.90	.0018	.88	97.4				
16	16	HUDSON		9	1111	1113	22.40	26.05	3.65	.0012	.88	101.8				
16	16	HUDSON		9	1111	1113	22.70	25.40	2.70	.0012	.55	101.8				
16	16	HUDSON		9	1111	1113	22.15	24.85	2.70	.0012	.55	101.8				
RUN: 1																
17	17	WOJCIECHOWSKI		27	1031	1035					.25	22.2	21.38			
17	17	WOJCIECHOWSKI		27	1037	1038					.47	330.5	330.5			
17	17	WOJCIECHOWSKI		27	1040	1044					.98	1322	1322			
17	17	WOJCIECHOWSKI		27	1043											
RUN: 1																
18	18	HUDSON		9	1030	1032	22.70	26.40	3.70	.0018	.55	2.4				
18	18	HUDSON		9	1038	1044	22.20	26.70	4.50	.0018	.82	9.4				
18	18	HUDSON		9	1040	1041	22.20	26.70	4.50	.0018	.82	9.4				
18	18	HUDSON		9	1111	1113	22.95	26.40	3.45	.0012	.6	3.88				
RUN: 1																
20	20	DOMONKOS		4	1030	1035					.25	50				
20	20	DOMONKOS		4	1030	1035					.27	50				
20	20	DOMONKOS		4	1030	1035					.17	13.2				
20	20	DOMONKOS		4	1030	1035					1.14	13.2				
RUN: 1																
21	21	TRUEBLOOD		26	1017	1018	19.90	25.00	5.10	.0012	1.05	11.3				
21	21	TRUEBLOOD		26	1020	1021	22.35	25.00	2.65	.0012	.88	11.3				
21	21	TRUEBLOOD		26	1027	1028	22.50	25.00	2.50	.0012	.88	11.3				
21	21	TRUEBLOOD		26	1033	1034	22.67	25.00	2.33	.0012	.77	11.3				
21	21	TRUEBLOOD		26	1033	1034	22.89	25.00	2.11	.0012	.65	11.3				
21	21	TRUEBLOOD		26	1036	1037	22.12	25.00	2.88	.0012	.55	11.3				
21	21	TRUEBLOOD		26	1045	1046	22.46	25.00	2.54	.0012	.55	11.3				
21	21	TRUEBLOOD		26	1048	1049	22.76	25.00	2.24	.0012	.32	11.3				
21	21	TRUEBLOOD		26	1059	1100	22.70	25.00	2.30	.0012	.32	11.3				
RUN: 1																
24	24	SERPOLAY		25	1020	1025	22.70	30.20	7.50	.0012	.77	0				
24	24	SERPOLAY		25	1050	1056	22.80	30.80	8.00	.0012	.37	0				
24	24	SERPOLAY		25	1103	1108	22.50	32.10	9.60	.0012	.77	280				
24	24	SERPOLAY		25	1130	1137	22.90	32.20	9.30	.0012	.77	150				
24	24	SERPOLAY		25	1143	1145	22.10	32.90	10.80	.0012	.77	150				
RUN: 1																
25	25	BORYS		3	1021	1023	19.20	26.00	6.80	.0012	1.9	2121	2121			
RUN: 1																

SAMP VOL X10**5

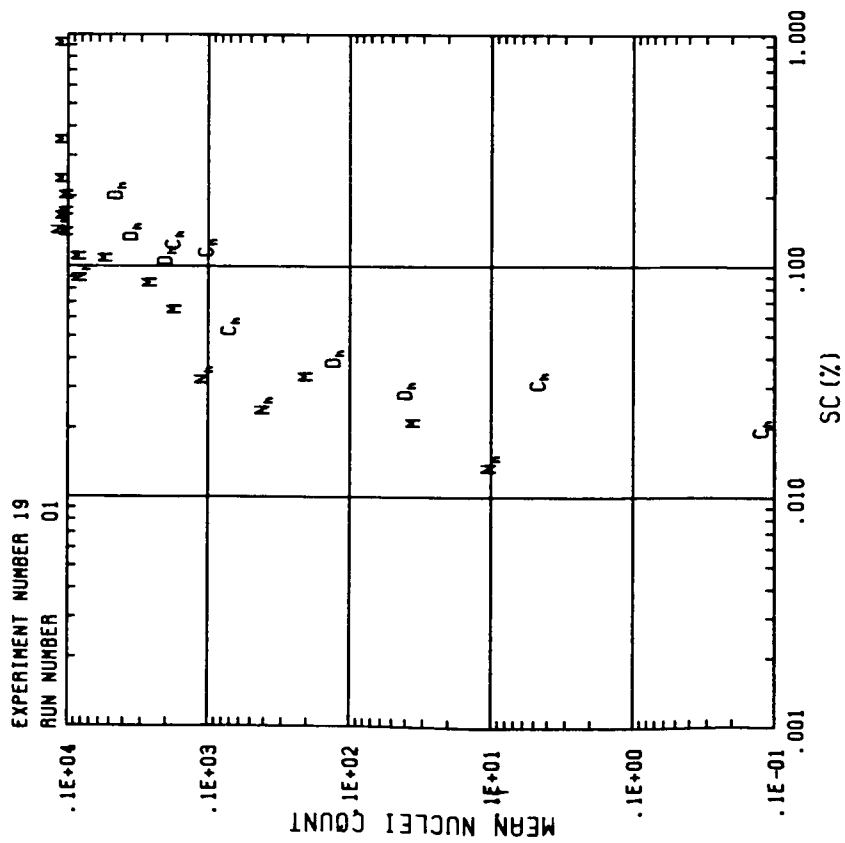
EXPERIMENT #: 18

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/09/26

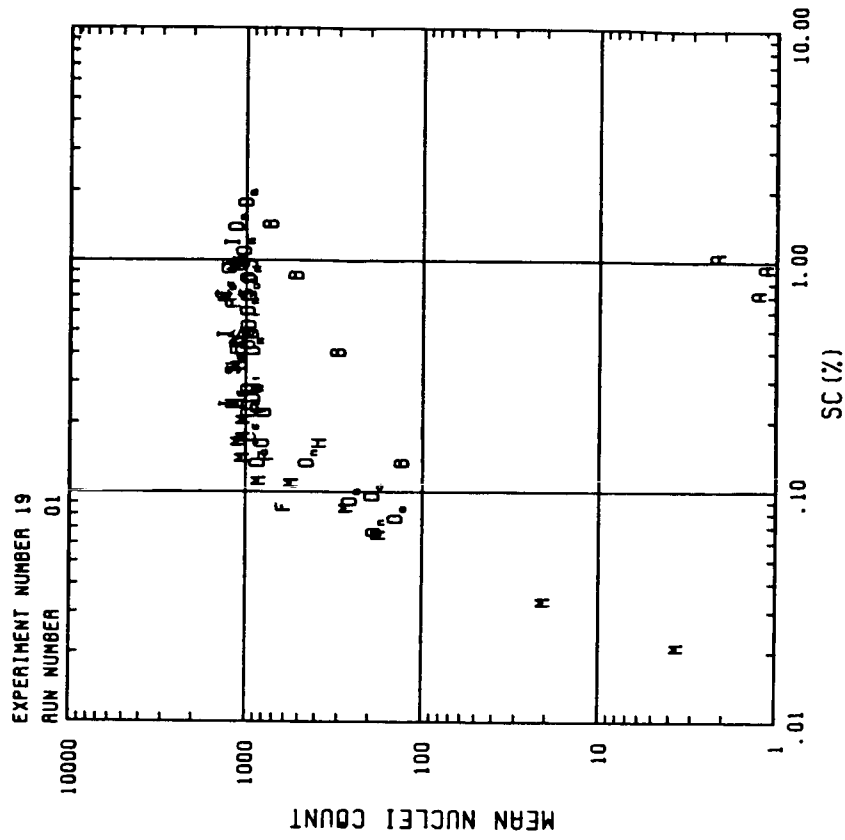
MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
							COLD	HOT (C)							
25	25	BORYS		3	1028	1031	20.60	26.08	5.40	6.36	1.2	542	542		
25	25	BORYS		3	1037	1045	23.70	26.10	2.40	12.72	1.78	59	59		
25	25	BORYS		3	1048	1055	22.70	26.20	3.50	21.22	.48	42	42		
25	25	BORYS		3	1107	1116	24.00	26.20	2.20	21.36	.19	28	28		
25	25	BORYS		3	1122	1126	20.70	25.70	5.00	6.36		365	365		
RUN: 1															
26	26	HINDMAN		6	1030	1121	26.00	26.00	0.00	1.2	.13	0	0		
26	26	HINDMAN		6	1030	1121	26.00	26.00	0.00		.12	0	0		
26	26	HINDMAN		6	1030	1121	26.00	26.00	0.00		.055	0	0		
26	26	HINDMAN		6	1030	1121	26.00	26.00	0.00		.032	0	0		
RUN: 1															
27	27	OHTAKE		19	1000	1000				7.3		.87			97% R.H.
27	27	OHTAKE		19	1000	1000				7.3		1.16			100% R.H.
27	27	OHTAKE		19	1000	1000				7.3		1.37			109% R.H.
27	27	OHTAKE		19	1000	1000				7.3		1.37			121% R.H. NOMINAL

INSTRUMENT COMPARISON



HAZE CHAMBER RESULTS

INSTRUMENT COMPARISON



SDC/CFD RESULTS

EXPERIMENT # 19

PURPOSE

INSTRUMENT COMPARISON WITH LARGE MONODISPERSE AEROSOL (FOR INCLUSION OF HAZE INSTRUMENT).

DESCRIPTION OF EXPERIMENT

DATE 13 OCT, 1980
 TIME START 1130
 TIME END 1320

NUCLEI TYPE

(NH₄)₂ SO₄, MONODISPERSE, D=0.14 UM.

GENERATION METHOD

DRI ATOMIZER, 70 PSI, 1.77 G/L SOLUTION.

SIZE DISTRIBUTION SHAPING

EC, 5600V, FLOWRATE 6 L/MIN, DILUTION FLOW 310 L/MIN.

REMARKS

NONE

WEATHER SYNOPSIS

NONE

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS		OUTSIDE CONDITIONS					
AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG WS (M/S)	AVG WD (DEG)
27.0	21.0	848					

EXPERIMENT #1 19

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/09/26

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	SAMPLE END	COLD	HOT (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
RUN: 1	2	ROGERS, DAVE	DAVE	22	1205	1206	21.20	23.40	2.20		.17		853		
	2	ROGERS, DAVE	DAVE	22	1209	1210	21.30	23.70	2.40		.23		831		
RUN: 2	2	ROGERS, DAVE	DAVE	22	1224	1225	22.40	24.50	2.10		.17		728		
	2	ROGERS, DAVE	DAVE	22	1226	1227	22.40	24.80	2.40		.23		941		
RUN: 1	5	KITCHEN		12	1138	1139	24.70	28.00	3.30		.42	359			
	5	KITCHEN		12	1139	1140	26.70	28.00	1.30		.14	126			
	5	KITCHEN		12	1140	1141	27.30	28.00	0.70		.42	268			
	5	KITCHEN		12	1141	1142	27.30	28.00	0.70		.13	541			
	5	KITCHEN		12	1142	1143	21.90	28.00	6.10		1.3	760			
RUN: 2	5	KITCHEN		12	1143	1144	20.50	28.00	7.50		2.4	102			
	5	KITCHEN		12	1144	1145	21.90	28.00	6.10		1.5	224			
	5	KITCHEN		12	1145	1146	22.30	28.00	5.70		1.4	224			
	5	KITCHEN		12	1146	1147	22.30	28.00	5.70		1.4	224			
	5	KITCHEN		12	1147	1148	21.90	28.00	6.10		1.5	224			
RUN: 1	8	ME		17	1140	1141	28.60	29.60	1.00		.075	570			
	8	ME		17	1141	1142	29.60	29.60	0.00		.075	771			
	8	ME		17	1142	1143	29.60	29.60	0.00		.075	1104			
	8	ME		17	1143	1144	29.60	29.60	0.00		.075	1104			
	8	ME		17	1144	1145	29.60	29.60	0.00		.075	1104			
	8	ME		17	1145	1146	29.60	29.60	0.00		.075	1104			
	8	ME		17	1146	1147	29.60	29.60	0.00		.075	1104			
	8	ME		17	1147	1148	29.60	29.60	0.00		.075	1104			
RUN: 1	9	GAGIN		5	1144	1145	21.70	26.70	5.00		1.05	1150			
	9	GAGIN		5	1147	1148	23.20	27.70	4.50		.70	900			
	9	GAGIN		5	1153	1154	25.40	27.60	2.20		.17	400			
RUN: 2	9	GAGIN		5	1157	1158	21.70	27.30	5.60			900			
	9	GAGIN		5	1159	1160	23.70	27.30	3.60			900			
	9	GAGIN		5	1206	1207	24.80	27.50	2.70			330			
RUN: 3	9	GAGIN		5	1210	1211	22.20	27.10	4.90		.93	1400			
	9	GAGIN		5	1212	1213	23.80	27.60	3.80		.54	960			
	9	GAGIN		5	1216	1217	24.90	27.70	2.80		.29	680			
	9	GAGIN		5	1218	1219	25.80	27.50	1.70		.13	330			
RUN: 4	9	GAGIN		5	1222	1223	22.10	27.40	5.30		1.09	1200			
	9	GAGIN		5	1224	1225	23.60	27.80	4.20		.66	1800			
	9	GAGIN		5	1228	1229	24.90	27.70	2.80		.29	700			
	9	GAGIN		5	1231	1232	26.00	27.50	1.50		.12	410			
RUN: 5	9	GAGIN		5	1237	1238	22.80	27.70	4.90		.93	1090			

EXPERIMENT #: 19

PROCESSING DATE: 81/09/26

MACH NO.		DUCT NO.		OBSERVER		SAMPLE TIME		PLATE TEMP		VOLUME SAMPLED		SUPER SAT		NUCLEI COUNT		OBS N		EST K		REMARKS		
				NO.		END		(C)		(L)		(%)										
9	9	9	9	GAGIN	5	1240	24.20	27.70	3.50	1.2	.47	910										
9	9	9	9	GAGIN	5	1242	25.20	27.60	2.40	1.2	.29	810										
9	9	9	9	GAGIN	5	1247	25.70	27.80	2.10	1.2	.17	380										
RUN: 6																						
9	9	9	9	GAGIN	5	1249	22.30	27.60	5.30	1.2	1.07	1430										
9	9	9	9	GAGIN	5	1251	23.70	27.40	2.70	1.2	.51	1230										
9	9	9	9	GAGIN	5	1255	23.00	28.00	2.70	1.2	.18	400										
9	9	9	9	GAGIN	5	1259	26.20	28.00	1.80	1.2												
RUN: 1																						
10	10	10	10	LALA	1	1440	23.93	26.50	2.55	.4	.25	1147.9										
10	10	10	10	LALA	1	1443	23.43	26.40	2.50	.4	.35	1251.3										
10	10	10	10	LALA	1	1445	23.77	26.40	2.50	.4	.35	1230.1										
10	10	10	10	LALA	1	1445	23.77	26.40	2.50	.4	.35	1136.1										
10	10	10	10	LALA	1	1446	23.77	26.40	2.50	.4	.75	1039.7										
10	10	10	10	LALA	1	1447	23.77	26.40	2.50	.4	.75	1039.7										
10	10	10	10	LALA	1	1448	21.98	26.26	4.37	.4	.11	991.1										
10	10	10	10	LALA	1	1448	21.98	26.26	4.37	.4	.11	991.1										
10	10	10	10	LALA	1	1449	21.98	26.26	4.37	.4	.11	991.1										
RUN: 2																						
10	10	10	10	LALA	1	1200	23.68	26.15	2.54	.4	.25	1183.8										
10	10	10	10	LALA	1	1202	23.09	26.07	2.54	.4	.35	1184.2										
10	10	10	10	LALA	1	1205	22.07	26.07	3.00	.4	.35	1125.7										
10	10	10	10	LALA	1	1206	22.07	26.06	3.58	.4	.5	1047.8										
10	10	10	10	LALA	1	1207	22.43	26.00	4.37	.4	.75	1108.3										
10	10	10	10	LALA	1	1209	21.60	25.96	4.36	.4	.75	1020.9										
10	10	10	10	LALA	1	1210	20.85	25.92	5.02	.4	.11	1094.6										
10	10	10	10	LALA	1	1210	20.85	25.92	5.02	.4	.11	942.6										
RUN: 1																						
11	11	11	11	FITZGERALD	28	1157					.15	1234										
11	11	11	11	FITZGERALD	28	1157					.04	1670										
11	11	11	11	FITZGERALD	28	1157					.03	1174										
11	11	11	11	FITZGERALD	28	1157					.02	1174										
11	11	11	11	FITZGERALD	28	1157					.01	1009										
RUN: 1																						
13	13	13	13	AYERS	2	1152	25.91	28.40	2.47	.25	.25	1375										
13	13	13	13	AYERS	2	1153	24.09	28.40	2.57	.25	.75	1355										
13	13	13	13	AYERS	2	1154	24.30	28.30	4.60	.25	.75	1100.5										
13	13	13	13	AYERS	2	1156	22.70	28.30	5.60	.25	1.25	1120.5										
RUN: 2																						
13	13	13	13	AYERS	2	1158	25.58	28.20	5.62	1.25	1.25	1200										
13	13	13	13	AYERS	2	1159	25.29	28.20	4.09	1.25	1.25	1200										
13	13	13	13	AYERS	2	1160	25.19	28.20	4.54	1.25	1.25	1200										
13	13	13	13	AYERS	2	1162	25.85	28.20	4.35	1.25	.75	1125.5										
13	13	13	13	AYERS	2	1163	25.67	28.20	4.32	1.25	.75	1125.5										
13	13	13	13	AYERS	2	1164	25.70	28.20	4.50	1.25	.25	1125.5										
RUN: 1																						
14	14	14	14	HUDSON	9	1156				.016	.21	500										
14	14	14	14	HUDSON	9	1156				.016	.21	500										
14	14	14	14	HUDSON	9	1156				.016	.21	500										
14	14	14	14	HUDSON	9	1156				.016	.21	500										

WATER IN ROYCO

F=40

EXPERIMENT #: 19

NUCLEI MEASUREMENTS

PROCESSING DATE: 8/10/9/26

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	SAMPLE TIME END	COLD	HOT (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (#)	NUCLEI COUNT	OBS N	EST K	REMARKS
RUN: 2															
14	14	HUDSON		9	1156	1233				.016	.11	403			F=40
14	14	HUDSON		9	1156	1233				.016	.04	243			F=40
14	14	HUDSON		9	1156	1233				.016	.04	183			F=40
14	14	HUDSON		9	1224	1300				.016	.21	591			F=40
14	14	HUDSON		9	1224	1300				.016	.11	357			F=40
14	14	HUDSON		9	1224	1300				.016	.11	187			F=40
14	14	HUDSON		9	1224	1300				.016	.09	262			F=40
RUN: 1															
14	14	HUDSON		9	1251	1318				.016	.21	562			
14	14	HUDSON		9	1251	1318				.016	.11	444			
14	14	HUDSON		9	1251	1318				.016	.11	226			
14	14	HUDSON		9	1251	1318				.016	.09	363			
RUN: 1															
15	15	ROGERS	FRED	23	1144	1236	1836	25.06	6.70		.86	1044			EACH PNT. IS AUG
15	15	ROGERS	FRED	23	1149	1236	1974	22.48	6.01		.15	1196			OF LAST 4 AT EACH
15	15	ROGERS	FRED	23	1152	1236	1911	22.44	5.33		.17	1077			SS. BACKGROUND'S
15	15	ROGERS	FRED	23	1200	1236	1997	22.31	4.64		.63	996			10 PER CC AT SS
15	15	ROGERS	FRED	23	1208	1236	2025	22.30	3.25		.265	1015			OF 1 PER CENT PLAT
15	15	ROGERS	FRED	23	1213	1236	2069	22.34	2.56		.14	960			LAST 3 PNTS.
15	15	ROGERS	FRED	23	1217	1236	2135	22.36	1.89		.07	470			LOWER 3 PNTS.
15	15	ROGERS	FRED	23	1233	1236	2141	22.71	1.30		.095	193			LAST 3 PNTS.
RUN: 1															
16	16	HUDSON		9	1145	1236				.024	.85	1038			
16	16	HUDSON		9	1145	1236				.024	.59	1036			
16	16	HUDSON		9	1150	1236				.024	.88	1002			
16	16	HUDSON		9	1204	1236				.024	.59	1002			
16	16	HUDSON		9	1208	1236				.024	.88	1002			
16	16	HUDSON		9	1223	1236				.024	.88	1002			
16	16	HUDSON		9	1223	1236				.024	.59	998			
RUN: 2															
16	16	HUDSON		9	1233	1236				.024	.85	1008			
16	16	HUDSON		9	1233	1236				.024	.59	1006			
16	16	HUDSON		9	1233	1236				.024	.88	1006			
16	16	HUDSON		9	1233	1236				.024	.59	1045			
16	16	HUDSON		9	1233	1236				.024	.85	1008			
RUN: 1															
17	17	WOJCIECHOWSKI		27	1147	1236				.024	.59	1008			
17	17	WOJCIECHOWSKI		27	1154	1236				.024	.59	1006			
17	17	WOJCIECHOWSKI		27	1159	1236				.024	.88	1045			
17	17	WOJCIECHOWSKI		27	1201	1236				.024	.59	1045			
RUN: 1															
18	18	HUDSON		9	1145	1236				.024	.75	1020			
18	18	HUDSON		9	1154	1236				.024	.59	1006			
18	18	HUDSON		9	1204	1236				.024	.88	1045			
18	18	HUDSON		9	1233	1236				.024	.59	1045			

EACH PNT. IS AUG
OF LAST 4 AT EACH
SS. BACKGROUND'S
10 PER CC AT SS
OF 1 PER CENT PLAT
GOOD DROP SIZE PLAT
LAST 3 PNTS.
LOWER 3 PNTS.
LAST 3 PNTS.

EXPERIMENT #: 19
 PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

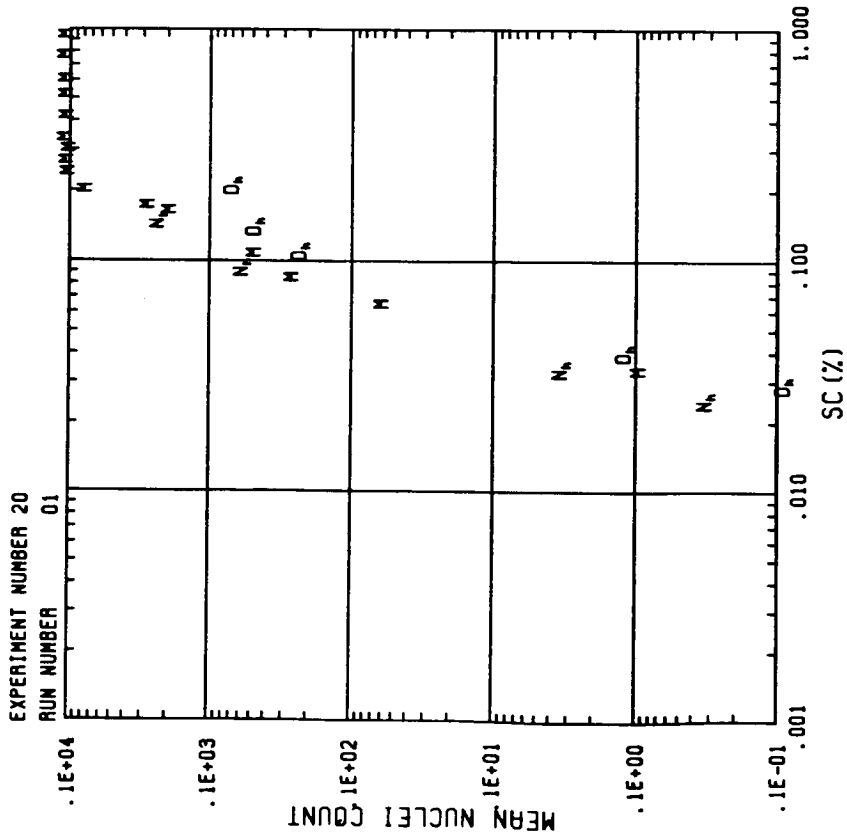
MACH NO.	DUCT NO.	NAME	USER	NO.	SAMPLE START	TIME END	PLATE TEMP		COLD	HOT (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
							23.75	25.65									
18	1A	HUDSON		9	1314	1317	23.75	25.65	1.90	.0018	.14	.897					
RUN: 1																	
21	21	TRUEBLOOD		26	1132	1133	20.05	25.00	4.95	.98	1158						
21	21	TRUEBLOOD		26	1146	1152	19.50	25.00	3.03	.37	1168						
21	21	TRUEBLOOD		26	1200	1201	22.69	25.00	2.50	.23	1186						
21	21	TRUEBLOOD		26	1204	1205	22.88	25.00	2.31	.18	1091						
21	21	TRUEBLOOD		26	1207	1208	22.30	25.00	1.92	.14	883						
21	21	TRUEBLOOD		26	1211	1212	22.30	25.00	1.70	.15	1064						
21	21	TRUEBLOOD		26	1223	1224	22.00	25.00	0.00	.19	578						
21	21	TRUEBLOOD		26	1223	1224	22.00	25.00	0.00	.05	280						
21	21	TRUEBLOOD		26	1231	1232	22.00	25.00	0.00	.05	280						
21	21	TRUEBLOOD		26	1231	1232	22.00	25.00	0.00	.05	280						
21	21	TRUEBLOOD		26	1231	1232	22.00	25.00	0.00	.05	280						
RUN: 1																	
24	24	SERPOLAY		25	1150	1157	29.20	30.70	1.50	.77	630						
24	24	SERPOLAY		25	1220	1230	29.40	32.80	3.40	.43	690						
24	24	SERPOLAY		25	1227	1231	29.60	34.00	4.40	.63	1130						
24	24	SERPOLAY		25	1241	1252	29.80	34.40	4.60	.67	1180						
24	24	SERPOLAY		25	1310	1312	30.10	32.10	2.00	.15	1300						
RUN: 1																	
25	25	BOKYS		3	1142	1145	20.60	25.50	4.90	.95	1301						
25	25	BOKYS		3	1150	1153	21.20	25.60	4.40	.46	1415						
25	25	BOKYS		3	1208	1216	23.70	26.70	3.00	.18	1104						
RUN: 1																	
26	26	HINDMAN		6	1210	1312	26.00	26.00	0.00	.13	190						
26	26	HINDMAN		6	1210	1312	26.00	26.00	0.00	.13	190						
26	26	HINDMAN		6	1210	1312	26.00	26.00	0.00	.13	190						
26	26	HINDMAN		6	1210	1312	26.00	26.00	0.00	.13	190						
RUN: 1																	
27	27	OHTAKE		19	1140	1140			3.7	.75	1.3						
27	27	OHTAKE		19	1140	1140			3.7	.75	1.3						
27	27	OHTAKE		19	1140	1140			3.7	.75	1.3						

VOL SAMP X10**5

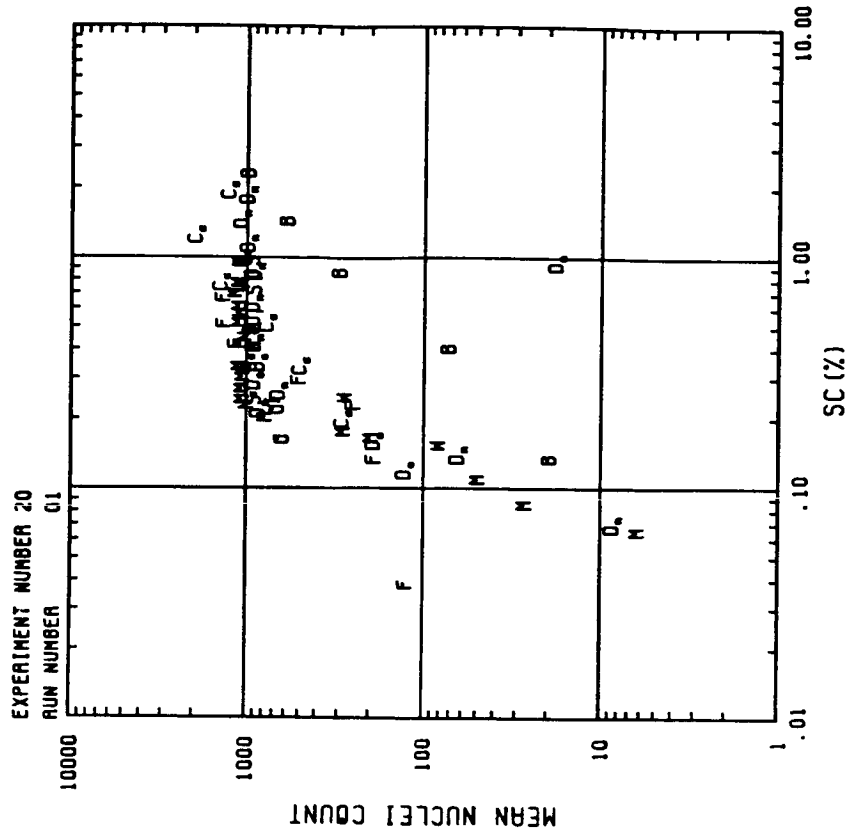
7-4.7
 7-4.2
 7-2.4
 7-0.042
 7-0.0032

75%
 97%
 109% NOMINAL

INSTRUMENT COMPARISON



INSTRUMENT COMPARISON



80/10/13

EXPERIMENT # 20

PURPOSE

INSTRUMENT COMPARISON WITH AEROSOL SIZE SUITABLE FOR CCN AND HAZE CHAMBERS.

DESCRIPTION OF EXPERIMENT

DATE 13 OCT 1980
TIME START 1352
TIME END 1545

NUCLEI TYPE

(NH₄)₂ SO₄, MONODISPERSE, D=0.08 UM.

GENERATION METHOD

DRI ATOMIZER, 70 PSI, 1.77 G/L SOLUTION.

SIZE DISTRIBUTION SHAPING

EC, 222V, FLOWRATE 4.4 L/MIN., DILUTION FLOW 370 L/MIN.

REMARKS

NONE

WEATHER SYNOPSIS

NONE

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS		OUTSIDE CONDITIONS					
AVG TEMP (C)	AVG RH (%)	AVG PHESS (MB)	AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG WS (M/S)	AVG WD (DEG)
29.0	21.0	848					

PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

EXPERIMENT #: 20

MACH NO.	DUCT NO.	OBSERVER NAME	OBSERVER NO.	PLATE TEMP		COLD	HOT (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
				SAMPLE START	SAMPLE END									
RUN: 1														
2	2	POLITOVICH	20	1444	1445	24.50	26.60	2.10		.17		662		
RUN: 2														
5	5	KITCHEN	2	1357	1358	25.60	27.50	1.90		.14		19		
5	5	KITCHEN	2	1358	1359	22.80	27.50	4.70		.16		228		
5	5	KITCHEN	2	1400	1401	21.40	27.50	6.10		1.5		1038		
5	5	KITCHEN	2	1402	1403	21.30	27.40	6.10		2.5		1578		
5	5	KITCHEN	2	1403	1404	22.80	27.50	4.70		1.3		296		
5	5	KITCHEN	2	1405	1405	25.60	27.50	1.90		.14		22		
RUN: 2														
5	5	KITCHEN	2	1406	1406	22.80	27.50	4.70		.9	304			
5	5	KITCHEN	2	1410	1410	22.80	27.50	4.70		.14	257			
5	5	KITCHEN	2	1411	1411	22.80	27.50	4.70		.14	119			
5	5	KITCHEN	2	1412	1412	22.80	27.50	4.70		.14	310			
5	5	KITCHEN	2	1413	1413	22.80	27.50	4.70		.14	313			
5	5	KITCHEN	2	1414	1414	22.80	27.50	4.70		.14	884			
RUN: 1														
10	10	LALA	13	1401	1401	23.88	26.43	2.55		.35	976	940.7		
10	10	LALA	13	1402	1402	23.34	26.38	3.04		.35	905.4	1000		
10	10	LALA	13	1403	1403	23.30	26.31	3.01		.35	1019.6	1009.8		
10	10	LALA	13	1404	1404	22.68	26.24	3.56		.35	1012.6	1007.4		
10	10	LALA	13	1405	1405	22.64	26.17	3.53		.35	972.1	945		
10	10	LALA	13	1407	1407	22.90	26.13	3.77		.35	917.93	943.3		
10	10	LALA	13	1409	1409	22.00	26.03	4.03		.35	933.6			
RUN: 2														
10	10	LALA	13	1408	1408	23.88	26.16	2.72		.35	728.4			
10	10	LALA	13	1409	1409	23.29	26.07	3.78		.35	1014			
10	10	LALA	13	1410	1410	23.66	26.06	3.60		.35	1018			
10	10	LALA	13	1411	1411	23.64	26.04	3.60		.35	1038.2			
10	10	LALA	13	1415	1415	22.75	26.12	3.37		.35	1016.3			
10	10	LALA	13	1415	1415	22.67	26.03	3.36		.35	1019.3			
RUN: 1														
11	11	FITZGERALD	28	1404	1421					.15	252	252		
11	11	FITZGERALD	28	1404	1421					.04	6336	6336		
11	11	FITZGERALD	28	1404	1421					.025	.034	.034		
11	11	FITZGERALD	28	1404	1421					.014	.034	.034		
RUN: 2														
11	11	FITZGERALD	28	1426	1443					.15	227	227		
11	11	FITZGERALD	28	1426	1443					.094	54.1	54.1		
11	11	FITZGERALD	28	1426	1443					.025	.069	.069		
11	11	FITZGERALD	28	1426	1443					.014	.069	.069		

PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

EXPERIMENT #: 20

MACH NO.	DUCT NO.	OBSERVER	NO.	SAMPLE START	TIME END	COLD	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
							HOT (C)	COLD							
16	16	HUDSON	9	1415	1418	22.15	25.85	2.70	.0018	.29	984				
16	16	HUDSON	9	1415	1419	22.15	26.05	4.65	.0018	.88	978				
16	16	HUDSON	9	1417	1423	22.15	26.40	3.70	.0018	.55	976				
16	16	HUDSON	9	1420	1423	22.15	26.40	4.65	.0018	.29	959				
16	16	HUDSON	9	1420	1423	22.15	26.85	2.70	.0018	.29	943				
16	16	HUDSON	9	1420	1423	22.15	26.85	2.70	.0018	.29	933				
RUN: 2															
16	16	HUDSON	9	1458	1500	21.40	26.10	4.70	.0012	.55	958				
16	16	HUDSON	9	1458	1500	21.40	25.40	3.70	.0012	.29	943				
16	16	HUDSON	9	1453	1516	21.40	26.10	4.70	.0024	.55	927				
16	16	HUDSON	9	1453	1516	21.40	25.40	3.70	.0024	.55	926				
16	16	HUDSON	9	1453	1516	21.40	24.90	2.70	.0024	.55	915				
RUN: 1															
17	17	WOJCIECHOWSKI	27	1422	1429	23.60	25.95	2.35	.0018	.22	899	1040			
17	17	WOJCIECHOWSKI	27	1427	1433	23.60	26.10	1.75	.0018	.22	961	1075			
17	17	WOJCIECHOWSKI	27	1431	1436	23.60	25.80	2.95	.0018	.16	198	1070			
17	17	WOJCIECHOWSKI	27	1437	1439	23.30	26.25	2.95	.0024	.35	906	1140			
RUN: 1															
18	18	HUDSON	9	1415	1418	23.60	25.95	2.35	.0018	.22	899	1040			
18	18	HUDSON	9	1427	1433	23.60	26.10	1.75	.0018	.22	961	1075			
18	18	HUDSON	9	1431	1436	23.60	25.80	2.95	.0018	.16	198	1070			
18	18	HUDSON	9	1437	1439	23.30	26.25	2.95	.0024	.35	906	1140			
RUN: 1															
20	20	DOMONKOS	4	1430	1432	20.00	25.00	5.00	.0000	.61	85	85			
20	20	DOMONKOS	4	1430	1433	20.00	25.00	4.00	.0000	.49	290	290			
20	20	DOMONKOS	4	1430	1433	20.00	25.00	3.00	.0000	.54	950	950			
20	20	DOMONKOS	4	1430	1433	20.00	25.00	2.00	.0000	.61	1150	1150			
RUN: 1															
21	21	TRUEBLOOD	26	1400	1401	20.00	25.00	5.00	.0000	.81	9	9			
21	21	TRUEBLOOD	26	1400	1403	20.00	25.00	4.00	.0000	.69	39	39			
21	21	TRUEBLOOD	26	1400	1404	20.00	25.00	3.00	.0000	.54	225	225			
21	21	TRUEBLOOD	26	1400	1405	20.00	25.00	2.00	.0000	.49	225	225			
21	21	TRUEBLOOD	26	1400	1405	20.00	25.00	1.00	.0000	.35	57	57			
21	21	TRUEBLOOD	26	1400	1405	20.00	25.00	0.00	.0000	.25	9	9			
21	21	TRUEBLOOD	26	1400	1405	20.00	25.00	0.00	.0000	.25	68	68			
21	21	TRUEBLOOD	26	1400	1405	20.00	25.00	0.00	.0000	.17	4	4			
21	21	TRUEBLOOD	26	1400	1405	20.00	25.00	0.00	.0000	.19	4	4			
21	21	TRUEBLOOD	26	1400	1405	20.00	25.00	0.00	.0000	.06	1	1			
21	21	TRUEBLOOD	26	1400	1405	20.00	25.00	0.00	.0000	.03	1	1			
RUN: 1															
24	24	SEKPOLAY	25	1402	1407	29.70	30.70	1.00	.77	.04	130	130			
24	24	SEKPOLAY	25	1426	1431	29.70	31.20	2.50	.77	.14	260	260			
24	24	SEKPOLAY	25	1435	1440	29.70	32.70	3.00	.77	.31	530	530			
24	24	SEKPOLAY	25	1450	1455	30.00	33.50	3.50	.77	.44	120	120			

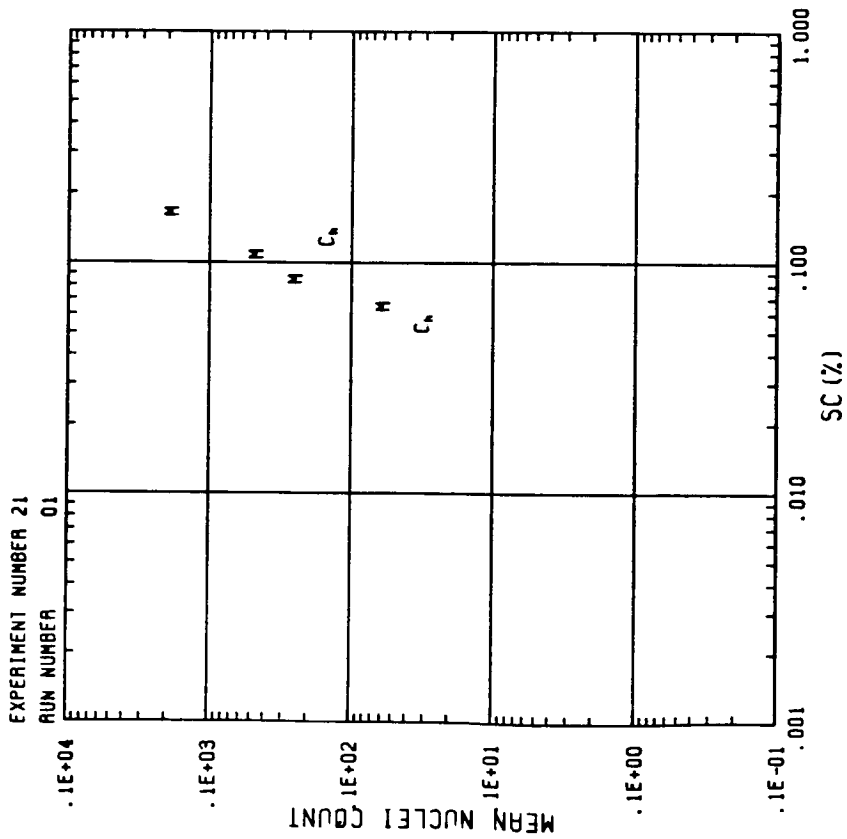
EXPERIMENT #: 20

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/09/26

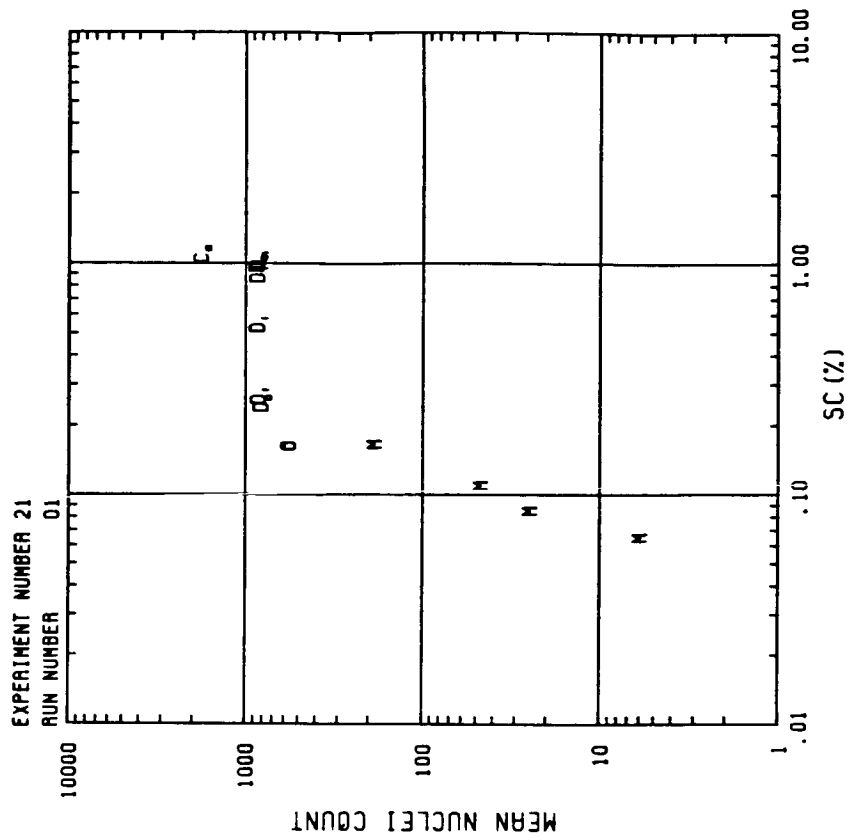
MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
							COLD	HOT (C)							
24	24	SERPOLAY		25	1505	1510	30.20	35.00	4.80	.77	.54	1410			
24	24	SERPOLAY		25	1520	1528	30.50	35.00	4.50	.77	.54	1430			
RUN: 1															
25	25	HORYS		3	1400	1405	19.00	26.00	7.00	5.3	1.95	1330			VOL SAMP X10**5
25	25	HORYS		3	1410	1415	20.50	28.90	5.40	5.36	1.52	2065			
25	25	HORYS		3	1422	1428	23.70	28.10	3.40	6.36	1.28	1485			
25	25	HORYS		3	1431	1435	22.60	28.50	3.60	10.6	.52	806			
25	25	HORYS		3	1441	1443	22.50	28.50	3.00	10.6	.52	823			
25	25	HORYS		3	1452	1453	24.50	28.80	2.30	15.9	.2	301			
RUN: 1															
27	27	OHTAKE		19	1438	1438				1.8		.277			107% NOMINAL

INSTRUMENT COMPARISON



HAZE CHAMBER RESULTS

INSTRUMENT COMPARISON



SDC/CFD RESULTS

EXPERIMENT # 21

PURPOSE

TIME VARIATION TEST TO DETERMINE REPRODUCIBILITY OF INSTRUMENT.

DESCRIPTION OF EXPERIMENT

DATE 13 OCT 1980 1545 1726

NUCLEI TYPE
SAME AS 20

GENERATION METHOD

SAME AS 20

SIZE DISTRIBUTION SHAPING

SAME AS 20

REMARKS

NONE

WEATHER SYNOPSIS

NONE

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS		OUTSIDE CONDITIONS					
AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG TEMP (C)	AVG RH (%)	AVG PRESS (MH)	AVG WS (M/S)	AVG WD (DEG)
28.0	21.0	848					

EXPERIMENT #: 21

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/09/26

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	SAMPLE TIME END	COLD	HOT (C)	PLATE TEMP	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
1	1	POLLITOV	CH	20	15:17	1626	19.90	22.00	20.30	5.20	1.00	0.17	902	612		
1	1	POLLITOV	CH	20	15:18	1709	19.10	22.50	21.00	5.20	1.00	0.17	884	659		
1	1	POLLITOV	CH	20	15:19	1631	19.80	22.00	20.30	5.20	1.00	0.17	920	588		
1	1	POLLITOV	CH	20	15:20	1631	19.30	22.50	21.00	5.20	1.00	0.17	910	582		
1	1	POLLITOV	CH	20	15:21	1631	19.30	22.50	21.00	5.20	1.00	0.17	911	637		
1	1	POLLITOV	CH	20	15:22	1631	19.80	22.00	20.30	5.20	1.00	0.17	876	585		
1	1	POLLITOV	CH	20	15:23	1631	19.50	22.50	21.00	5.20	1.00	0.17	884	609		
1	1	POLLITOV	CH	20	15:24	1631	19.70	22.00	20.30	5.20	1.00	0.17	893	623		
1	1	POLLITOV	CH	20	15:25	1631	19.90	22.50	21.00	5.20	1.00	0.17	909	623		
1	1	POLLITOV	CH	20	15:26	1631	19.70	22.00	20.30	5.20	1.00	0.17	887	627		
1	1	POLLITOV	CH	20	15:27	1631	19.30	22.50	21.00	5.20	1.00	0.17	892	627		
1	1	POLLITOV	CH	20	15:28	1631	19.50	22.00	20.30	5.20	1.00	0.17	897	629		
1	1	POLLITOV	CH	20	15:29	1631	19.30	22.50	21.00	5.20	1.00	0.17	936	629		
1	1	POLLITOV	CH	20	15:30	1631	19.70	22.00	20.30	5.20	1.00	0.17	943	629		
1	1	POLLITOV	CH	20	15:31	1631	19.30	22.50	21.00	5.20	1.00	0.17	886	629		
1	1	POLLITOV	CH	20	15:32	1631	19.50	22.00	20.30	5.20	1.00	0.17	939	629		
1	1	POLLITOV	CH	20	15:33	1631	19.70	22.50	21.00	5.20	1.00	0.17	883	629		

STD. DEV. = 64.23
COUNT AROUND
NOISE LEVEL=100CM-3

RUN: 1

12 12

12 12

12 12

EXPERIMENT #: 21

NUCLEI MEASUREMENTS

PROCESSING DATE: 01/09/26

MACH NO. DUCT NO. NAME OBSERVER NO. SAMPLE TIME START END COLD HOT PLATE TEMP DELTA VOLUME SAMPLED SUPER SAT (%) NUCLEI COUNT OBS EST K REMARKS

10 10 LALA 13 1600 1625 21.37 26.25 5.04 4 910.2 913.3 919.4 920.0 926.4 905.4 905.6 N AVG = 906.7 STD. DEV. = 26.5

Table with columns: MACH NO., DUCT NO., NAME, OBSERVER NO., SAMPLE TIME START, END, COLD, HOT, PLATE TEMP, DELTA, VOLUME SAMPLED, SUPER SAT (%), NUCLEI COUNT, OBS, EST, K, REMARKS. Contains multiple rows of data for different runs and samples.

38 POINTS (1 POINT PER MINUTE) STD. DEV. = 79.9
40 POINTS (1 POINT PER MINUTE) STD. DEV. = 96.8
BACKGROUND 10 PER

EXPERIMENT #1 21

NUCLEI MEASUREMENTS

PROCESSING DATE: 01/09/20

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	SAMPLE END	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS	EST	REMARKS
							COLD	HOT (C)							
15	15	ROGERS	FRED	23	1605	1609	24.03	25.06	1.03	1.03	895	N	K	CC. AVERAGING OVER 1 MINUTE.	
15	15	ROGERS	FRED	23	1607	1608	24.03	25.06	1.03	1.03	901	N	K	GOOD DROP SIZE PLATE AUS.	
15	15	ROGERS	FRED	23	1608	1609	24.03	25.06	1.03	1.03	898	N	K	MAIN FLOW, ABOUT 1300 COUNTS.	
15	15	ROGERS	FRED	23	1609	1610	24.03	25.06	1.03	1.03	899	N	K	ROOM TEMP 27.75 C.	
15	15	ROGERS	FRED	23	1610	1611	24.03	25.06	1.03	1.03	897	N	K		
15	15	ROGERS	FRED	23	1611	1612	24.03	25.06	1.03	1.03	898	N	K		
15	15	ROGERS	FRED	23	1612	1613	24.03	25.06	1.03	1.03	903	N	K		
RUN: 2															
15	15	ROGERS	FRED	23	1614	1615	24.03	25.06	1.03	1.03	902	N	K	K	AVERAGING OVER 1 MINUTE.
15	15	ROGERS	FRED	23	1615	1616	24.03	25.06	1.03	1.03	899	N	K	K	
15	15	ROGERS	FRED	23	1616	1617	24.03	25.06	1.03	1.03	899	N	K	K	
15	15	ROGERS	FRED	23	1617	1618	24.03	25.06	1.03	1.03	899	N	K	K	
15	15	ROGERS	FRED	23	1618	1619	24.03	25.06	1.03	1.03	890	N	K	K	
15	15	ROGERS	FRED	23	1619	1620	24.03	25.06	1.03	1.03	906	N	K	K	
15	15	ROGERS	FRED	23	1620	1621	24.03	25.06	1.03	1.03	881	N	K	K	
15	15	ROGERS	FRED	23	1621	1622	24.03	25.06	1.03	1.03	506	N	K	K	
15	15	ROGERS	FRED	23	1622	1623	20.70	23.19	2.49	2.47	487	N	K	K	
15	15	ROGERS	FRED	23	1636	1639	20.70	23.19	2.49	2.47	487	N	K	K	
RUN: 3															
15	15	ROGERS	FRED	23	1640	1641	20.70	23.19	2.49	2.47	481	N	K	K	POOR DROP SIZE PLAT. MAIN FLOW MAY BE TOO HIGH AT 1310 COUNTS.
15	15	ROGERS	FRED	23	1642	1643	20.70	23.19	2.49	2.47	464	N	K	K	READING ROYCO CHANNEL 2 MAY BE APPLYING EXCES-SIVE BACKGROUND CORRECTION AT THIS LOWER SS.
15	15	ROGERS	FRED	23	1644	1645	20.70	23.19	2.49	2.47	472	N	K	K	
15	15	ROGERS	FRED	23	1646	1649	20.70	23.19	2.49	2.47	493	N	K	K	
15	15	ROGERS	FRED	23	1650	1651	20.70	23.19	2.49	2.47	484	N	K	K	
15	15	ROGERS	FRED	23	1652	1653	20.70	23.19	2.49	2.47	486	N	K	K	
15	15	ROGERS	FRED	23	1654	1655	20.70	23.19	2.49	2.47	480	N	K	K	
15	15	ROGERS	FRED	23	1658	1659	20.70	23.19	2.49	2.47	480	N	K	K	
RUN: 4															
15	15	ROGERS	FRED	23	1700	1701	20.70	23.19	2.49	2.47	500	N	K	K	AVERAGING OVER ONE MINUTE.
15	15	ROGERS	FRED	23	1702	1705	20.70	23.19	2.49	2.47	499	N	K	K	
15	15	ROGERS	FRED	23	1704	1705	20.70	23.19	2.49	2.47	491	N	K	K	
15	15	ROGERS	FRED	23	1708	1709	20.70	23.19	2.49	2.47	488	N	K	K	
15	15	ROGERS	FRED	23	1710	1711	20.70	23.19	2.49	2.47	495	N	K	K	
15	15	ROGERS	FRED	23	1712	1713	20.70	23.19	2.49	2.47	495	N	K	K	
RUN: 1															
16	16	HUDSON	HUDSON	9	1545	1625	21.40	25.18	4.70	1.3	903	N	K	K	STAND. DEV 9
16	16	HUDSON	HUDSON	9	1545	1625	22.20	25.40	3.70	1.3	892	N	K	K	STAND. DEV 9
16	16	HUDSON	HUDSON	9	1635	1715	21.40	25.40	3.70	1.3	880	N	K	K	STAND. DEV 10
16	16	HUDSON	HUDSON	9	1635	1715	21.70	25.40	3.70	1.3	879	N	K	K	STAND. DEV 8
16	16	HUDSON	HUDSON	9	1635	1715	22.20	25.40	3.70	1.3	870	N	K	K	STAND. DEV. 9
RUN: 1															
18	18	HUDSON	HUDSON	9	1545	1625	23.30	27.30	5.00	1.3	913	N	K	K	STAND DEV. 10
18	18	HUDSON	HUDSON	9	1635	1715	23.55	26.05	2.50	1.3	851	N	K	K	STAND DEV. 21
RUN: 1															
21	21	ALOFS	ALOFS	1	1548	1549	25.00	25.00	0.00	1.73	194.2	N	K	K	
21	21	ALOFS	ALOFS	1	1548	1549	25.00	25.00	0.00	1.73	49.2	N	K	K	
21	21	ALOFS	ALOFS	1	1548	1549	25.00	25.00	0.00	1.73	25.7	N	K	K	
RUN: 2															
21	21	ALOFS	ALOFS	1	1553	1554	25.00	25.00	0.00	1.73	6.15	N	K	K	
21	21	ALOFS	ALOFS	1	1553	1554	25.00	25.00	0.00	1.73	196	N	K	K	

EXPERIMENT #: 21

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/09/26

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	COLD	HOT (C)	PLATE TEMP	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
21	21	ALOFS		1	1553	1554	25.00	25.00	25.00	0.00	0.00	.114	52			
21	21	ALOFS		1	1553	1554	25.00	25.00	25.00	0.00	0.00	.068	6.67			
RUN: 3																
21	21	ALOFS		1	1558	1559	25.00	25.00	25.00	0.00	0.00	.173	198.4			
21	21	ALOFS		1	1558	1559	25.00	25.00	25.00	0.00	0.00	.114	51.3			
21	21	ALOFS		1	1558	1559	25.00	25.00	25.00	0.00	0.00	.089	27.3			
21	21	ALOFS		1	1558	1559	25.00	25.00	25.00	0.00	0.00	.068	8.12			
RUN: 4																
21	21	ALOFS		1	1603	1604	25.00	25.00	25.00	0.00	0.00	.173	193			
21	21	ALOFS		1	1603	1604	25.00	25.00	25.00	0.00	0.00	.114	47			
21	21	ALOFS		1	1603	1604	25.00	25.00	25.00	0.00	0.00	.089	24.7			
21	21	ALOFS		1	1603	1604	25.00	25.00	25.00	0.00	0.00	.068	6.76			
RUN: 5																
21	21	ALOFS		1	1608	1609	25.00	25.00	25.00	0.00	0.00	.173	190			
21	21	ALOFS		1	1608	1609	25.00	25.00	25.00	0.00	0.00	.114	46.9			
21	21	ALOFS		1	1608	1609	25.00	25.00	25.00	0.00	0.00	.089	26.76			
21	21	ALOFS		1	1608	1609	25.00	25.00	25.00	0.00	0.00	.068	6.59			
RUN: 6																
21	21	ALOFS		1	1612	1613	25.00	25.00	25.00	0.00	0.00	.173	194			
21	21	ALOFS		1	1612	1613	25.00	25.00	25.00	0.00	0.00	.114	47.9			
21	21	ALOFS		1	1612	1613	25.00	25.00	25.00	0.00	0.00	.089	26.6			
21	21	ALOFS		1	1612	1613	25.00	25.00	25.00	0.00	0.00	.068	6.4			
RUN: 7																
21	21	ALOFS		1	1615	1615	25.00	25.00	25.00	0.00	0.00	.173	186			
21	21	ALOFS		1	1615	1616	25.00	25.00	25.00	0.00	0.00	.114	48.3			
21	21	ALOFS		1	1615	1616	25.00	25.00	25.00	0.00	0.00	.089	28.4			
21	21	ALOFS		1	1615	1616	25.00	25.00	25.00	0.00	0.00	.068	6.69			
RUN: 8																
21	21	ALOFS		1	1619	1620	25.00	25.00	25.00	0.00	0.00	.173	187.7			
21	21	ALOFS		1	1619	1620	25.00	25.00	25.00	0.00	0.00	.114	47.5			
21	21	ALOFS		1	1619	1620	25.00	25.00	25.00	0.00	0.00	.089	24.8			
21	21	ALOFS		1	1619	1620	25.00	25.00	25.00	0.00	0.00	.068	6.54			
RUN: 9																
21	21	ALOFS		1	1622	1623	25.00	25.00	25.00	0.00	0.00	.173	193			
21	21	ALOFS		1	1622	1623	25.00	25.00	25.00	0.00	0.00	.114	50.6			
21	21	ALOFS		1	1622	1623	25.00	25.00	25.00	0.00	0.00	.089	26.8			
21	21	ALOFS		1	1622	1623	25.00	25.00	25.00	0.00	0.00	.068	6.32			
RUN: 10																
21	21	ALOFS		1	1624	1625	25.00	25.00	25.00	0.00	0.00	.173	187.8			
21	21	ALOFS		1	1624	1625	25.00	25.00	25.00	0.00	0.00	.114	45			
21	21	ALOFS		1	1624	1625	25.00	25.00	25.00	0.00	0.00	.089	24.1			
21	21	ALOFS		1	1624	1625	25.00	25.00	25.00	0.00	0.00	.068	6.1			
RUN: 11																
21	21	ALOFS		1	1627	1628	25.00	25.00	25.00	0.00	0.00	.173	191.3			
21	21	ALOFS		1	1627	1628	25.00	25.00	25.00	0.00	0.00	.114	50.3			
21	21	ALOFS		1	1627	1628	25.00	25.00	25.00	0.00	0.00	.089	26.3			
21	21	ALOFS		1	1627	1628	25.00	25.00	25.00	0.00	0.00	.068	7.68			
RUN: 12																
21	21	ALOFS		1	1629	1630	25.00	25.00	25.00	0.00	0.00	.173	189.9			
21	21	ALOFS		1	1629	1630	25.00	25.00	25.00	0.00	0.00	.114	50.7			

PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

EXPERIMENT #: 21

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	SAMPLE END	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
							COLD	HOT (C)							
21	21	ALOFS			1629	1630	25.00	25.00	0.00	.068	7.19				
RUN: 13															
21	21	ALOFS			1631	1632	25.00	25.00	0.00	.173	188				
21	21	ALOFS			1631	1632	25.00	25.00	0.00	.114	45.2				
21	21	ALOFS			1631	1632	25.00	25.00	0.00	.089	5.08				
RUN: 14															
21	21	ALOFS			1634	1635	25.00	25.00	0.00	.173	185				
21	21	ALOFS			1634	1635	25.00	25.00	0.00	.114	46.4				
21	21	ALOFS			1634	1635	25.00	25.00	0.00	.089	5.97				
RUN: 15															
21	21	ALOFS			1637	1638	25.00	25.00	0.00	.173	197.4				
21	21	ALOFS			1637	1638	25.00	25.00	0.00	.114	51.2				
21	21	ALOFS			1637	1638	25.00	25.00	0.00	.089	7.56				
RUN: 16															
21	21	ALOFS			1641	1642	25.00	25.00	0.00	.173	184				
21	21	ALOFS			1641	1642	25.00	25.00	0.00	.114	43.5				
21	21	ALOFS			1641	1642	25.00	25.00	0.00	.089	5.56				
RUN: 17															
21	21	ALOFS			1644	1645	25.00	25.00	0.00	.173	187.1				
21	21	ALOFS			1644	1645	25.00	25.00	0.00	.114	48.7				
21	21	ALOFS			1644	1645	25.00	25.00	0.00	.089	5.6				
RUN: 18															
21	21	ALOFS			1650	1651	25.00	25.00	0.00	.173	184				
21	21	ALOFS			1650	1651	25.00	25.00	0.00	.114	42.8				
21	21	ALOFS			1650	1651	25.00	25.00	0.00	.089	6.2				
RUN: 19															
21	21	ALOFS			1654	1655	25.00	25.00	0.00	.173	186.5				
21	21	ALOFS			1654	1655	25.00	25.00	0.00	.114	50.9				
21	21	ALOFS			1654	1655	25.00	25.00	0.00	.089	25.8				
RUN: 20															
21	21	ALOFS			1658	1659	25.00	25.00	0.00	.173	189				
21	21	ALOFS			1658	1659	25.00	25.00	0.00	.114	48.4				
21	21	ALOFS			1658	1659	25.00	25.00	0.00	.089	26.8				
RUN: 21															
21	21	ALOFS			1701	1702	25.00	25.00	0.00	.173	187.4				
21	21	ALOFS			1701	1702	25.00	25.00	0.00	.114	52.5				
21	21	ALOFS			1701	1702	25.00	25.00	0.00	.089	28.4				
RUN: 22															
21	21	ALOFS			1704	1705	25.00	25.00	0.00	.173	186.3				
21	21	ALOFS			1704	1705	25.00	25.00	0.00	.114	48.6				
21	21	ALOFS			1704	1705	25.00	25.00	0.00	.089	26.5				

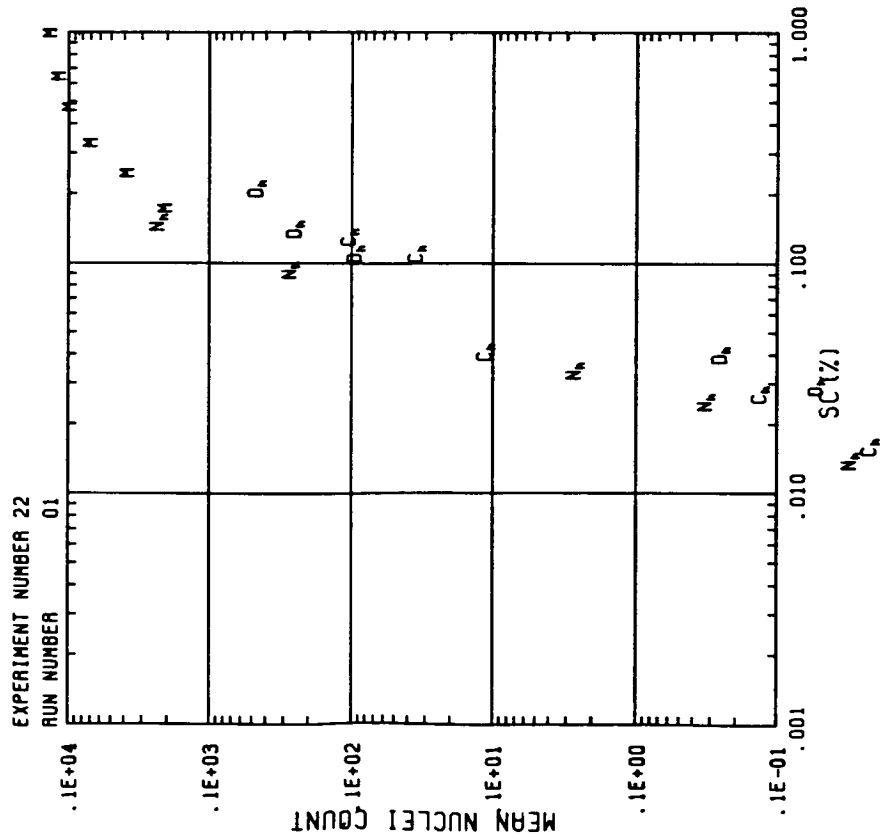
PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

EXPERIMENT #: 21

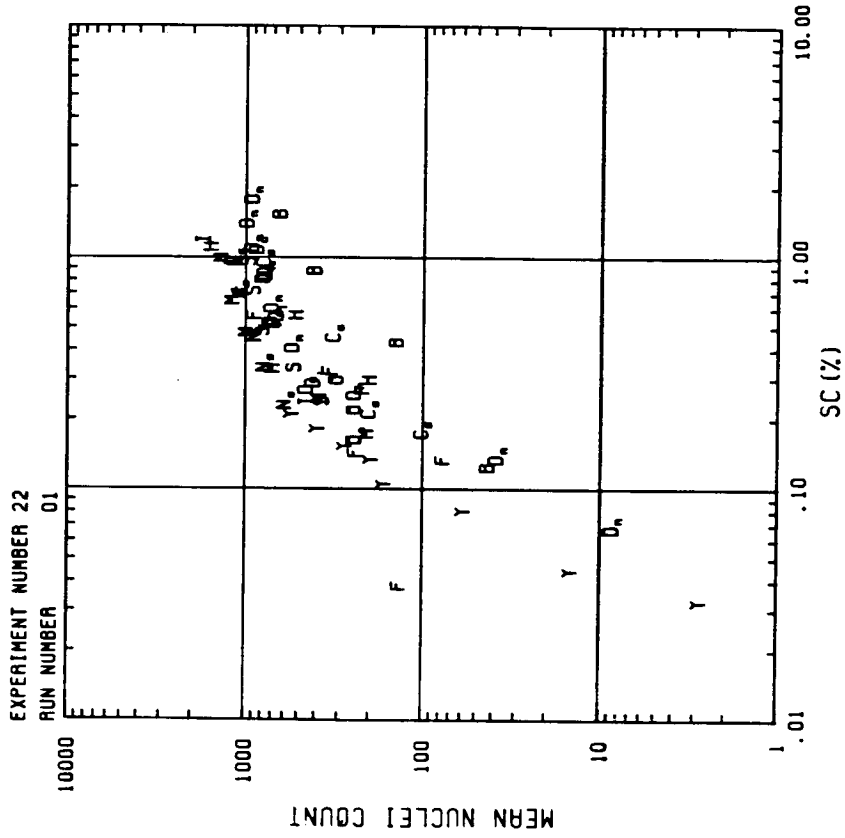
MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
							COLD	HOT (C)							
21	21	ALOFS		1	1704	1705	25.00	25.00	0.00		.068	6.3			
RUN: 23															
21	21	ALOFS		1	1707	1708	25.00	25.00	0.00		.173	198.3			
21	21	ALOFS		1	1707	1708	25.00	25.00	0.00		.114	53.6			
21	21	ALOFS		1	1707	1708	25.00	25.00	0.00		.068	27.4			
RUN: 24															
21	21	ALOFS		1	1711	1712	25.00	25.00	0.00		.173	189			
21	21	ALOFS		1	1711	1712	25.00	25.00	0.00		.114	48.26			
21	21	ALOFS		1	1711	1712	25.00	25.00	0.00		.068	7.4			
RUN: 25															
21	21	ALOFS		1	1714	1715	25.00	25.00	0.00		.173	185.1			
21	21	ALOFS		1	1714	1715	25.00	25.00	0.00		.114	47.3			
21	21	ALOFS		1	1714	1715	25.00	25.00	0.00		.068	25.1			
RUN: 1															
25	25	HORYS		3			21.60	26.70	5.10	5.3	1.1	1856			STD. DEV. = 24.6 VOL. SAMP. X 10 ^{0.5} INST. CALIB.
RUN: 2															
25	25	HORYS		3			24.30	26.90	2.60	10.6	.26	317			STD. DEV. = 35 VOL. SAMP. X 10 ^{0.5} INST. CALIB.
RUN: 1															
26	26	HINDMAN		6	1549	1626	26.00	26.00	0.00	3.4	.063	15.6			+/- 4.5 +/- 1.0
26	26	HINDMAN		6	1627	1751	26.00	26.00	0.00	1.9		3.3			

INSTRUMENT COMPARISON



HAZE CHAMBER RESULTS

INSTRUMENT COMPARISON



SDC/CFD RESULTS

EXPERIMENT # 22

PURPOSE

INSTRUMENT COMPARISON WITH PULYDISPERSE AEROSOL, MEDIUM CONC.

DESCRIPTION OF EXPERIMENT

DATE 14 OCT, 1980 TIME START 1020 END 1233

NUCLEI TYPE (NH4)2 SO4

GENERATION METHOD

DRI ATOMIZER, 30 PSI, 0.065 G/L SOLUTION, HIGH DILUTION.

SIZE DISTRIBUTION SHAPING

NONE

REMARKS

1020-1040 DRIFT UP 10%, 1040 DROP APPROX. 20%, 1052 DROP APPROX. 20%.

WEATHER SYNOPSIS

NONE

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS		OUTSIDE CONDITIONS					
AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG WS (M/S)	AVG WD (DEG)
25.0	22.0	849					

EXPERIMENT #: 22
 PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

MACH NO.	DUCT NO.	OBSERVER NAME	NO.	SAMPLE START	SAMPLE END	COLD	PLATE TEMP HOT (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (R)	NUCLEI COUNT	OBS N	EST K	REMARKS
RUN: 1														
2	2	ROGERS, DAVE	22	1057	1058	19.90	21.10	2.10		.17		259		
2	2	ROGERS, DAVE	22	1059	1100	18.90	21.30	2.40		.31		256		
2	2	ROGERS, DAVE	22	1102	1103	18.80	21.60	2.80				323		
RUN: 2														
2	2	ROGERS, DAVE	22	1128	1129	20.50	24.60	2.10		.17		244		
2	2	ROGERS, DAVE	22	1130	1131	20.50	22.60	2.10		.23		309		
2	2	ROGERS, DAVE	22	1132	1133	19.90	22.70	2.80		.31		338		
RUN: 1														
5	5	KITCHEN	12	1029	1029	21.60	23.50	1.90		.13	45			
5	5	KITCHEN	12	1030	1030	20.50	23.50	3.00		.45	148			
5	5	KITCHEN	12	1031	1031	18.80	23.50	4.70		.92	429			
5	5	KITCHEN	12	1032	1032	17.50	23.50	6.00		1.6	678			
RUN: 2														
5	5	KITCHEN	12	1034	1034	17.40	23.50	6.10		1.92	885			
5	5	KITCHEN	12	1035	1035	18.30	23.50	4.70		.45	143			
5	5	KITCHEN	12	1036	1036	20.70	23.50	3.30		.13	152			
5	5	KITCHEN	12	1037	1037	21.50	23.50	3.30		.92	142			
5	5	KITCHEN	12	1038	1038	19.90	23.50	3.30		.92	408			
5	5	KITCHEN	12	1039	1039	17.50	23.50	6.10		1.6	785			1.00
RUN: 3														
5	5	KITCHEN	12	1112	1112	22.00	23.90	1.90		.12	59			
5	5	KITCHEN	12	1113	1113	19.20	23.90	4.70		.92	275			
5	5	KITCHEN	12	1114	1114	17.80	23.90	6.10		.92	721			
5	5	KITCHEN	12	1115	1115	16.40	23.90	7.50		1.6	551			
5	5	KITCHEN	12	1116	1116	17.90	23.90	6.10		.92	571			
5	5	KITCHEN	12	1117	1117	17.40	23.90	6.10		1.92	148			
5	5	KITCHEN	12	1118	1118	19.20	23.90	4.30		.45	148			
5	5	KITCHEN	12	1119	1119	20.60	23.90	3.30		.45	148			
5	5	KITCHEN	12	1120	1120	20.60	23.90	3.30		.45	148			
RUN: 1														
7	7	LEAITCH	14	1108	1109	19.90	9.00		.283	.29	572			HOT TEMPERATURE IS ACTUALLY BETWEEN OFFER AND AMBIENT. TURB AND HUMIDITY IS 70%
7	7	LEAITCH	14	1110	1110	17.40	7.40		.283	.16	411			
7	7	LEAITCH	14	1112	1112	17.40	7.40		.283	.16	209			
7	7	LEAITCH	14	1113	1113	16.40	6.40		.283	.11	203			
7	7	LEAITCH	14	1138	1139	6.40	6.40		.283	.046	161			
7	7	LEAITCH	14	1148	1149	5.15	5.15		.283	.034	15			
7	7	LEAITCH	14	1208	1209	5.20	5.20		.283	.034	15			
7	7	LEAITCH	14	1228	1229	5.20	5.20		.283	.034	15			
RUN: 1														
9	9	GAGIN	5	1032	1032	22.30	27.80	5.50	1.22	1.16	1650			
9	9	GAGIN	5	1035	1035	24.10	24.00	3.90	1.22	.31	540			
9	9	GAGIN	5	1038	1038	25.10	28.00	2.90	1.22		210			
RUN: 2														
9	9	GAGIN	5	1049	1049	22.50	27.80	5.30	1.22	1.09	1250			
9	9	GAGIN	5	1055	1055	24.60	27.80	3.30	1.22	.41	350			
9	9	GAGIN	5	1058	1058	25.80	25.80	0.00	1.22	.11	140			
RUN: 3														
9	9	GAGIN	5	1102	1102	23.40	27.80	5.40	1.22	1.12	1360			
9	9	GAGIN	5	1105	1105	23.80	27.80	4.00	1.22	.64	500			
9	9	GAGIN	5	1108	1108	24.80	28.00	3.20	1.22		140			

EXPERIMENT #: 22
 PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	COLD	PLATE TEMP HOT (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
9	9	GAGIN		5	1112		25.80	28.10	2.30	1.2	.19	70			
RUN: 4															
9	9	GAGIN		5	1116		22.30	27.80	5.50	1.2	1.16	1200			
9	9	GAGIN		5	1121		25.10	27.80	2.70	1.2	.29	50			
9	9	GAGIN		5	1126		25.40	27.60	2.20	1.2	.17	120			
RUN: 1															
10	10	LALA		13	1028		21.27	23.77	2.50	.4	.25	412.4	390.7		
10	10	LALA		13	1030		20.82	23.77	2.95	.4	.35	365.6			
10	10	LALA		13	1032		20.80	23.75	2.94	.4	.35	571.4	558.7		
10	10	LALA		13	1033		20.78	23.75	3.01	.4	.35	853.2			
10	10	LALA		13	1034		20.37	23.69	4.28	.4	.75	905.6	844.2		
10	10	LALA		13	1035		19.34	23.62	4.93	.4	.75	990.7	948.2		
10	10	LALA		13	1036		18.65	23.53	4.93	.4	.75	1012.7	1009		
10	10	LALA		13	1037		18.60	23.53	4.93	.4	.75	1012.7			
RUN: 2															
10	10	LALA		13	1038		21.51	23.75	2.24	.4	.23	306.7			
10	10	LALA		13	1039		20.74	23.72	2.75	.4	.45	482.6			
10	10	LALA		13	1041		20.18	23.66	3.81	.4	.67	831.2			
10	10	LALA		13	1042		19.50	23.64	4.14	.4	.67	1022.1			
10	10	LALA		13	1043		18.57	23.59	4.68	.4	.67	1985.8			
10	10	LALA		13	1044		18.58	23.56	4.93	.4	.67	1115.8			
10	10	LALA		13	1045		17.47	23.51	6.00	.4	1.5	1115.8			
RUN: 3															
10	10	LALA		13	1047		21.07	23.51	2.51	.4	.25	315.2	313.9		CHANGING AEROSOL
10	10	LALA		13	1048		20.20	23.47	2.54	.4	.25	312.4			AFTER 1052
10	10	LALA		13	1049		20.06	23.47	2.59	.4	.25	493.4	478		
10	10	LALA		13	1050		19.87	23.47	2.59	.4	.25	523.2	608		
10	10	LALA		13	1051		19.92	23.47	2.59	.4	.25	727.7	714.9		
10	10	LALA		13	1052		19.21	23.47	4.05	.4	.75	767.6	747.9		
10	10	LALA		13	1053		19.21	23.47	4.05	.4	.75	758.9			
RUN: 4															
10	10	LALA		13	1054		21.83	23.07	2.27	.4	.23	240			
10	10	LALA		13	1055		20.57	23.07	2.33	.4	.23	354.8	20		
10	10	LALA		13	1056		20.57	23.05	2.33	.4	.23	593.7	247		
10	10	LALA		13	1057		20.57	23.05	2.33	.4	.23	659.2	28.8		
10	10	LALA		13	1058		19.52	23.05	4.47	.4	.23	706.5	28		
10	10	LALA		13	1059		19.22	23.05	4.47	.4	.23	767.6	.034		
10	10	LALA		13	1060		17.81	23.05	4.63	.4	1.5	723.5	.003		
RUN: 1															
11	11	FITZGERALD		28	1033	1052					.15	247	154		
11	11	FITZGERALD		28	1033	1052					.094	28.8	28		
11	11	FITZGERALD		28	1033	1052					.025	.034	.034		
11	11	FITZGERALD		28	1033	1052					.014	.003	.003		
RUN: 2															
11	11	FITZGERALD		28	1105	1125					.15	154	154		

PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

EXPERIMENT #: 22

MACH NO.	DUCT NO.	NAME	OSERVER	NO.	SAMPLE START	SAMPLE END	COLD	HOT	DELTA	VOLUME SAMPLED	SUPER SAT	NUCLEI COUNT	OBS N	EST K	REMARKS
15	15	KOCHOND		30	1138	1139	20.63	23.19	2.56	.0024	.265	259			OVER 4 PNTS EA SS.
15	15	KOCHOND		30	1142	1143	20.99	23.68	1.69	.0024	.17	49			AVG LAST 2 PNTS.
15	15	KOCHOND		30	1146	1147	21.35	23.54	1.19	.0018	.07	40			AVG LAST 2 PNTS.
15	15	KOCHOND		30	1207	1208	18.36	25.04	6.70	.0012	1.86	1018			RAISED MAIN FLO.
RUN: 1															
16	16	HUDSON		9	1054	1057	21.40	26.10	4.70	.0024	.9	783			
16	16	HUDSON		9	1054	1057	21.70	25.50	3.80	.0024	.58	680			
16	16	HUDSON		9	1054	1057	22.20	24.90	2.70	.0024	.3	429			
16	16	HUDSON		9	1146	1148	21.40	25.05	3.65	.0018	.9	701			
16	16	HUDSON		9	1146	1148	22.20	25.00	2.80	.0018	.58	436			
16	16	HUDSON		9	1153	1154	21.70	26.05	4.35	.0012	.3	452			
16	16	HUDSON		9	1153	1154	22.20	25.00	2.80	.0012	.53	713			
RUN: 2															
16	16	HUDSON		9	1200	1203	21.40	26.05	4.65	.0024	.9	848			
16	16	HUDSON		9	1200	1203	22.20	25.00	2.80	.0024	.58	731			
16	16	HUDSON		9	1210	1213	21.70	26.05	4.35	.0018	.3	452			
16	16	HUDSON		9	1210	1213	22.20	25.00	2.80	.0018	.58	713			
16	16	HUDSON		9	1224	1226	22.20	25.00	2.80	.0018	.53	456			
RUN: 1															
17	17	WOJCIECHOWSKI		27	1039	1043	20.63	23.19	2.56	.0024	.265	631.5			
17	17	WOJCIECHOWSKI		27	1047	1049	20.99	23.68	1.69	.0024	.17	828.8			
17	17	WOJCIECHOWSKI		27	1050	1054	21.35	23.54	1.19	.0018	.07	932.4			
17	17	WOJCIECHOWSKI		27	1052	1054	18.36	25.04	6.70	.0012	1.86	1228.4			
RUN: 2															
17	17	WOJCIECHOWSKI		27	1134	1136	20.63	23.19	2.56	.0024	.265	384.8			
17	17	WOJCIECHOWSKI		27	1137	1138	22.20	24.90	2.70	.0012	.9	828.8			
17	17	WOJCIECHOWSKI		27	1139	1142	21.40	25.05	3.65	.0018	.3	452			
17	17	WOJCIECHOWSKI		27	1143	1147	22.20	25.00	2.80	.0018	.58	713			
17	17	WOJCIECHOWSKI		27	1149	1151	22.20	25.00	2.80	.0018	.53	456			
RUN: 1															
18	18	HUDSON		9	1054	1057	21.40	26.10	4.70	.0024	.9	783			
18	18	HUDSON		9	1146	1148	21.70	25.50	3.80	.0012	.58	429			
18	18	HUDSON		9	1200	1203	22.20	24.90	2.70	.0024	.3	452			
18	18	HUDSON		9	1224	1226	22.20	25.00	2.80	.0018	.53	456			
RUN: 1															
21	21	TRUEBLOOD		26	1026	1027	19.89	25.00	5.11	.0024	1.04	1249			
21	21	TRUEBLOOD		26	1031	1035	20.63	25.00	4.37	.0012	.69	1048			
21	21	TRUEBLOOD		26	1039	1040	22.20	25.00	2.80	.0018	.74	405			
21	21	TRUEBLOOD		26	1042	1045	22.20	25.00	2.80	.0018	.74	405			
21	21	TRUEBLOOD		26	1047	1048	22.20	25.00	2.80	.0018	.74	405			
RUN: 2															
21	21	TRUEBLOOD		26	1104	1105	25.00	25.00	0.00	.0018	.173	178			AFTER CONC CHANGE 1052

PROCESSING DATE: 81/09/26

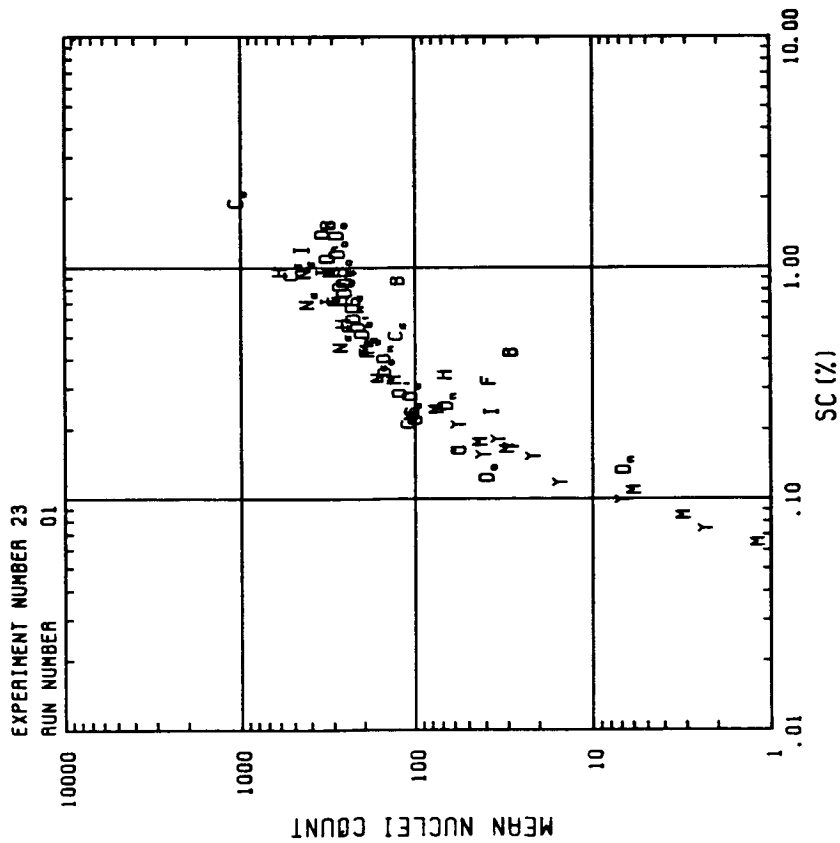
NUCLEI MEASUREMENTS

EXPERIMENT #: 22

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
							COLD	HOT (C)							
21	21	TRUEBLOOD		26	1104	1105	25.00	25.00	0.00	1.4	45.2				
21	21	TRUEBLOOD		26	1104	1105	25.00	25.00	0.00	.08	27.8				
21	21	TRUEBLOOD		26	1112	1114	25.00	25.00	0.00	.05	7.3				
21	21	TRUEBLOOD		26	1135	1135	25.00	25.00	0.00	.02	1017				
21	21	TRUEBLOOD		26	1140	1140	19.91	25.00	5.09	1.036	1017				
21	21	TRUEBLOOD		26	1144	1144	20.72	25.00	4.28	.773	691				
21	21	TRUEBLOOD		26	1148	1148	21.57	25.00	3.43	.47	472				
21	21	TRUEBLOOD		26	1152	1152	22.00	25.00	3.00	.46	267.8				
21	21	TRUEBLOOD		26	1157	1157	22.5H	25.00	2.42	.23	472				
21	21	TRUEBLOOD		26	1157	1157	22.92	25.00	2.08	.173	267.8				
RUN: 1															
24	24	SERPOLAY		25	1025	1034	25.30	26.30	1.00	.77	200				
24	24	SERPOLAY		25	1037	1045	25.70	26.70	1.00	.45	350				
24	24	SERPOLAY		25	1116	1120	25.00	30.00	5.00	.77	930				
24	24	SERPOLAY		25	1130	1135	26.70	30.70	4.00	.77	1980				
24	24	SERPOLAY		25	1135	1135	26.00	30.00	4.00	.77	250				
24	24	SERPOLAY		25	1135	1135	26.00	30.00	4.00	.77	250				
24	24	SERPOLAY		25	1201	1207	26.00	28.00	2.00	.77	80				
24	24	SERPOLAY		25	1225	1227	26.90	28.90	2.00	.77	80				
RUN: 1															
25	25	BOHYS		3	1107	1107	21.50	23.90	2.40	.2	212			.92	VOL SAMP X 10**5
25	25	BOHYS		3	1203	1208	21.90	24.10	2.20	.58	109				
25	25	BOHYS		3	1225	1225	11.40	23.90	12.50	.9	835				
25	25	BOHYS		3	1225	1225	11.00	23.90	12.90	.6	835				
RUN: 1															
26	26	HINDMAN		6	1150	1216	26.00	26.00	0.00	.13	11				-0.45
26	26	HINDMAN		6	1150	1216	26.00	26.00	0.00	.11	3.1				-0.19
26	26	HINDMAN		6	1150	1216	26.00	26.00	0.00	.07	1.2				-0.08
26	26	HINDMAN		6	1150	1216	26.00	26.00	0.00	.02	1.1				-0.0043
26	26	HINDMAN		6	1150	1216	26.00	26.00	0.00	.016	.0022				-0.0016
RUN: 1															
27	27	OHYAKE		19	1031	1031	5.5	5.5	0.00	5.5	1.43				100% R.H. NOMINAL
27	27	OHYAKE		19	1031	1031	5.5	5.5	0.00	5.5	1.71				100% R.H. NOMINAL
27	27	OHYAKE		19	1123	1123	5.5	5.5	0.00	5.5	1.54				100% R.H. NOMINAL
27	27	OHYAKE		19	1123	1123	5.5	5.5	0.00	5.5	1.91				105% R.H. NOMINAL

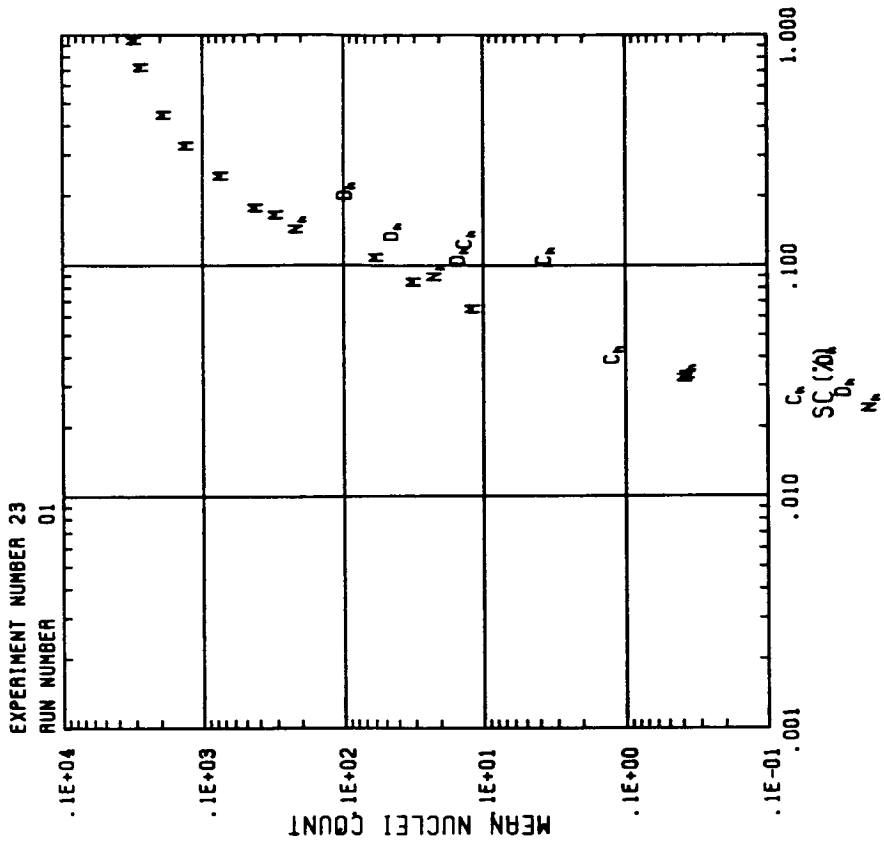
CONCENTRATION VARIATIONS

INSTRUMENT COMPARISON



M
SDC/CFD RESULTS

INSTRUMENT COMPARISON



HAZE CHAMBER RESULTS

80/10/14

EXPERIMENT # 23

PURPOSE

INSTRUMENT COMPARISON WITH LOW CONCENTRATION POLYDISPERSE ARTIFICIAL AFROSOL.

DESCRIPTION OF EXPERIMENT

DATE DATE TIME START END

14 OCT. 1980 1323 1516

NUCLEI TYPE

(NH₄)₂ SO₄, POLYDISPERSE.

GENERATION METHOD

DRI ATOMIZER, 40 PSI, HIGH DILUTION, 0.065 G/L SOLUTION.

SIZE DISTRIBUTION SHAPING

NONE

REMARKS

NONE

WEATHER SYNOPSIS

NONE

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS		OUTSIDE CONDITIONS					
AVG TEMP (C)	AVG RH (%)	AVG PRESS (MM)	AVG TEMP (C)	AVG RH (%)	AVG PRESS (MM)	AVG WS (M/S)	AVG WD (DEG)
25.0	21.0	849					

EXPERIMENT #: 23

NUCLEI MEASUREMENTS

PROCESSING DATE: 01/09/26

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	SAMPLE END	PLATE TEMP		COLD	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
							HOT (C)	HOT (C)								
RUN: 1																
2	2	ROGERS, DAVE	DAVE	22	1336	1337	24.30	24.30	22.20	2.10	.17	59	59			
2	2	ROGERS, DAVE	DAVE	22	1339	1340	24.40	24.40	22.00	2.40	.23	103	103			
RUN: 2																
2	2	POLITOVICH	POLITOVICH	20	1412	1413	24.60	24.60	22.50	2.10	.17	53	53			
2	2	POLITOVICH	POLITOVICH	20	1415	1416	24.60	24.60	22.20	2.40	.23	106	106			
RUN: 3																
5	5	KITCHEN	KITCHEN	12	1322	1323	22.10	22.10	19.80	3.30	.45	28	28			
5	5	KITCHEN	KITCHEN	12	1323	1324	22.10	22.10	17.60	4.50	.92	127	127			
5	5	KITCHEN	KITCHEN	12	1324	1325	22.10	22.10	16.00	6.10	1.02	305	305			
5	5	KITCHEN	KITCHEN	12	1325	1326	22.10	22.10	17.40	4.70	1.02	101	101			
5	5	KITCHEN	KITCHEN	12	1326	1328	22.20	22.20	18.90	3.30	.45	22	22			
RUN: 4																
7	7	LEAITCH	LEAITCH	14	1322	1323	7.60	7.60	25.20	2.30	.176	29.7	29.7			HOT TEMPERATURE IS ACTUALLY DIFFERENCE BETWEEN TUBE AND AMBIENT. INPUT HUMIDITY IS 70%
7	7	LEAITCH	LEAITCH	14	1323	1324	8.30	8.30	24.80	2.70	.256	76	76			
7	7	LEAITCH	LEAITCH	14	1325	1326	7.90	7.90	24.30	2.70	.229	58.9	58.9			
7	7	LEAITCH	LEAITCH	14	1355	1407	7.60	7.60	23.40	2.30	.164	35	35			
7	7	LEAITCH	LEAITCH	14	1406	1411	7.60	7.60	23.30	2.30	.16	42.3	42.3			
7	7	LEAITCH	LEAITCH	14	1410	1427	7.05	7.05	21.60	2.05	.124	15.7	15.7			
7	7	LEAITCH	LEAITCH	14	1426	1437	7.05	7.05	21.60	2.05	.124	15.7	15.7			
7	7	LEAITCH	LEAITCH	14	1436	1447	6.35	6.35	20.40	2.05	.1078	2.4	2.4			
RUN: 5																
8	8	MEE	MEE	17	1400	1400	27.50	27.50	25.20	2.30	.2	220	220			
8	8	MEE	MEE	17	1400	1400	27.50	27.50	24.80	2.70	.3	300	300			
8	8	MEE	MEE	17	1400	1400	27.50	27.50	23.40	2.30	.65	481	481			
8	8	MEE	MEE	17	1400	1400	27.50	27.50	23.30	2.30	1.1	373	373			
8	8	MEE	MEE	17	1400	1400	27.50	27.50	21.60	2.05	1.4	518	518			
RUN: 6																
9	9	GAGIN	GAGIN	5	1327	1327	28.20	28.20	23.10	5.10	1	620	620			
9	9	GAGIN	GAGIN	5	1331	1331	28.20	28.20	25.40	2.80	.36	270	270			
9	9	GAGIN	GAGIN	5	1335	1335	28.20	28.20	25.40	2.80	.36	270	270			
RUN: 7																
9	9	GAGIN	GAGIN	5	1347	1347	24.50	24.50	23.50	5.00	.98	560	560			
9	9	GAGIN	GAGIN	5	1352	1352	24.50	24.50	25.00	2.90	.57	350	350			
9	9	GAGIN	GAGIN	5	1359	1359	24.60	24.60	26.30	2.30	.39	100	100			
RUN: 8																
9	9	GAGIN	GAGIN	5	1425	1425	28.50	28.50	23.40	5.10	1.01	600	600			
9	9	GAGIN	GAGIN	5	1430	1430	28.50	28.50	25.60	2.90	.39	360	360			
9	9	GAGIN	GAGIN	5	1439	1439	28.60	28.60	26.40	2.20	.19	70	70			
RUN: 9																
10	10	LALA	LALA	13	1325	1325	25.12	25.12	22.60	2.52	.25	105.9	105.9			
10	10	LALA	LALA	13	1326	1326	25.05	25.05	22.54	2.52	.25	108.6	108.6			
10	10	LALA	LALA	13	1327	1327	25.02	25.02	22.04	2.98	.35	157.7	157.7			

EXPERIMENT #: 23

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/09/26

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	COLD	HOT (C)	PLATE TEMP	DELTA	VOLUME SAMPLED (L)	SUPER SA (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
10	10	LALA		13	1318		21.98	24.96		3.98	.4	.35	161.5	158.6		
10	10	LALA		13	1319		21.36	24.90		3.54	.4	.5	192.3			
10	10	LALA		13	1320		21.29	24.84		3.55	.4	.5	176.9	186.1		
10	10	LALA		13	1321		20.45	24.78		4.32	.4	.75	221.2			
10	10	LALA		13	1322		20.39	24.71		4.32	.4	.75	254.1	237.7		
10	10	LALA		13	1323		19.63	24.60		4.96	.4	.1	293.1			
10	10	LALA		13	1324		19.56	24.52		4.96	.4	.1	268.7	280.9		
RUN: 2																
10	10	LALA		13	1325		22.33	24.54		2.25	.4	.2	102.2			
10	10	LALA		13	1326		21.80	24.54		2.74	.4	.3	135.1			
10	10	LALA		13	1327		21.38	24.54		3.16	.4	.5	194.6			
10	10	LALA		13	1328		20.95	24.48		3.53	.4	.6	258.4			
10	10	LALA		13	1329		20.53	24.46		4.17	.4	.8	338.3			
10	10	LALA		13	1330		20.23	24.40		4.44	.4	.9	404.9			
10	10	LALA		13	1331		19.59	24.30		4.71	.4	.9	504.9			
10	10	LALA		13	1332		19.29	24.24		4.95	.4	.1	552.7			
10	10	LALA		13	1333		18.15	24.19		6.04	.4	.1	611.0			
RUN: 3																
10	10	LALA		13	1334		18.10	24.14		6.05	.4	.1	607.5			
10	10	LALA		13	1335		19.27	23.97		4.73	.4	.9	422.9			
10	10	LALA		13	1336		19.29	23.92		4.73	.4	.7	422.5			
10	10	LALA		13	1337		19.73	23.88		3.95	.4	.5	305.2			
10	10	LALA		13	1338		20.40	23.91		3.51	.4	.5	305.2			
10	10	LALA		13	1339		20.74	23.84		3.74	.4	.2	181.0			
10	10	LALA		13	1340		21.66	23.89		2.23	.4	.2	181.0			
RUN: 1																
11	11	FITZGERALD		28	1409	1426						.15	23	23		
11	11	FITZGERALD		28	1426	1426						.094	23	23		
11	11	FITZGERALD		28	1426	1426						.035	23	23		
11	11	FITZGERALD		28	1426	1426						.014	23	23		
RUN: 2																
11	11	FITZGERALD		28	1336	1352						.15	23	23		
11	11	FITZGERALD		28	1336	1352						.094	23	23		
11	11	FITZGERALD		28	1336	1352						.035	23	23		
11	11	FITZGERALD		28	1336	1352						.014	23	23		
RUN: 1																
13	13	AVERS		22	1347	1348	19.00	24.00		4.00	.4	.25	51	51		
13	13	AVERS		22	1349	1349	19.50	24.50		4.50	.4	.5	162	162		
13	13	AVERS		22	1401	1401	19.00	24.00		4.00	.4	.75	174	174		
13	13	AVERS		22	1403	1403	19.00	24.00		4.00	.4	.75	390	390		
13	13	AVERS		22	1403	1403	19.00	24.00		4.00	.4	.75	390	390		
13	13	AVERS		22	1405	1405	19.00	24.00		4.00	.4	.75	390	390		
13	13	AVERS		22	1406	1406	19.00	24.00		4.00	.4	.75	390	390		
13	13	AVERS		22	1407	1407	19.00	24.00		4.00	.4	.75	390	390		
13	13	AVERS		22	1408	1408	19.00	24.00		4.00	.4	.75	390	390		
13	13	AVERS		22	1409	1409	19.00	24.00		4.00	.4	.75	390	390		
13	13	AVERS		22	1410	1410	19.00	24.00		4.00	.4	.75	390	390		
13	13	AVERS		22	1411	1411	19.00	24.00		4.00	.4	.75	390	390		
13	13	AVERS		22	1412	1412	19.00	24.00		4.00	.4	.75	390	390		
13	13	AVERS		22	1413	1413	19.00	24.00		4.00	.4	.75	390	390		
13	13	AVERS		22	1414	1414	19.00	24.00		4.00	.4	.75	390	390		
13	13	AVERS		22	1415	1415	19.00	24.00		4.00	.4	.75	390	390		

EXPERIMENT #: 23

NUCLEI MEASUREMENTS

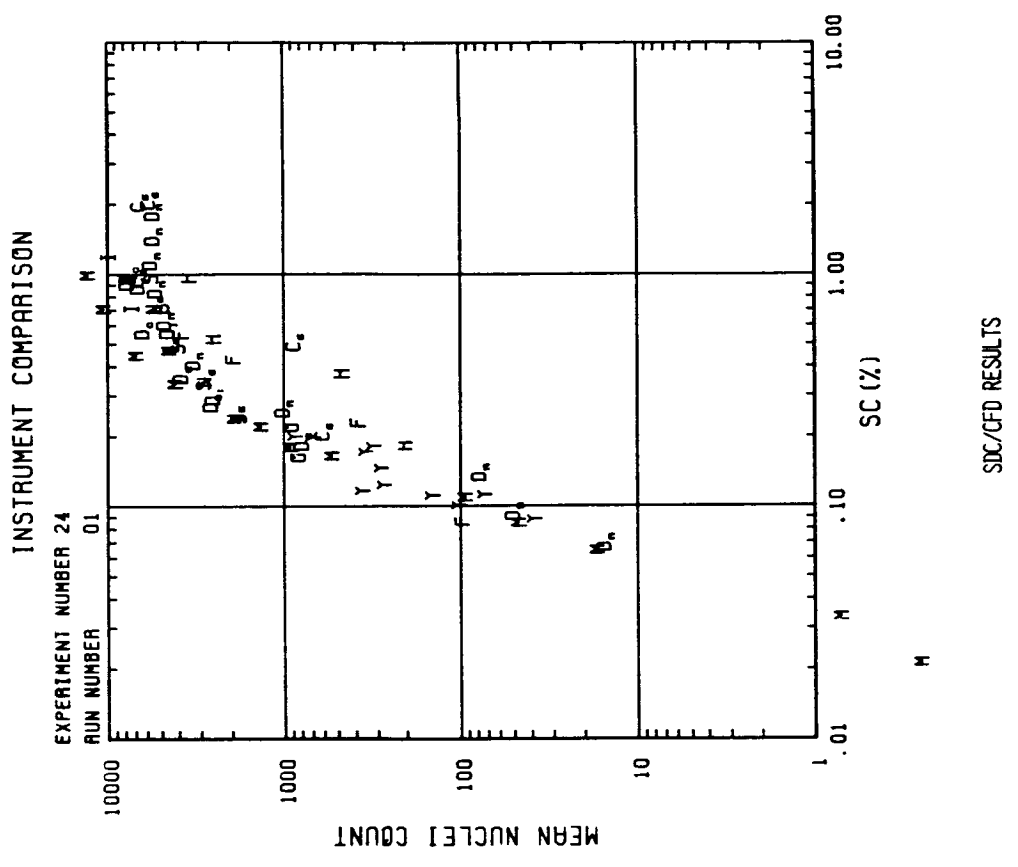
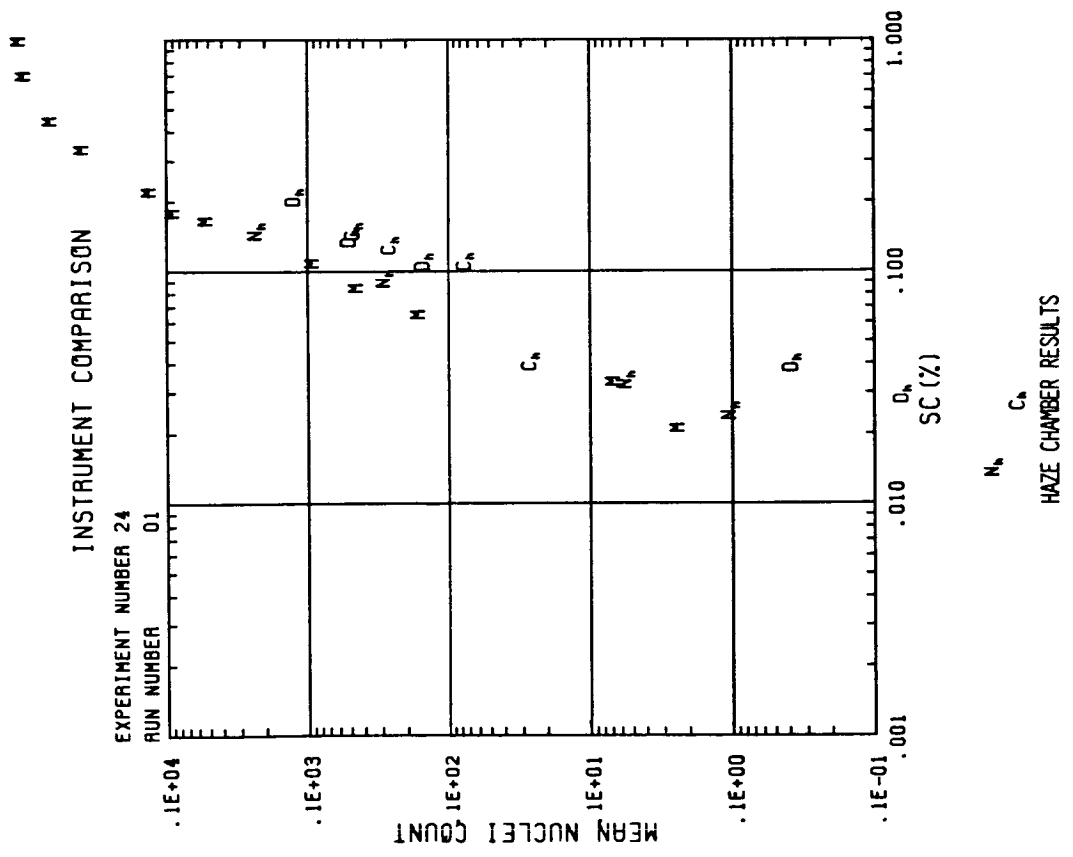
PROCESSING DATE: 81/09/26

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	COLD	HOT (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
RUN: 1															
14	14	HUDSON		9	1451	1517				.0166	.21	10.25			F=4
14	14	HUDSON		9	1451	1517				.0166	.17	4.75			F=4
14	14	HUDSON		9	1451	1517				.0166	.04	1.52			F=4
14	14	HUDSON		9	1451	1517				.0166	.029	.003			F=4
RUN: 1															
15	15	ROGERS	FRED	23	1335	1332	18.74	24.75	6.01		1.46	356			USING HIGHER MAIN FLOW USING THIS MORNINGS BACKGROUND AS FCN OF SS, MULT TIMES 2
15	15	ROGERS	FRED	23	1339	1440	19.49	24.44	4.95		1.17	331			LOWERING MAIN FLO.
15	15	ROGERS	FRED	23	1347	1448	20.25	24.11	3.86		.87	283			
15	15	ROGERS	FRED	23	1351	1452	20.63	23.50	2.87		.63	239			
15	15	ROGERS	FRED	23	1355	1456	20.99	22.88	1.89		.265	159			
RUN: 1															
16	16	HUDSON		9	1326	1329	21.35	26.05	4.70	.0024	.58	293			
16	16	HUDSON		9	1326	1329	22.20	25.20	3.00	.0024	.39	132			
16	16	HUDSON		9	1335	1339	22.13	28.05	5.92	.0024	.58	212			
16	16	HUDSON		9	1335	1339	22.25	25.20	2.95	.0024	.39	125			
16	16	HUDSON		9	1345	1348	22.17	26.05	3.88	.0024	.58	223			
16	16	HUDSON		9	1345	1348	22.42	24.25	1.83	.0024	.39	123			
RUN: 2															
16	16	HUDSON		9	1355	1359	21.35	25.05	3.70	.0024	.58	261			
16	16	HUDSON		9	1355	1359	22.20	23.33	1.13	.0024	.39	122			
16	16	HUDSON		9	1406	1410	22.57	24.58	2.01	.0024	.39	122			
16	16	HUDSON		9	1406	1410	22.57	24.58	2.01	.0024	.39	122			
16	16	HUDSON		9	1416	1418	22.17	24.58	2.41	.0024	.39	122			
16	16	HUDSON		9	1416	1418	22.42	24.25	1.83	.0024	.39	121			
RUN: 3															
16	16	HUDSON		9	1438	1441	21.35	25.05	3.70	.0024	.58	220			
16	16	HUDSON		9	1438	1441	22.20	23.33	1.13	.0024	.39	122			
16	16	HUDSON		9	1447	1451	22.13	26.05	3.92	.0024	.58	225			
16	16	HUDSON		9	1447	1451	22.25	25.20	2.95	.0024	.39	124			
16	16	HUDSON		9	1458	1451	22.17	26.05	3.88	.0024	.58	225			
16	16	HUDSON		9	1458	1451	22.42	24.25	1.83	.0024	.39	122			
RUN: 1															
17	17	WOJCIECHOWSKI		27	1349	1350					.24	108.5			108.5
17	17	WOJCIECHOWSKI		27	1352	1347					.35	122.6			172.6
17	17	WOJCIECHOWSKI		27	1400	1405					.47	281.2			281.2
17	17	WOJCIECHOWSKI		27	1404	1405					.78	453.8			453.8
RUN: 1															
18	18	HUDSON		9	1326	1329	22.25	27.25	5.00	.0024	.19	279			
18	18	HUDSON		9	1335	1339	22.45	27.15	4.70	.0024	.19	266			

EXPERIMENT #: 23
 PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
							COLD	HOT (C)							
18	18	HUDSON		9	1345	1348	26.95	26.60	3.65	.024	213				
18	18	HUDSON		9	1356	1359	26.70	26.10	2.70	.024	110				
18	18	HUDSON		9	1406	1410	26.25	25.85	3.15	.024	41				
18	18	HUDSON		9	1416	1418	26.25	26.30	3.05	.018	155				
18	18	HUDSON		9	1430	1432	25.50	27.00	4.50	.024	292				
18	18	HUDSON		9	1438	1441	25.05	27.50	5.45	.024	241				
18	18	HUDSON		9	1447	1451	21.80	26.95	5.15	.024	291				
18	18	HUDSON		9	1458	1501	21.80	27.75	5.95	.024	294				
RUN: 1															
21	21	TRUEBL00D		26	1326	1324	20.00	25.00	5.00		321.6				
21	21	TRUEBL00D		26	1330	1322	20.95	25.00	4.95		285.6				
21	21	TRUEBL00D		26	1335	1326	20.57	25.00	4.43		196.9				
21	21	TRUEBL00D		26	1339	1343	20.47	25.00	4.53		178.7				
21	21	TRUEBL00D		26	1345	1348	20.45	25.00	4.55		44.5				
21	21	TRUEBL00D		26	1401	1404	20.00	25.00	0.00		31.7				
21	21	TRUEBL00D		26	1401	1404	20.00	25.00	0.00		9.1				
21	21	TRUEBL00D		26	1401	1404	20.00	25.00	0.00		1.8				
21	21	TRUEBL00D		26	1401	1404	20.00	25.00	0.00		1.23				
21	21	TRUEBL00D		26	1401	1404	20.00	25.00	0.00		1.05				
21	21	TRUEBL00D		26	1401	1404	20.00	25.00	0.00		1.05				
RUN: 1															
24	24	SERPOLAY		25	1405	1420	26.10	27.10	1.00	.77	0				
24	24	SERPOLAY		25	1425	1430	26.20	27.20	1.00	.77	0				
24	24	SERPOLAY		25	1437	1442	26.40	27.90	1.50	.77	40				
24	24	SERPOLAY		25	1450	1455	26.60	30.60	4.00	.77	200				
24	24	SERPOLAY		25	1505	1512	26.90	31.40	4.50	.77	300				
RUN: 1															
25	25	BORYS		3	1403	1406	26.00	24.40	1.60	.22	113			.86	
25	25	BORYS		3	1423	1426	26.80	24.40	1.60	.53	134				
25	25	BORYS		3	1430	1436	19.30	24.10	4.80	1.99	537				
25	25	BORYS		3	1500	1510	17.10	24.00	6.90	1.99	1103				
RUN: 1															
26	26	HINDMAN		6	1339	1405	26.00	26.00	0.00	.13	1.4				
26	26	HINDMAN		6	1339	1405	26.00	26.00	0.00	.11	.39				
26	26	HINDMAN		6	1339	1405	26.00	26.00	0.00	.07	.13				
26	26	HINDMAN		6	1339	1405	26.00	26.00	0.00	.02	.0067				
26	26	HINDMAN		6	1339	1405	26.00	26.00	0.00	.016	.0				
RUN: 1															
27	27	OMTAKF		19	1424	1425	5.5	5.5	0.0		19			97% R.H.	
27	27	OMTAKF		19	1424	1425	5.5	5.5	0.0		154			98% R.H.	
27	27	OMTAKF		19	1424	1425	5.5	5.5	0.0		24			103% R.H. NOMINAL	



80/10/14

EXPERIMENT # 24

PURPOSE

TEST HIGH PARTICLE CONCENTRATION EFFECTS.

DESCRIPTION OF EXPERIMENT

DATE 14 OCT 1980
TIME START 1525
TIME END 1800

NUCLEI TYPE

(NH₄)₂ SO₄, POLYDISPERSE, HIGH CONC.

GENERATION METHOD

DRI ATOMIZER, 50 PSI, 0.065 G/L SOLUTION.

SIZE DISTRIBUTION SHAPING

NONE

REMARKS

NONE

WEATHER SYNOPSIS

NONE

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS		OUTSIDE CONDITIONS					
AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG WS (M/S)	AVG WD (DEG)
24.0	21.0	849					

EXPERIMENT #: 24

PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

MACH NO.	DUCT NO.	NAME	SERVER	NO.	SAMPLE START	SAMPLE END	COLD	HOT (C)	PLATE TFMP	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	ORS - N	EST K	REMARKS
RUN: 1																
2	2	ROGERS, DAVE	ROGERS, DAVE	22	1624	1625	22.00	24.10	2.10	2.10	.17		864			
2	2	ROGERS, DAVE	ROGERS, DAVE	22	1628	1629	21.80	24.20	2.40	2.40	.23		948			
RUN: 1																
7	7	LEAITCH		14	1553	1554		7.70			.283		326			HOT TEMPERATURE IS ACTUALLY DIFFERENCE BETWEEN TUBE AND AMBIENT INPUT HUMIDITY IS 72-80 %
7	7	LEAITCH		14	1603	1604		7.50			.208		371			
7	7	LEAITCH		14	1611	1612		7.75			.283		665			
7	7	LEAITCH		14	1620	1621		7.50			.121		292			
7	7	LEAITCH		14	1630	1631		7.20			.122		272			
7	7	LEAITCH		14	1639	1640		6.85			.123		542			
7	7	LEAITCH		14	1649	1651		6.80			.106		104			
7	7	LEAITCH		14	1659	1660		6.55			.108		104			
7	7	LEAITCH		14	1709	1710		6.55			.092		73.2			
7	7	LEAITCH		14	1718	1731		5.50								
RUN: 1																
8	8	MEE		17			25.00	20		3.20						
8	8	MEE		17			27.50	20		3.50						
8	8	MEE		17			27.50	20		3.50						
8	8	MEE		17			27.50	20		4.20						
8	8	MEE		17			27.50	20		5.50						
8	8	MEE		17			27.50	20		5.50						
8	8	MEE		17			27.50	20		5.50						
RUN: 1																
9	9	GAGIN		5	1552		22.40	27.50		5.10	1.2	1.01	3580			
9	9	GAGIN		5	1602		24.50	28.30		3.80	1.2	.55	2600			
9	9	GAGIN		5	1606		25.30	28.50		3.20	1.2	.39	490			
9	9	GAGIN		5	1610		26.30	28.50		2.20	1.2	.19	210			
RUN: 2																
9	9	GAGIN		5	1614		23.00	26.40		5.40	1.2	1.15	4530			
9	9	GAGIN		5	1617		24.50	28.50		4.00	1.2	.36	1750			
9	9	GAGIN		5	1625		26.20	28.60		2.40	1.2	.21	170			
RUN: 1																
10	10	LALA		13	1552		1.85	5		5.1	.4	.25	1901.4	1870.9		
10	10	LALA		13	1553		21.42	34		2.2	.4	.25	1840.3	1870.9		
10	10	LALA		13	1554		1.42	34		2.2	.4	.25	1935.3	2982.5		
10	10	LALA		13	1555		20.80	34		3.5	.4	.25	3025.7	4166.2		
10	10	LALA		13	1556		20.80	34		3.5	.4	.25	4157.6	4166.2		
10	10	LALA		13	1557		20.80	34		4.3	.4	.25	5140.9	5117		
10	10	LALA		13	1558		20.80	34		4.3	.4	.25	5140.9	5117		
10	10	LALA		13	1559		20.80	34		4.3	.4	.25	5140.9	5117		
10	10	LALA		13	1600		19.50	24		4.3	.4	.25	5140.9	5117		
10	10	LALA		13	1601		19.50	24		4.3	.4	.25	5140.9	5117		
RUN: 2																
10	10	LALA		13	1602		2.11	34		3.5	.4	.25	1144.1	1870.9		
10	10	LALA		13	1603		1.8	34		2.7	.4	.25	1425.2	1870.9		
10	10	LALA		13	1604		20.4	34		3.5	.4	.25	3319.4	4166.2		
10	10	LALA		13	1605		20.4	34		3.5	.4	.25	4650.4	4650.4		
10	10	LALA		13	1609		20.4	34		4.1	.4	.25	4650.4	4650.4		

EXPERIMENT #: 24

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/09/26

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	COLD	HOT (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
RUN: 3															
10	10	LALA		13	1608	1611	19.79	24.23	4.44	.4	.8	5218			
10	10	LALA		13	1609	1611	19.50	24.21	4.71	.4	.9	5609.3			
10	10	LALA		13	1610	1611	19.22	24.17	4.95	.4	1.5	5633.4			
10	10	LALA		13	1611	1611	18.10	24.13	6.03	.4		5916.9			
RUN: 1															
11	11	FITZGERALD		20	1603	1644	21.93	24.17	2.24	.4	.3	1083.3	250		
11	11	FITZGERALD		20	1603	1644	21.42	24.16	2.74	.4	.3	3388.5	250		
11	11	FITZGERALD		20	1603	1644	20.90	24.15	3.25	.4	.4	4103.3	31		
11	11	FITZGERALD		20	1603	1644	20.65	24.13	3.48	.4	.5	4185.5	19		
11	11	FITZGERALD		20	1603	1644	19.25	24.07	4.82	.4	.6	5159.7	19		
11	11	FITZGERALD		20	1603	1644	19.23	24.08	4.85	.4	.8	5319.7	19		
11	11	FITZGERALD		20	1603	1644	19.23	23.95	4.70	.4	.9	5544.9	19		
11	11	FITZGERALD		20	1603	1644	17.88	23.91	6.03	.4	1.5	5953.9	19		
RUN: 1															
13	13	AYERS		2	1601	1612	26.50	29.10	2.60	.5	.5	1901	250		
13	13	AYERS		2	1602	1603	26.30	29.00	2.70	.5	.5	1905	31		
13	13	AYERS		2	1604	1605	25.22	28.88	3.66	.5	.5	4455	19		
13	13	AYERS		2	1606	1607	24.43	28.87	4.44	.5	.5	8100	19		
13	13	AYERS		2	1607	1608	24.33	28.70	4.37	.5	.7	7050	19		
13	13	AYERS		2	1608	1609	23.36	28.60	5.24	.5	.7	8340	19		
13	13	AYERS		2	1609	1610	23.68	28.50	4.82	.5	1.2	7920	19		
13	13	AYERS		2	1611	1612	22.81	28.40	5.59	.5	1.2	10575	1020		
RUN: 2															
13	13	AYERS		2	1619	1620	25.92	28.20	2.28	.5	.5	1850	19		
13	13	AYERS		2	1621	1622	25.66	28.20	2.54	.5	.5	1953	19		
13	13	AYERS		2	1622	1623	24.54	28.10	3.64	.5	.5	4620	19		
13	13	AYERS		2	1623	1624	23.75	28.10	4.36	.5	.5	7125	19		
13	13	AYERS		2	1624	1625	23.01	28.00	4.99	.5	.7	7170	19		
13	13	AYERS		2	1625	1626	23.01	27.90	5.02	.5	1.1	7365	19		
13	13	AYERS		2	1626	1627	22.31	27.90	5.59	.5	1.2	7425	19		
13	13	AYERS		2	1627	1628	22.29	27.90	5.61	.5	1.2	8715	9390		
RUN: 3															
13	13	AYERS		2	1645	1646	25.29	27.80	2.51	.5	.5	2315	19		
13	13	AYERS		2	1646	1647	25.25	27.80	2.55	.5	.5	4605	19		
13	13	AYERS		2	1647	1648	24.36	27.80	3.44	.5	.5	7380	19		
13	13	AYERS		2	1648	1649	23.91	27.70	3.79	.5	.7	7020	19		
13	13	AYERS		2	1649	1650	23.41	27.60	4.19	.5	1.1	7575	19		
13	13	AYERS		2	1650	1651	23.03	27.60	4.57	.5	1.2	9450	1020		
14	14	HUDSON		9	1607	1637				.0189	.21	142			F=41

EXPERIMENT #: 24

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/09/26

MACH NO.	DUCT NO.	NAME	OBSERVER	SAMPLE NO.	START	END	COLD	PLATE TEMP HOT (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OHS N	EST K	REMARKS
14	14	HUDSON		9	1607	1637	18.36	25.06	6.70	.0189	.14	58.1			F=41
14	14	HUDSON		9	1607	1637	18.74	24.15	6.01	.0189	.04	17.5			F=41
14	14	HUDSON		9	1607	1637	19.19	24.44	5.40	.0189	.07	17.5			F=41
14	14	HUDSON		9	1638	1708	19.87	23.81	3.94	.0189	.29	124.5			F=41
14	14	HUDSON		9	1638	1708	20.25	23.50	3.30	.0189	.11	49.9			F=41
14	14	HUDSON		9	1638	1708	20.63	23.18	2.26	.0189	.11	14.8			F=41
14	14	HUDSON		9	1638	1708	20.99	22.86	1.89	.0189	.04	14.8			F=41
14	14	HUDSON		9	1638	1708	21.41	22.54	1.30	.0189	.029	.0062			F=41
RUN: 2															
14	14	HUDSON		9	1709	1751	18.36	25.06	6.70	.0182	.21	104.1			F=41
14	14	HUDSON		9	1709	1751	18.74	24.15	6.01	.0182	.11	41.1			F=41
14	14	HUDSON		9	1709	1751	19.19	24.44	5.40	.0182	.04	12.0			F=41
14	14	HUDSON		9	1709	1751	19.87	23.81	3.94	.0182	.029	.0073			F=41
RUN: 1															
15	15	ROGERS, FRED		23	1709	1710	18.36	25.06	6.70	.0182	.14	5792			:12/.62 REPLACED
15	15	ROGERS, FRED		23	1717	1714	18.74	24.15	6.01	.0182	.11	5796			CAPillary HIGH
15	15	ROGERS, FRED		23	1721	1722	19.19	24.44	5.40	.0182	.07	5921			SS PITS AT HIGH
15	15	ROGERS, FRED		23	1725	1730	19.87	23.81	3.94	.0182	.63	4984			MAIN FLOW.
15	15	ROGERS, FRED		23	1723	1730	20.25	23.50	3.30	.0182	.265	3396			LOWERED MAIN FLOW.
15	15	ROGERS, FRED		23	1737	1738	20.63	23.18	2.26	.0182	.17	1061			LOWERED MAIN FLOW.
15	15	ROGERS, FRED		23	1741	1742	20.99	22.86	1.89	.0182	.095	152			TO 1070 CNTS.
RUN: 1															
16	16	HUDSON		9	1522	1554	21.70	26.06	4.30	.0018	.58	6951			
16	16	HUDSON		9	1552	1554	21.35	25.50	3.85	.0018	.3	4092			
16	16	HUDSON		9	1609	1622	21.35	26.06	4.70	.0018	.58	7013			
16	16	HUDSON		9	1609	1622	21.70	26.06	4.05	.0018	.58	5033			
16	16	HUDSON		9	1619	1620	21.35	26.06	3.70	.0025	.3	7206			
16	16	HUDSON		9	1619	1620	22.12	25.50	3.85	.0025	.58	5150			
RUN: 2															
16	16	HUDSON		9	1629	1632	21.30	26.06	4.70	.0004	.58	5917			
16	16	HUDSON		9	1629	1640	21.70	26.06	3.85	.0004	.58	4628			
16	16	HUDSON		9	1638	1640	21.30	26.06	4.70	.0003	.58	6350			
16	16	HUDSON		9	1638	1640	21.70	26.06	3.85	.0004	.58	4974			
16	16	HUDSON		9	1655	1658	21.30	26.06	4.70	.0004	.58	2538			
16	16	HUDSON		9	1655	1658	22.12	25.50	3.85	.0004	.58	6393			
RUN: 3															
16	16	HUDSON		9	1711	1713	21.30	26.06	4.70	.0004	.58	4981			
16	16	HUDSON		9	1711	1713	21.70	26.06	3.85	.0004	.58	5711			
16	16	HUDSON		9	1720	1722	21.30	26.06	4.70	.0004	.58	5254			
16	16	HUDSON		9	1720	1722	21.70	26.06	3.85	.0004	.58	6439			
16	16	HUDSON		9	1729	1732	21.30	26.06	4.70	.0004	.58	5146			
16	16	HUDSON		9	1729	1732	21.70	26.06	3.85	.0004	.58	2305			
16	16	HUDSON		9	1729	1732	22.12	25.50	3.85	.0004	.58	4821			
RUN: 4															
16	16	HUDSON		9	1737	1740	21.30	26.06	4.70	.0004	.92	6346			

EXPERIMENT #: 24

NUCLEI MEASUREMENTS

PROCESSING DATE: 01/09/26

MACH NO.	DICT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	COLD	HOT	PLATE TEMP	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
16	16	HUDSON		9	1737	1740	21.70	25.50		3.80	.0004	.58	4732			
16	16	HUDSON		9	1737	1740	22.20	24.90		2.70	.0004	.59	2534			
RUN: 1																
17	17	WOJCIECHOWSKI		27	1558	1556							1983	1983		
17	17	WOJCIECHOWSKI		27	1602	1600							2932	2932		
17	17	WOJCIECHOWSKI		27	1605	1606							4602	4602		
17	17	WOJCIECHOWSKI		27	1605	1606							5742	5742		
RUN: 1																
18	18	HUDSON		9	1609	1610	22.30	27.10		4.80	.0018	.94	8269			
18	18	HUDSON		9	1639	1630	22.30	27.10		4.80	.0018	.94	7693			
18	18	HUDSON		9	1639	1630	22.30	27.10		4.80	.0018	.94	8063			
18	18	HUDSON		9	1655	1645	22.30	26.65		2.65	.0005	.58	2658			
18	18	HUDSON		9	1700	1650	22.30	26.80		3.50	.0008	.38	8693			
18	18	HUDSON		9	1700	1650	22.30	26.80		3.50	.0008	.38	8693			
18	18	HUDSON		9	1700	1650	22.30	26.80		3.50	.0008	.38	8693			
18	18	HUDSON		9	1737	1740	22.25	27.25		5.00	.0018	.11	4014			
18	18	HUDSON		9	1737	1740	22.25	27.25		5.00	.0018	.11	7924			
RUN: 1																
21	21	TRUEHLOOD		6	1553	1557	19.69	25.00		5.31	.0018	1.04	1347			
21	21	TRUEHLOOD		6	1604	1601	22.58	25.00		3.42	.0018	.68	1101			
21	21	TRUEHLOOD		6	1604	1601	22.58	25.00		3.42	.0018	.68	1101			
21	21	TRUEHLOOD		6	1606	1608	22.53	25.00		4.47	.0018	.52	4355			
21	21	TRUEHLOOD		6	1611	1612	22.43	25.00		0.00	.0018	.88	1952			
21	21	TRUEHLOOD		6	1620	1621	22.50	25.00		0.00	.0018	.73	5528			
21	21	TRUEHLOOD		6	1620	1621	22.50	25.00		0.00	.0018	.73	5528			
21	21	TRUEHLOOD		6	1620	1621	22.50	25.00		0.00	.0018	.73	5528			
21	21	TRUEHLOOD		6	1620	1621	22.50	25.00		0.00	.0018	.73	5528			
21	21	TRUEHLOOD		6	1626	1621	22.50	25.00		0.00	.0018	.22	4843			
21	21	TRUEHLOOD		6	1626	1621	22.50	25.00		0.00	.0018	.22	4843			
21	21	TRUEHLOOD		6	1626	1621	22.50	25.00		0.00	.0018	.22	4843			
21	21	TRUEHLOOD		6	1626	1621	22.50	25.00		0.00	.0018	.22	4843			
RUN: 1																
24	24	SERPOLAY		25	1615	1620	26.80	26.30		1.50	.0018	.09	100			
24	24	SERPOLAY		25	1630	1640	26.70	26.20		3.50	.0018	.45	400			
24	24	SERPOLAY		25	1648	1652	26.80	26.80		4.00	.0018	.58	2010			
24	24	SERPOLAY		25	1658	1701	26.90	31.40		4.50	.0018	.74	5050			
RUN: 1																
25	25	BORYS		3	1629	1638	21.60	23.90		2.30	.0018	.21	632	632	1.01	SAMP VOL X 10**5
25	25	BORYS		3	1643	1652	19.90	23.60		3.70	.0018	.51	919	919		
25	25	BORYS		3	1705	1705	18.40	22.30		4.90	.0018	.97	7120	7120		
25	25	BORYS		3	1713	1714	16.30	23.30		7.00	.0018	2.05	6932	6932		
25	25	BORYS		3	1713	1714	16.40	23.30		6.90	.0018	2.05	6932	6932		
RUN: 1																
26	26	HINDMAN		6	1600	1625	26.00	26.00		0.00	.0018	.13	28	28		
26	26	HINDMAN		6	1600	1625	26.00	26.00		0.00	.0018	.13	28	28		
26	26	HINDMAN		6	1600	1625	26.00	26.00		0.00	.0018	.13	28	28		
26	26	HINDMAN		6	1600	1625	26.00	26.00		0.00	.0018	.13	28	28		
26	26	HINDMAN		6	1600	1625	26.00	26.00		0.00	.0018	.13	28	28		
26	26	HINDMAN		6	1731	1747	26.00	26.00		0.00	.0018	.15	51	51		
RUN: 1																
27	27	ONTAKE		19	1601	1601					.73		9.43			95% R.H.

EXPERIMENT #: 24
 MACH DUCT NO. 27
 OBSERVER
 NAME OHTAKE
 NO. 19
 SAMPLE START TIME 1601
 END TIME 1601
 COLD
 PLATE TEMP
 HOT (C)
 DELTA
 VOLUME SAMPLED (L) 0.73
 SUPER SAT (%)
 NUCLEI COUNT 10.48
 OBS N
 EST K
 REMARKS 101% R.H. NOMINAL
 PROCESSING DATE: 81/09/26

PURPOSE

CLEAN AIR FOR NOISE CHECK IN INSTRUMENT.

DESCRIPTION OF EXPERIMENT

DATE 15 OCT. 1980 0820 0921
 TIME START END

NUCLEI TYPE NA

GENERATION METHOD NA

SIZE DISTRIBUTION SHAPING NA

REMARKS

NONE

WEATHER SYNOPSIS

NONE

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS		OUTSIDE CONDITIONS					
AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG WS (M/S)	AVG WD (DEG)
27.0	22.0	841					

PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

EXPERIMENT #: 25

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE TIME START	SAMPLE TIME END	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
							COLD	HOT (C)							
RUN: 1															
8	8	WEE		17	0850	0900	23.70	30.30	1.60	.07	0				
8	8	WEE		17	0850		23.30	30.30	1.40	.19	20				
8	8	WEE		17	0850		23.30	30.30	3.30	.32	19				
8	8	WEE		17	0850		23.30	30.30	4.20	.47	18				
8	8	WEE		17	0850		23.30	30.30	4.50	.71	20				
8	8	WEE		17	0850		23.30	30.30	5.20	.73	19				
8	8	WEE		17	0850		23.30	30.30	5.60	1.14	16				
8	8	WEE		17	0850		23.30	30.30	5.50	1.67	11				
8	8	WEE		17	0850		23.30	30.30	6.40	1.71	16				
8	8	WEE		17	0850		23.30	30.30	6.70	1.67	11				
8	8	WEE		17	0850	0900	23.30	30.30	7.30	2.24	5				
8	8	WEE		17	0850		23.30	30.30	7.00	2.2	5				
RUN: 1															
9	9	GAGIN		5	0949		22.20	27.50	5.30	1.11	20				
9	9	GAGIN		5	0952		23.50	27.40	3.90	.61	60				
9	9	GAGIN		5	0954		24.20	27.50	2.20	.25	26				
9	9	GAGIN		5	0957		25.20	27.50	2.30	.42	70				
RUN: 2															
9	9	GAGIN		5	0902		22.40	27.50	5.10	1.01	40				
9	9	GAGIN		5	0904		23.40	27.40	4.00	.64	50				
9	9	GAGIN		5	0906		23.50	27.60	3.40	.42	16				
9	9	GAGIN		5	0908		25.30	27.50	2.20	.2	1				
RUN: 1															
10	10	LALA		13	0823		19.22	22.24	2.46	.25	1				
10	10	LALA		13	0824		19.22	22.39	2.97	.35	8				
10	10	LALA		13	0825		19.28	22.50	2.92	.44	4				
10	10	LALA		13	0826		19.38	22.71	3.29	.35	4				
10	10	LALA		13	0827		19.38	22.61	3.48	.35	3				
10	10	LALA		13	0828		19.38	22.61	3.48	.35	3				
10	10	LALA		13	0829		18.35	22.34	4.26	.72	8				
10	10	LALA		13	0830		18.35	22.34	4.26	.72	7				
10	10	LALA		13	0831		18.35	22.34	4.91	1	1				
10	10	LALA		13	0832		18.35	22.34	4.91	1	1				
RUN: 1															
11	11	FITZGERALD		28	0910	0920				.15	34				
11	11	FITZGERALD		28	0910	0920				.04	32				
11	11	FITZGERALD		28	0910	0920				.034	30				
11	11	FITZGERALD		28	0910	0920				.025	0				
11	11	FITZGERALD		28	0910	0920				.014	0				
RUN: 1															
13	13	AYERS		2	0830	0831	17.40	22.80	5.40	1.25	0				
13	13	AYERS		2	0832	0833	17.40	22.80	4.85	.75	0				
13	13	AYERS		2	0834	0835	18.35	22.80	4.25	.75	0				
13	13	AYERS		2	0836	0837	19.35	22.80	3.45	.25	0				
13	13	AYERS		2	0838	0839	20.35	22.80	2.45	.25	0				
RUN: 1															
14	14	HUDSON		9	0914	0938				.21	1.44			F=41	
14	14	HUDSON		9	0914	0938				.14	.36			F=41	

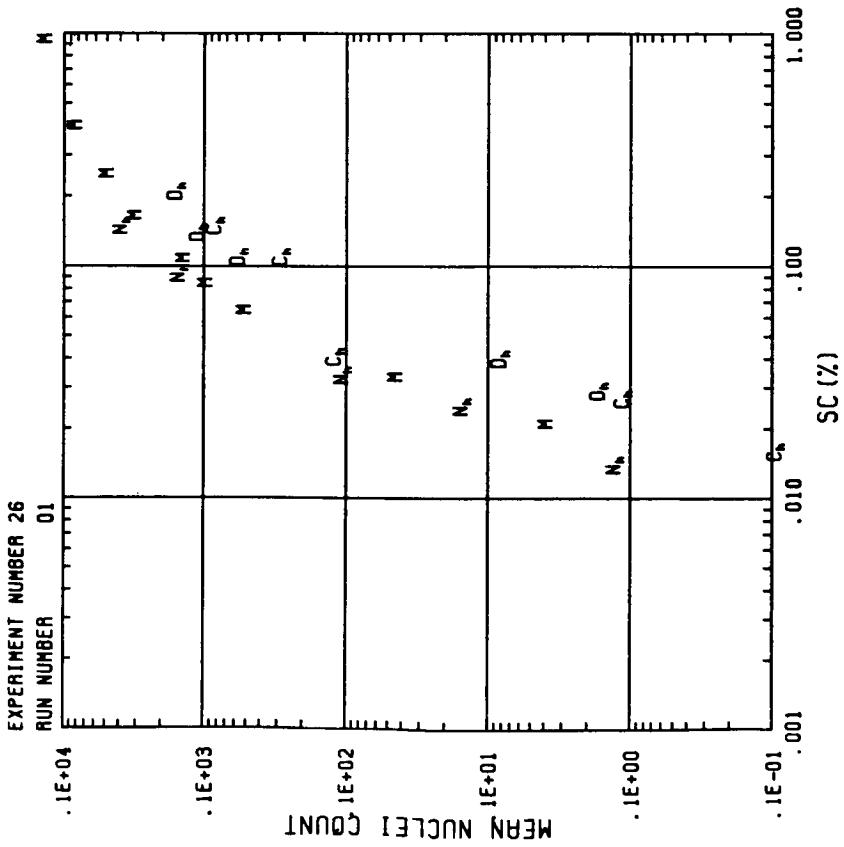
EXPERIMENT #: 25

NUCLEI MEASUREMENTS

PROCESSING DATE: R1/09/26

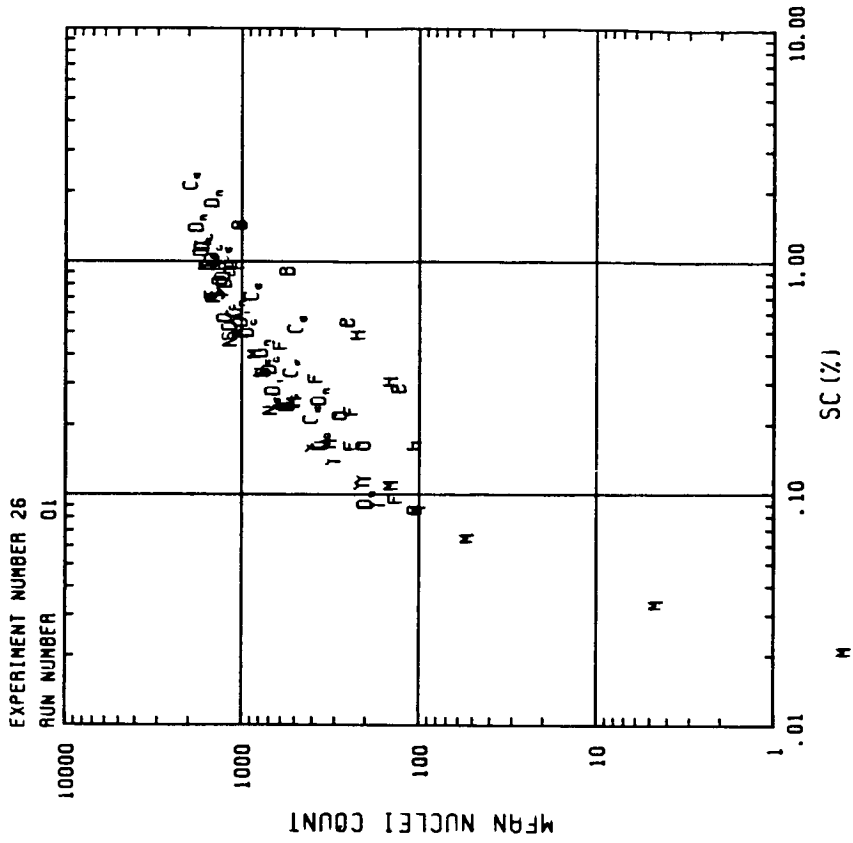
MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	COLD	HOT (C)	PLATE TEMP	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
14	14	HUDSON	HUDSON	9	0914	0938						.11	14			F=41
14	14	HUDSON	HUDSON	9	0914	0938						.029	.001			F=41
14	14	HUDSON	HUDSON	9	0914	0938						.029	.001			F=41
RUN: 1																
15	15	ROGERS	FRED	23	0833	0834	18.36	25.06		6.70		1.86	7			MAIN FLOW ABOUT
15	15	ROGERS	FRED	23	0833	0834	18.36	25.06		6.70		1.86	7			1600 COUNTS
15	15	ROGERS	FRED	23	0833	0834	18.36	25.06		6.70		1.86	7			THESE VALUES ARE
15	15	ROGERS	FRED	23	0833	0834	18.36	25.06		6.70		1.86	7			THE HIGHEST YET
15	15	ROGERS	FRED	23	0833	0834	18.36	25.06		6.70		1.86	7			OBSERVED DURING THE
15	15	ROGERS	FRED	23	0833	0834	18.36	25.06		6.70		1.86	7			WORKSHOP.
15	15	ROGERS	FRED	23	0833	0834	18.36	25.06		6.70		1.86	7			
15	15	ROGERS	FRED	23	0833	0834	18.36	25.06		6.70		1.86	7			
15	15	ROGERS	FRED	23	0833	0834	18.36	25.06		6.70		1.86	7			
RUN: 1																
16	16	HUDSON	HUDSON	9	0904	0913	21.45	25.15		4.70	.005	.9	1.67			
16	16	HUDSON	HUDSON	9	0904	0913	21.45	25.15		4.70	.005	.9	1.67			
16	16	HUDSON	HUDSON	9	0904	0913	21.45	25.15		4.70	.005	.9	1.67			
RUN: 1																
17	17	NRL	NRL	32	0835							.25		0		
17	17	NRL	NRL	32	0835							.25		0		
17	17	NRL	NRL	32	0835							.25		0		
17	17	NRL	NRL	32	0835	0830						.25		0		
17	17	NRL	NRL	32	0835	0830						.25		0		
17	17	NRL	NRL	32	0835	0830						.25		0		
RUN: 1																
18	18	HUDSON	HUDSON	9	0904	0913	22.45	27.60		5.15	.005	1.08	.34			
18	18	HUDSON	HUDSON	9	0904	0913	22.45	27.60		5.15	.005	1.08	.34			
18	18	HUDSON	HUDSON	9	0904	0913	22.45	27.60		5.15	.005	1.08	.34			
RUN: 1																
21	21	ALOFS	ALOFS	1	0842	0846	20.00	25.00		5.00		1	.307			
21	21	ALOFS	ALOFS	1	0842	0846	20.00	25.00		5.00		1	.307			
21	21	ALOFS	ALOFS	1	0842	0846	20.00	25.00		5.00		1	.307			
21	21	ALOFS	ALOFS	1	0842	0846	20.00	25.00		5.00		1	.307			
21	21	ALOFS	ALOFS	1	0842	0846	20.00	25.00		5.00		1	.307			
21	21	ALOFS	ALOFS	1	0842	0846	20.00	25.00		5.00		1	.307			
21	21	ALOFS	ALOFS	1	0842	0846	20.00	25.00		5.00		1	.307			
21	21	ALOFS	ALOFS	1	0842	0846	20.00	25.00		5.00		1	.307			
21	21	ALOFS	ALOFS	1	0842	0846	20.00	25.00		5.00		1	.307			
RUN: 1																
25	25	BORYS	BORYS	3	0857	0908	22.80	23.80		1.00	10.6	.04	57			VOL SAMP X 10**5
25	25	BORYS	BORYS	3	0910	0913	16.10	24.80		3.60	4.24	2.53	955	57		ACTUALLY < 47
25	25	BORYS	BORYS	3	0910	0913	20.40	26.00		3.60	2.12	.53	47	955		
RUN: 1																
26	26	HINDMAN	HINDMAN	6	0832	0857	26.00	26.00		0.00	1.2	.15	76			
26	26	HINDMAN	HINDMAN	6	0832	0857	26.00	26.00		0.00	1.2	.15	76			
26	26	HINDMAN	HINDMAN	6	0832	0857	26.00	26.00		0.00	1.2	.15	76			
26	26	HINDMAN	HINDMAN	6	0832	0857	26.00	26.00		0.00	1.2	.15	76			
26	26	HINDMAN	HINDMAN	6	0832	0857	26.00	26.00		0.00	1.2	.15	76			
26	26	HINDMAN	HINDMAN	6	0832	0857	26.00	26.00		0.00	1.2	.15	76			
26	26	HINDMAN	HINDMAN	6	0832	0857	26.00	26.00		0.00	1.2	.15	76			
26	26	HINDMAN	HINDMAN	6	0832	0857	26.00	26.00		0.00	1.2	.15	76			
26	26	HINDMAN	HINDMAN	6	0832	0857	26.00	26.00		0.00	1.2	.15	76			

INSTRUMENT COMPARISON



HAZE CHAMBER RESULTS

INSTRUMENT COMPARISON



SDC/CFD RESULTS

PURPOSE

INSTRUMENT COMPARISON WITH AMBIENT AIR.

DESCRIPTION OF EXPERIMENT

DATE DATE TIME START END
15 OCT. 1980 0930 1130

NUCLEI TYPE

NA

GENERATION METHOD

NA

SIZE DISTRIBUTION SHAPING

NA

REMARKS

AIR THROUGH OPEN WINDOW IN AEROSOL LAB DUCT ELBOWS IN LAB OPENED.

WEATHER SYNOPSIS

NONE

MEAN WEATHER CONDITIONS DURING EXPERIMENT

Table with columns: ROOM CONDITIONS, OUTSIDE CONDITIONS. Rows include AVG TEMP (C), AVG RH (%), AVG PRESS (MB), AVG TEMP (C), AVG RH (%), AVG PRESS (MB), AVG WS (M/S), AVG WD (DEG).

EXPERIMENT #: 26

PROCESSING DATE: 81/09/26

MACH NO.		DICT NO.	NAME	SERVER	NO.	SAMPLE START	SAMPLE END	COLD	PLATE TEMP HOT (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
RUN: 1																
2	2		POLITOVICH		20	1015	1017	22.70	24.80	2.10		.17		216		
						1018	1019	22.60	25.00	2.40		.23		294		
RUN: 1																
5	5		KITICHEN		12	1005	1005	25.50	27.10	1.60		.09	110			PHOTOCELL AMP.
5	5		KITICHEN		13	1007	1007	24.50	27.70	3.20		.28	150			RESET
5	5		KITICHEN		14	1008	1008	23.50	27.50	4.00		.25	130			
5	5		KITICHEN		15	1009	1010	23.20	27.70	4.50		.25	920			
5	5		KITICHEN		16	1010	1011	23.70	27.70	4.00		.58	720			
5	5		KITICHEN		17	1011	1012	23.70	27.70	4.00		.58	720			
5	5		KITICHEN		18	1012	1020	21.60	27.50	5.90		1.15	1080			
RUN: 1																
7	7		LEAITCH		14	0931	0932		4.50		.283	.25	542			HOT TEMPERATURE IS ACTUALLY TEMPERATURE DIFFERENCE BETWEEN TUBE AND AMBIENT
7	7		LEAITCH		14	0937	0938		7.95		.283	.166	411			
7	7		LEAITCH		14	0945	0946		7.65		.283	.122	315			
7	7		LEAITCH		14	0953	0960		7.55		.283	.114	212			
7	7		LEAITCH		14	1007	1008		6.10		.283	.104	217			
RUN: 2																
7	7		LEAITCH		14	1033	1034		5.45		.283	.045	163			
7	7		LEAITCH		14	1035	1036		5.50		.283	.036	148			
7	7		LEAITCH		14	1039	1040		5.00		.283	.03	115			
7	7		LEAITCH		14	1041	1042		4.90		.283	.026	109			
7	7		LEAITCH		14	1047	1049		4.45		.283	.023	85			
7	7		LEAITCH		14	1048	1053		4.25		.283	.014	62			
RUN: 1																
8	8		MEEE		17	0945		27.10	28.00	1.90		.15	0			
8	8		MEEE		17	0945		25.00	28.00	3.00		.68	518			
8	8		MEEE		17	0945		24.00	28.00	4.00		.9	637			
8	8		MEEE		17	0945		23.70	28.00	4.30		1.23	293			
8	8		MEEE		17	0945		23.40	28.00	4.60		1.49	210			
8	8		MEEE		17	0945		23.20	28.00	4.80		1.3	303			
8	8		MEEE		17	0945	0955	22.90	28.00	5.70		1.49	203			
8	8		MEEE		17	0945		22.10	28.00	6.00		1.4	420			
RUN: 1																
9	9		GAGIN		5	0945		23.30	27.40	5.10		1.01	100			
9	9		GAGIN		5	0948		23.80	27.00	3.60		.51	230			
9	9		GAGIN		5	0952		24.60	27.50	2.90		.32	150			
9	9		GAGIN		5	0954		25.50	27.60	2.10		.17	110			
RUN: 2																
9	9		GAGIN		5	1059		23.30	27.50	5.20		1.08	950			
9	9		GAGIN		5	1001		23.90	27.00	3.80		.55	210			
9	9		GAGIN		5	1006		24.60	27.40	2.80		.32	120			
9	9		GAGIN		5	1008		25.60	27.40	1.80		.13	140			
RUN: 3																
9	9		GAGIN		5	1015		22.00	27.50	5.50		1.18	880			

EXPERIMENT #: 26

NUCLEI MEASUREMENTS

PROCESSING DATE: 01/09/26

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	COLD	PLATE TEMP HOT (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
9	9	GAGIN		5	1017		23.70	27.40	3.70	1.2	.51	320			
9	9	GAGIN		5	1020		24.70	27.50	2.80	1.2	.32	170			
9	9	GAGIN		5	1024		25.60	27.70	2.10	1.2	.1	110			
RUN: 4															
9	9	GAGIN		5	1038		22.40	27.50	5.10	1.2	1.01	490			
9	9	GAGIN		5	1041		23.70	27.50	3.80	1.2	.58	250			
9	9	GAGIN		5	1044		23.70	27.70	3.00	1.2	.35	110			
9	9	GAGIN		5	1047		25.60	27.60	2.00	1.2	.31	100			
RUN: 1															
10	10	LALA		13	0937		19.82	22.46	2.64	.4	.25	619.8	603.4		
10	10	LALA		13	0939		19.92	22.39	2.47	.4	.25	586.9	603.4		
10	10	LALA		13	0939		19.59	22.50	2.91	.4	.35	815.3	790.6		
10	10	LALA		13	0940		19.68	22.60	2.92	.4	.35	765.3	790.6		
10	10	LALA		13	0941		19.22	22.71	3.49	.4	.5	947.9	1217.7		
10	10	LALA		13	0942		19.35	22.81	3.48	.4	.5	1487.4	1217.7		
10	10	LALA		13	0943		18.65	22.91	4.26	.4	.75	1523.2	1432.8		
10	10	LALA		13	0944		18.72	22.98	4.26	.4	.75	1523.2	1432.8		
10	10	LALA		13	0945		18.19	23.06	4.91	.4	1	1520.7	1273.4		N=1520.7 S**0.60
10	10	LALA		13	0946		18.19	23.10	4.91	.4	1	1520.7	1273.4		
RUN: 2															
10	10	LALA		13	0947		23.50	26.06	2.56	.4	.35	683.7			
10	10	LALA		13	0948		22.45	26.04	3.59	.4	.35	80.9			
10	10	LALA		13	0949		22.45	26.04	3.59	.4	.35	80.9			
10	10	LALA		13	0950		22.45	26.04	3.59	.4	.35	80.9			
10	10	LALA		13	0951		22.45	26.04	3.59	.4	.35	80.9			
10	10	LALA		13	0952		22.45	26.04	3.59	.4	.35	80.9			
10	10	LALA		13	0953		22.45	26.04	3.59	.4	.35	80.9			
10	10	LALA		13	0954		22.45	26.04	3.59	.4	.35	80.9			
10	10	LALA		13	0955		22.45	26.04	3.59	.4	.35	80.9			
10	10	LALA		13	0956		22.45	26.04	3.59	.4	.35	80.9			
RUN: 3															
10	10	LALA		13	1035		24.61	27.23	2.62	.4	.25	778.3	770.7		
10	10	LALA		13	1036		22.45	27.12	3.03	.4	.35	916.5	918		
10	10	LALA		13	1037		24.07	27.05	3.03	.4	.35	1213.66	1189.9		
10	10	LALA		13	1038		23.40	27.01	3.61	.4	.5	1335.1	1358.1		
10	10	LALA		13	1039		23.40	27.01	3.61	.4	.5	1335.1	1358.1		
10	10	LALA		13	1040		22.51	26.94	4.40	.4	.75	1321.3	1358.1		
10	10	LALA		13	1041		22.51	26.94	4.40	.4	.75	1321.3	1358.1		
10	10	LALA		13	1042		21.76	26.82	5.06	.4	1	1475.3	1443.8		N=1521.1 S**0.47
10	10	LALA		13	1043		21.76	26.82	5.06	.4	1	1475.3	1443.8		
RUN: 4															
10	10	LALA		13	1040		24.39	26.97	2.56	.4	.25	820.1	838.2		
10	10	LALA		13	1041		23.91	26.95	2.92	.4	.25	846.2	838.2		
10	10	LALA		13	1042		23.91	26.95	2.92	.4	.25	846.2	838.2		
10	10	LALA		13	1043		23.91	26.95	2.92	.4	.25	846.2	838.2		
10	10	LALA		13	1044		23.91	26.95	2.92	.4	.25	846.2	838.2		
10	10	LALA		13	1045		23.91	26.95	2.92	.4	.25	846.2	838.2		
10	10	LALA		13	1046		23.91	26.95	2.92	.4	.25	846.2	838.2		
10	10	LALA		13	1047		23.91	26.95	2.92	.4	.25	846.2	838.2		
10	10	LALA		13	1048		23.91	26.95	2.92	.4	.25	846.2	838.2		
10	10	LALA		13	1049		23.91	26.95	2.92	.4	.25	846.2	838.2		
RUN: 1															
11	11	FITZGERALD		28	0950		1.5	420			.15	420	420		
11	11	FITZGERALD		28	0950		1.5	420			.024	11.3	1.23		
11	11	FITZGERALD		28	0950		1.5	420			.025	1.62	1.62		
11	11	FITZGERALD		28	0950		1.5	420			.025	1.62	1.62		

EXPERIMENT #: 26

PROCESSING DATE: 01/09/76

NUCLEI MEASUREMENTS

MACH NO.	DUCT NO.	NAM	OBSERVER	NO.	SAMPLE START	SAMPLE END	COLD	HOT	PLATE T	DELTA	VOLUME S	SUPER SAT	NUCLEI COUNT	ORS	EST	REMARKS
11	11	FITZGERALD		28	0950	1008										
RUN: 1																
13	13	AYERS			0943	0945	23.77	25.20		2.41	.0187	.25	693			
13	13	AYERS			0945	0947	23.70	25.20		3.50	.0187	.25	750			
13	13	AYERS			0947	0949	21.05	25.30		3.25	.0187	.25	1075			
13	13	AYERS			0949	0951	21.03	25.30		4.27	.0187	.25	1375			
13	13	AYERS			0951	0952	20.36	25.30		4.84	.0187	.25	1590			
13	13	AYERS			0952	0954	19.84	25.30		5.47	.0187	.25	1545			
13	13	AYERS			0954	0954	19.90	25.40		5.50	.0187	.25	1725			
RUN: 2																
13	13	AYERS			1059	1100	26.40	28.90		3.50	.0187	.25	720			
13	13	AYERS			1100	1103	24.53	28.90		4.37	.0187	.25	1365			
13	13	AYERS			1103	1104	23.73	28.80		4.07	.0187	.25	1725			
13	13	AYERS			1104	1105	23.17	28.80		5.63	.0187	.25	2070			
RUN: 1																
14	14	HUDSON			1001	1030					.0187	.21	168			F=42
14	14	HUDSON			1001	1030					.0187	.21	117.2			F=22
14	14	HUDSON			1001	1030					.0187	.21	62.09			F=22
14	14	HUDSON			1001	1030					.0187	.21	1.34			F=22
14	14	HUDSON			1001	1030					.0187	.21	176.9			F=22
14	14	HUDSON			1001	1030					.0187	.21	120.3			F=22
14	14	HUDSON			1001	1030					.0187	.21	60.8			F=22
14	14	HUDSON			1001	1030					.0187	.21	1.12			F=22
RUN: 2																
14	14	HUDSON			1122	1130					.0187	.21	206.2			F=22
14	14	HUDSON			1122	1130					.0187	.21	171.2			F=22
14	14	HUDSON			1122	1130					.0187	.21	172.2			F=22
14	14	HUDSON			1122	1130					.0187	.21	1.15			F=22
RUN: 1																
15	15	KOCMOND			0939	0940	18.36	25.06		6.70	.0187	.21	1546			MAIN FLOW ABOUT
15	15	KOCMOND			0939	0942	19.49	25.44		8.21	.0187	.21	1900			1570 COUNTS.
15	15	KOCMOND			0942	0952	19.40	25.44		9.24	.0187	.21	1382			DROP SIZE PLAT GOOD
15	15	KOCMOND			0952	1000	19.87	23.81		3.84	.0187	.21	1142			ATTEN COUNT RISING
15	15	KOCMOND			1000	1004	20.63	23.59		2.56	.0187	.21	882			ATTEN 5 PER CENT PER
15	15	KOCMOND			1004	1004					.0187	.21	210			MIN. SLOWED MF
15	15	KOCMOND			1004	1042	21.41	22.71		1.30	.0187	.21				USING FLTS MORNING
RUN: 2																
15	15	ROGERS	FRED		1026	1059	19.74	24.76		6.01	.0187	.21	1837			SECOND RUN AT
15	15	ROGERS	FRED		1059	1107	19.49	24.43		4.37	.0187	.21	1590			HIGHER MAIN FLOW,
15	15	ROGERS	FRED		1107	1159	19.87	23.59		3.25	.0187	.21	1347			ABOUT 230 CNTS.
15	15	ROGERS	FRED		1159	1127	20.82	22.88		2.56	.0187	.21	967			GOOD DROP SIZE PLAT.
15	15	ROGERS	FRED		1127	1127	20.99	22.88		2.21	.0187	.21	364			DROP SIZE PLAT GOING
15	15	ROGERS	FRED		1127	1127	21.35	22.56		1.21	.0187	.21	285			BAD.

EXPERIMENT #: 26

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/09/26

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	PLATE TEMP		COLD	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
							HOT (C)	TEMP (C)								
RUN: 1																
16	16	HUDSON		9	0936	0943	26.05	27.75	21.30	4.75	.0042	1369				
16	16	HUDSON		9	0936	0943	25.40	27.70	1.65	3.20	.0042	1174				
16	16	HUDSON		9	0936	0943	26.05	27.70	2.20	4.75	.0042	1915				
16	16	HUDSON		9	0955	0959	26.05	27.70	2.20	4.75	.0042	1310				
16	16	HUDSON		9	0955	0959	26.05	27.70	2.20	4.75	.0042	1587				
16	16	HUDSON		9	1008	1012	26.05	27.70	2.20	4.75	.0042	1308				
16	16	HUDSON		9	1008	1012	26.05	27.70	2.20	4.75	.0042	1986				
16	16	HUDSON		9	1008	1012	26.05	27.70	2.20	4.75	.0042	571				
RUN: 2																
16	16	HUDSON		9	1018	1022	26.05	27.70	21.30	4.75	.0042	1442				
16	16	HUDSON		9	1018	1022	26.05	27.70	2.20	4.75	.0042	1086				
16	16	HUDSON		9	1018	1022	26.05	27.70	2.20	4.75	.0042	1436				
16	16	HUDSON		9	1035	1038	26.05	27.70	2.20	4.75	.0042	1490				
16	16	HUDSON		9	1035	1038	26.05	27.70	2.20	4.75	.0042	1105				
16	16	HUDSON		9	1045	1049	26.05	27.70	2.20	4.75	.0042	1473				
16	16	HUDSON		9	1045	1049	26.05	27.70	2.20	4.75	.0042	1161				
RUN: 3																
16	16	HUDSON		9	1101	1105	26.05	27.70	21.30	4.75	.0042	1572				
16	16	HUDSON		9	1101	1105	26.05	27.70	2.20	4.75	.0042	1710				
16	16	HUDSON		9	1101	1105	26.05	27.70	2.20	4.75	.0042	1574				
16	16	HUDSON		9	1114	1116	26.05	27.70	2.20	4.75	.0042	1713				
16	16	HUDSON		9	1114	1116	26.05	27.70	2.20	4.75	.0042	1520				
16	16	HUDSON		9	1123	1126	26.05	27.70	2.20	4.75	.0042	1700				
16	16	HUDSON		9	1123	1126	26.05	27.70	2.20	4.75	.0042	1544				
16	16	HUDSON		9	1130	1132	26.05	27.70	21.30	4.75	.0042	1725				
16	16	HUDSON		9	1130	1132	26.05	27.70	2.20	4.75	.0042	1544				
RUN: 4																
17	17	WOJCIECHOWSKI		27	0944	0945	26.05	27.70	21.30	4.75	.0042	720	720			
17	17	WOJCIECHOWSKI		27	0949	0950	26.05	27.70	2.20	4.75	.0042	804.6	804.6			
17	17	WOJCIECHOWSKI		27	0951	0952	26.05	27.70	2.20	4.75	.0042	1213.6	1213.6			
17	17	WOJCIECHOWSKI		27	0953	0954	26.05	27.70	2.20	4.75	.0042	1167.6	1167.6			
RUN: 1																
18	18	HUDSON		9	0936	0943	26.05	27.70	21.30	4.75	.0042	1537				
18	18	HUDSON		9	1008	1012	26.05	27.70	2.20	4.75	.0042	1303				
18	18	HUDSON		9	1018	1022	26.05	27.70	2.20	4.75	.0042	1977				
18	18	HUDSON		9	1035	1038	26.05	27.70	2.20	4.75	.0042	571				
18	18	HUDSON		9	1045	1049	26.05	27.70	2.20	4.75	.0042	1300				
18	18	HUDSON		9	1101	1105	26.05	27.70	2.20	4.75	.0042	1227				
18	18	HUDSON		9	1123	1126	26.05	27.70	2.20	4.75	.0042	1167				
18	18	HUDSON		9	1130	1132	26.05	27.70	2.20	4.75	.0042	1149				
RUN: 1																
21	21	TRUEBLOOD		26	0938	0939	25.00	25.00	20.00	5.00	.0042	1488				
21	21	TRUEBLOOD		26	0943	0944	25.00	25.00	21.75	3.25	.0042	1886				

EXPERIMENT #1 26

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/09/26

MACH NO.	DUCT NO.	OBSERVER	NAME	NO.	SAMPLE START	TIME END	PLATE TEMP		COLD	HOT	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
							(C)	(C)									
21	21	TRUEBLOOD	26	0949	0949	25.00	25.00	22.44	25.00	2.56	530.8	.62	530.8				
21	21	TRUEBLOOD	26	0959	0959	25.00	25.00	25.00	25.00	0.00	150.7	.14	150.7				
21	21	TRUEBLOOD	26	0959	0959	25.00	25.00	25.00	25.00	0.00	194.3	.08	194.3				
21	21	TRUEBLOOD	26	1006	1006	25.00	25.00	25.00	25.00	0.00	4.77	.35	4.77				
21	21	TRUEBLOOD	26	1006	1006	25.00	25.00	25.00	25.00	0.00	4.11	.22	4.11				
RUN: 2																	
21	21	TRUEBLOOD	26	1025	1025	25.00	25.00	19.88	25.00	5.12	181.7	.49	181.7				
21	21	TRUEBLOOD	26	1037	1037	25.00	25.00	25.00	25.00	0.00	253.7	.25	253.7				
21	21	TRUEBLOOD	26	1046	1047	25.00	25.00	25.00	25.00	0.00	167.4	.14	167.4				
21	21	TRUEBLOOD	26	1046	1047	25.00	25.00	25.00	25.00	0.00	139.5	.09	139.5				
21	21	TRUEBLOOD	26	1053	1054	25.00	25.00	25.00	25.00	0.00	5.85	.33	5.85				
21	21	TRUEBLOOD	26	1053	1054	25.00	25.00	25.00	25.00	0.00	5.65	.25	5.65				
RUN: 1																	
24	24	SERPOLAY	25	1000	1000	27.70	27.70	27.80	27.80	1.10	11.0	.1	11.0				
24	24	SERPOLAY	25	1019	1019	27.40	27.40	27.80	27.80	1.50	11.0	.2	11.0				
24	24	SERPOLAY	25	1023	1023	27.10	27.10	27.80	27.80	1.50	15.0	.75	15.0				
24	24	SERPOLAY	25	1050	1050	27.60	27.60	27.70	27.70	0.10	15.0	.69	15.0				
24	24	SERPOLAY	25	1050	1050	27.70	27.70	27.80	27.80	0.10	15.0	.59	15.0				
24	24	SERPOLAY	25	1053	1053	27.80	27.80	27.80	27.80	1.10	11.0	.1	11.0				
RUN: 1																	
25	25	POXY	25	1059	1059	26.10	26.10	21.70	26.10	4.40	42.4	.22	42.4			.54 VOL SAMP X 10 ^{0.5}	
25	25	POXY	25	1059	1059	26.50	26.50	22.50	26.50	4.00	25.3	.25	25.3				
25	25	POXY	25	1059	1059	26.50	26.50	22.50	26.50	4.30	25.9	.25	25.9				
25	25	POXY	25	1059	1059	26.50	26.50	22.50	26.50	4.30	25.9	.25	25.9				
RUN: 1																	
26	26	HINDMAN	6	0947	1032	27.00	27.00	27.00	27.00	0.00	90	.15	90			7.9	
26	26	HINDMAN	6	0947	1032	27.00	27.00	27.00	27.00	0.00	31	.4	31			1.4	
26	26	HINDMAN	6	0947	1032	27.00	27.00	27.00	27.00	0.00	11	.2	11			0.62	
26	26	HINDMAN	6	0947	1032	27.00	27.00	27.00	27.00	0.00	10	.1	10			0.013	
RUN: 1																	
27	27	ORTAKE	19	1050	1050	105.0	105.0	105.0	105.0	0.00	6.3		6.3			75% R.H. NOMINAL	
27	27	ORTAKE	19	1050	1050	105.0	105.0	105.0	105.0	0.00	2.5		2.5			106% R.H. NOMINAL	
27	27	ORTAKE	19	1050	1050	105.0	105.0	105.0	105.0	0.00	12.5		12.5			125% R.H. NOMINAL	

80/10/15

EXPERIMENT # 27

PURPOSE

INSTRUMENT COMPARISON WITH INSOLUBLE, WETTABLE AEROSOL.

DESCRIPTION OF EXPERIMENT

DATE TIME
 START END

15 OCT, 1980 1151 1415

NUCLEI TYPE

AGI

GENERATION METHOD

THERMAL, N2 CARRIER GAS, HIGH DILUTION.

SIZE DISTRIBUTION SHAPING

NA

REMARKS

SLOW UPWARD DRIFT OF TOTAL NUMBER CONC.

WEATHER SYNOPSIS

NONE

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS

OUTSIDE CONDITIONS

AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG WS (M/S)	AVG WD (DEG)
25.0	22.0	844					

EXPERIMENT #: 27

PROCESSING DATE: 01/09/76

MACH NO.	DUCT NO.	OBSERVER	NO.	SAMPLE START	SAMPLE END	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	ORS N	EST K	REMARKS
						COLD	HOT (C)							
RUN: 1														
2	2	POLITOVICH	20	1316	1317	23.10	25.20	2.10	.17	.23		15		
RUN: 2														
5	5	KITCHEN	12	1245	1245	23.70	27.50	3.80	.6	.77	77			
5	5	KITCHEN	12	1246	1246	23.70	27.50	3.80	.7	.77	77			
5	5	KITCHEN	12	1247	1247	23.70	27.50	3.80	1.6	.77	77			
5	5	KITCHEN	12	1248	1248	23.70	27.50	3.80	1.6	.77	77			
5	5	KITCHEN	12	1249	1249	23.70	27.50	3.80	.5	.77	77			
5	5	KITCHEN	12	1250	1250	23.70	27.50	3.80	.5	.77	77			
5	5	KITCHEN	12	1251	1251	23.70	27.50	3.80	.5	.77	77			
RUN: 3														
7	7	LEAITCH	14	1313	1314	7.35	7.35		.22	.283	283	73		
7	7	LEAITCH	14	1314	1315	7.35	7.35		.22	.283	283	1		
7	7	LEAITCH	14	1315	1316	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1316	1317	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1317	1318	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1318	1319	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1319	1320	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1320	1321	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1321	1322	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1322	1323	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1323	1324	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1324	1325	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1325	1326	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1326	1327	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1327	1328	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1328	1329	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1329	1330	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1330	1331	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1331	1332	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1332	1333	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1333	1334	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1334	1335	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1335	1336	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1336	1337	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1337	1338	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1338	1339	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1339	1340	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1340	1341	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1341	1342	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1342	1343	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1343	1344	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1344	1345	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1345	1346	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1346	1347	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1347	1348	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1348	1349	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1349	1350	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1350	1351	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1351	1352	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1352	1353	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1353	1354	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1354	1355	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1355	1356	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1356	1357	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1357	1358	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1358	1359	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1359	1360	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1360	1361	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1361	1362	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1362	1363	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1363	1364	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1364	1365	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1365	1366	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1366	1367	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1367	1368	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1368	1369	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1369	1370	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1370	1371	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1371	1372	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1372	1373	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1373	1374	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1374	1375	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1375	1376	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1376	1377	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1377	1378	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1378	1379	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1379	1380	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1380	1381	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1381	1382	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1382	1383	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1383	1384	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1384	1385	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1385	1386	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1386	1387	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1387	1388	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1388	1389	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1389	1390	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1390	1391	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1391	1392	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1392	1393	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1393	1394	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1394	1395	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1395	1396	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1396	1397	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1397	1398	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1398	1399	7.35	7.35		.15	.283	283	4		
7	7	LEAITCH	14	1399	1400	7.35	7.35		.15	.283	283	4		

PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

EXPERIMENT #: 27

MACH NO.	DUCT NO.	NAME	OSERVER	NO.	SAMPLE START	TIME END	COLD	HOT (C)	PLATE TEMP	DELTA	VOLUME SAMPLED (L)	SUPER SAT (S)	NUCLEI COUNT	OBS N	EST K	REMARKS
10	10	LALA		13	1215		24.610	27.113	27.113	2.57	4	.25	19.6	147.8		
10	10	LALA		13	1216		24.033	27.033	27.033	3.03	4	.35	18.8			
10	10	LALA		13	1217		23.332	26.992	26.992	3.60	4	.35	55.7	274.3		
10	10	LALA		13	1219		22.325	26.85	26.85	3.60	4	.75	48.3	518.5		
10	10	LALA		13	1221		22.38	26.77	26.77	4.39	4	.75	820	856.5		
10	10	LALA		13	1222		21.56	26.61	26.61	5.05	4	1	1125.3	1075.5		
10	10	LALA		13	1225		23.26	26.53	26.53	2.79	4	.3	125.6			
10	10	LALA		13	1226		22.40	26.40	26.40	3.92	4	.5	227			
10	10	LALA		13	1227		22.40	26.32	26.32	3.92	4	.5	518			
10	10	LALA		13	1228		21.94	26.24	26.24	4.21	4	.8	657			
10	10	LALA		13	1230		21.90	26.15	26.15	4.51	4	.9	761.1			
10	10	LALA		13	1231		20.37	26.08	26.08	4.78	4	1	922.1			
10	10	LALA		13	1232		19.77	26.00	26.00	5.02	4	1.2	961.6			
10	10	LALA		13	1233		18.77	25.81	25.81	7.04	4	1.2	1471.4			
10	10	LALA		13	1235		22.64	25.99	25.99	2.77	4	.3	97			
10	10	LALA		13	1236		22.64	25.91	25.91	3.20	4	.4	186.7			
10	10	LALA		13	1237		22.64	25.81	25.81	3.57	4	.5	383.5			
10	10	LALA		13	1238		22.64	25.71	25.71	4.20	4	.7	575.3			
10	10	LALA		13	1239		22.64	25.63	25.63	4.75	4	.9	725.9			
10	10	LALA		13	1240		22.64	25.57	25.57	5.09	4	1.2	1092.7			
10	10	LALA		13	1241		21.8	25.47	25.47	7.00	4	1.2	1237			
10	10	LALA		13	1242		21.8	25.37	25.37	7.00	4	1.2	1092.7			
10	10	LALA		13	1243		21.8	25.27	25.27	7.00	4	1.2	1237			
10	10	LALA		13	1244		21.8	25.17	25.17	7.00	4	1.2	1092.7			
10	10	LALA		13	1245		21.8	25.07	25.07	7.00	4	1.2	1237			
10	10	LALA		13	1246		21.8	24.97	24.97	7.00	4	1.2	1092.7			
10	10	LALA		13	1247		21.8	24.87	24.87	7.00	4	1.2	1237			
10	10	LALA		13	1248		21.8	24.77	24.77	7.00	4	1.2	1092.7			
10	10	LALA		13	1249		21.8	24.67	24.67	7.00	4	1.2	1237			
10	10	LALA		13	1250		21.8	24.57	24.57	7.00	4	1.2	1092.7			
10	10	LALA		13	1251		21.8	24.47	24.47	7.00	4	1.2	1237			
10	10	LALA		13	1252		21.8	24.37	24.37	7.00	4	1.2	1092.7			
10	10	LALA		13	1253		21.8	24.27	24.27	7.00	4	1.2	1237			
10	10	LALA		13	1254		21.8	24.17	24.17	7.00	4	1.2	1092.7			
10	10	LALA		13	1255		21.8	24.07	24.07	7.00	4	1.2	1237			
10	10	LALA		13	1256		21.8	23.97	23.97	7.00	4	1.2	1092.7			
10	10	LALA		13	1257		21.8	23.87	23.87	7.00	4	1.2	1237			
10	10	LALA		13	1258		21.8	23.77	23.77	7.00	4	1.2	1092.7			
10	10	LALA		13	1259		21.8	23.67	23.67	7.00	4	1.2	1237			
10	10	LALA		13	1260		21.8	23.57	23.57	7.00	4	1.2	1092.7			
10	10	LALA		13	1261		21.8	23.47	23.47	7.00	4	1.2	1237			
10	10	LALA		13	1262		21.8	23.37	23.37	7.00	4	1.2	1092.7			
10	10	LALA		13	1263		21.8	23.27	23.27	7.00	4	1.2	1237			
10	10	LALA		13	1264		21.8	23.17	23.17	7.00	4	1.2	1092.7			
10	10	LALA		13	1265		21.8	23.07	23.07	7.00	4	1.2	1237			
10	10	LALA		13	1266		21.8	22.97	22.97	7.00	4	1.2	1092.7			
10	10	LALA		13	1267		21.8	22.87	22.87	7.00	4	1.2	1237			
10	10	LALA		13	1268		21.8	22.77	22.77	7.00	4	1.2	1092.7			
10	10	LALA		13	1269		21.8	22.67	22.67	7.00	4	1.2	1237			
10	10	LALA		13	1270		21.8	22.57	22.57	7.00	4	1.2	1092.7			
10	10	LALA		13	1271		21.8	22.47	22.47	7.00	4	1.2	1237			
10	10	LALA		13	1272		21.8	22.37	22.37	7.00	4	1.2	1092.7			
10	10	LALA		13	1273		21.8	22.27	22.27	7.00	4	1.2	1237			
10	10	LALA		13	1274		21.8	22.17	22.17	7.00	4	1.2	1092.7			
10	10	LALA		13	1275		21.8	22.07	22.07	7.00	4	1.2	1237			
10	10	LALA		13	1276		21.8	21.97	21.97	7.00	4	1.2	1092.7			
10	10	LALA		13	1277		21.8	21.87	21.87	7.00	4	1.2	1237			
10	10	LALA		13	1278		21.8	21.77	21.77	7.00	4	1.2	1092.7			
10	10	LALA		13	1279		21.8	21.67	21.67	7.00	4	1.2	1237			
10	10	LALA		13	1280		21.8	21.57	21.57	7.00	4	1.2	1092.7			
10	10	LALA		13	1281		21.8	21.47	21.47	7.00	4	1.2	1237			
10	10	LALA		13	1282		21.8	21.37	21.37	7.00	4	1.2	1092.7			
10	10	LALA		13	1283		21.8	21.27	21.27	7.00	4	1.2	1237			
10	10	LALA		13	1284		21.8	21.17	21.17	7.00	4	1.2	1092.7			
10	10	LALA		13	1285		21.8	21.07	21.07	7.00	4	1.2	1237			
10	10	LALA		13	1286		21.8	20.97	20.97	7.00	4	1.2	1092.7			
10	10	LALA		13	1287		21.8	20.87	20.87	7.00	4	1.2	1237			
10	10	LALA		13	1288		21.8	20.77	20.77	7.00	4	1.2	1092.7			
10	10	LALA		13	1289		21.8	20.67	20.67	7.00	4	1.2	1237			
10	10	LALA		13	1290		21.8	20.57	20.57	7.00	4	1.2	1092.7			
10	10	LALA		13	1291		21.8	20.47	20.47	7.00	4	1.2	1237			
10	10	LALA		13	1292		21.8	20.37	20.37	7.00	4	1.2	1092.7			
10	10	LALA		13	1293		21.8	20.27	20.27	7.00	4	1.2	1237			
10	10	LALA		13	1294		21.8	20.17	20.17	7.00	4	1.2	1092.7			
10	10	LALA		13	1295		21.8	20.07	20.07	7.00	4	1.2	1237			
10	10	LALA		13	1296		21.8	19.97	19.97	7.00	4	1.2	1092.7			
10	10	LALA		13	1297		21.8	19.87	19.87	7.00	4	1.2	1237			
10	10	LALA		13	1298		21.8	19.77	19.77	7.00	4	1.2	1092.7			
10	10	LALA		13	1299		21.8	19.67	19.67	7.00	4	1.2	1237			
10	10	LALA		13	1300		21.8	19.57	19.57	7.00	4	1.2	1092.7			
10	10	LALA		13	1301		21.8	19.47	19.47	7.00	4	1.2	1237			
10	10	LALA		13	1302		21.8	19.37	19.37	7.00	4	1.2	1092.7			
10	10	LALA		13	1303		21.8	19.27	19.27	7.00	4	1.2	1237			
10	10	LALA		13	1304		21.8	19.17	19.17	7.00	4	1.2	1092.7			
10	10	LALA		13	1305		21.8	19.07	19.07	7.00	4					

EXPERIMENT #: 27

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/09/26

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	COLD	HOT (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
RUN: 1															
14	14	HUDSON		9	1306	1306				.023	.11	.071			
14	14	HUDSON		9	1306	1306				.023	.003				
14	14	HUDSON		9	1307	1309				.021	.029	1.076			
14	14	HUDSON		9	1307	1309				.021	.11	.124			
14	14	HUDSON		9	1307	1309				.021	.029	.0016			
RUN: 2															
14	14	HUDSON		9	1340	1411				.028	.21	.806			
14	14	HUDSON		9	1340	1411				.028	.11	.134			
14	14	HUDSON		9	1340	1411				.028	.11	.064			
14	14	HUDSON		9	1340	1411				.028	.029	.0088			
RUN: 1															
15	15	ROGERS	FRED	23	1226	1226	18.36	25.06	6.70		.06	1367			MAIN FLOW ABOUT 2300 USING THIS MORNING'S BACKGROUND USING SMALLER CAPILLARY GOOD DROP SIZE PLATS
15	15	ROGERS	FRED	23	1230	1230	18.74	24.75	6.03		.05	1241			
15	15	ROGERS	FRED	23	1237	1241	19.19	24.44	5.25		.11	636			
15	15	ROGERS	FRED	23	1240	1245	19.87	23.61	3.74		.03	397			
15	15	ROGERS	FRED	23	1248	1249	20.25	23.19	2.94		.025	59			
15	15	ROGERS	FRED	23	1253	1254	20.69	22.88	2.19		.021	15			SIZE PLAT GOING RAD. MAIN FLOW ART. 2300.
RUN: 2															
15	15	ROGERS	FRED	23	1334	1334	18.36	25.06	6.70		.06	1711			SECOND RUN AT LOWER MAIN FLOW, ART. 1800
15	15	ROGERS	FRED	23	1338	1338	18.74	24.75	6.03		.05	1538			AVGAGING LAST 4 COUNTS PER SEC. USING SMALLER CAPILLARY.
15	15	ROGERS	FRED	23	1346	1346	19.19	24.44	5.25		.11	543			
15	15	ROGERS	FRED	23	1357	1357	19.87	23.61	3.74		.03	236			
15	15	ROGERS	FRED	23	1357	1401	20.25	23.19	2.94		.025	19			
15	15	ROGERS	FRED	23	1400	1401	20.69	22.88	2.19		.021				
RUN: 1															
16	16	HUDSON		9	1200	1200	25.5	0	25.5	.024	.91	1205			
16	16	HUDSON		9	1200	1200	25.5	0	25.5	.024	.28	875			
16	16	HUDSON		9	1200	1200	25.5	0	25.5	.024	.91	1080			
16	16	HUDSON		9	1200	1200	25.5	0	25.5	.024	.28	952			
16	16	HUDSON		9	1200	1200	25.5	0	25.5	.024	.68	438			
16	16	HUDSON		9	1200	1200	25.5	0	25.5	.024	.28	51			
RUN: 2															
16	16	HUDSON		9	1250	1251	25.5	0	25.5	.024	.91	953			
16	16	HUDSON		9	1251	1251	25.5	0	25.5	.024	.28	406			
16	16	HUDSON		9	1251	1251	25.5	0	25.5	.024	.91	399			
16	16	HUDSON		9	1251	1251	25.5	0	25.5	.024	.28	421			
16	16	HUDSON		9	1303	1303	25.5	0	25.5	.024	.91	339			
16	16	HUDSON		9	1303	1303	25.5	0	25.5	.024	.28	329			
RUN: 3															
16	16	HUDSON		9	1310	1312	25.30	0.05	25.35	.018	.91	871			
16	16	HUDSON		9	1312	1312	25.65	0.05	25.70	.018	.28	404			
16	16	HUDSON		9	1321	1321	25.30	0.05	25.35	.018	.91	398			

PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

EXPERIMENT #: 27

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
							COLD	HOT (C)							
16	16	HUDSON		9	1329	1331	21.65	22.40	3.75	.0018	366				
16	16	HUDSON		9	1329	1331	21.65	22.40	3.75	.0018	337				
16	16	HUDSON		9	1343	1349	21.30	22.05	4.75	.0036	898				
16	16	HUDSON		9	1343	1349	21.05	22.40	3.75	.0036	338				
RUN: 4															
16	16	HUDSON		9	1358	1402	21.30	26.00	4.70	.0024	997				
16	16	HUDSON		9	1358	1402	21.30	26.00	4.70	.0024	360				
16	16	HUDSON		9	1358	1402	21.30	26.00	4.70	.0024	348				
16	16	HUDSON		9	1411	1412	21.05	22.80	3.75	.0012	481				
16	16	HUDSON		9	1411	1412	21.10	22.80	2.70	.0012	251				
RUN: 1															
17	17	WOJCIECHOWSKI		27	1235	1236						19.7			
17	17	WOJCIECHOWSKI		27	1235	1236						19.5			
17	17	WOJCIECHOWSKI		27	1235	1236						187.5			
17	17	WOJCIECHOWSKI		27	1235	1236						202.2			
RUN: 2															
17	17	WOJCIECHOWSKI		27	1328	1329						0			
17	17	WOJCIECHOWSKI		27	1332	1333						0			
17	17	WOJCIECHOWSKI		27	1332	1333						74			
17	17	WOJCIECHOWSKI		27	1334	1335						152.9			
RUN: 1															
18	18	HUDSON		9	1235	1235	15	22.55	5.45	.0024	1707				
18	18	HUDSON		9	1243	1243	22.90	22.55	3.65	.0018	1086				
18	18	HUDSON		9	1255	1255	22.80	22.55	3.75	.0018	3304				
18	18	HUDSON		9	1310	1312	22.25	22.85	3.65	.0018	3385				
18	18	HUDSON		9	1329	1331	22.35	22.15	3.65	.0018	3228				
18	18	HUDSON		9	1358	1402	22.70	22.80	3.80	.0024	1463				
RUN: 2															
18	18	HUDSON		9	1411	1412	21.50	26.00	6.50	.0012	2530				
RUN: 1															
21	21	TRUEBLOOD		26	1235	1236	19.98	26.00	5.02	1.01	1622				
21	21	TRUEBLOOD		26	1235	1236	19.60	26.00	4.47	1.77	1981				
21	21	TRUEBLOOD		26	1235	1236	22.00	26.00	3.08	.36	250				
21	21	TRUEBLOOD		26	1238	1239	22.92	26.00	2.08	.173	1.39				
RUN: 2															
21	21	TRUEBLOOD		26	1300	1301	19.76	26.00	5.24	1.009	1522				
21	21	TRUEBLOOD		26	1308	1309	20.77	26.00	4.63	.877	884				
21	21	TRUEBLOOD		26	1312	1312	20.75	26.00	4.26	.733	545				
21	21	TRUEBLOOD		26	1316	1317	21.48	26.00	3.52	.466	119				
21	21	TRUEBLOOD		26	1324	1325	22.90	26.00	2.45	.325	2916				
RUN: 1															
24	24	SEKPOLAY		25	1245	1250	26.80	24.30	1.50	.77	0				

EXPERIMENT #: 27

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/09/26

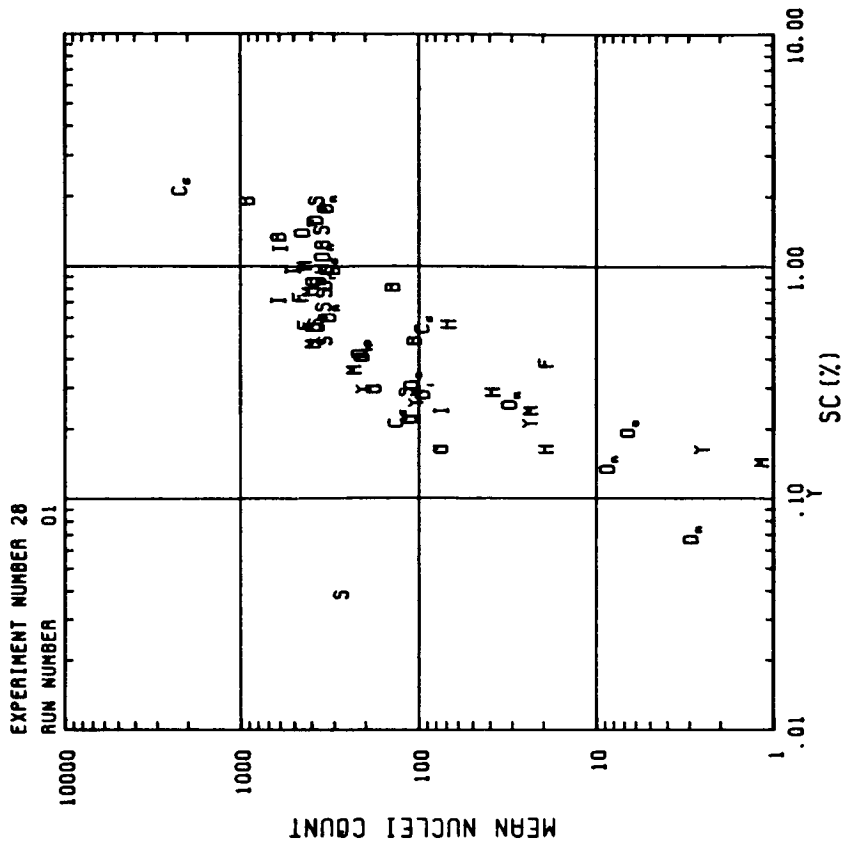
MACH NO.	DUCT NO.	NAME	OBSERVER	N.O.	SAMPLE START	TIME END	COLD	HOT	PLATE TEMP (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
24	24	SERPOLAY		25	1310	1310	26.90	30.40	3.50	.77	.45	110				
24	24	SERPOLAY		25	1315	1318	26.90	30.90	4.00	.77	.58	630				
24	24	SERPOLAY		25	1325	1330	27.10	31.70	4.60	.77	.75	870				
24	24	SERPOLAY		25	1340	1347	27.30	31.30	4.00	.77	.57	540				
24	24	SERPOLAY		25	1355	1400	27.40	30.40	3.00	.77	.33	40				
24	24	SERPOLAY		25	1402	1404	27.40	30.70	3.30	.77	.41	40				
RUN: 1																
25	25	ROYS		3	1244	1255	23.20	25.60	2.40	10.6	.23	57		.82		VOL SAMP X 10**5
25	25	ROYS		3	1249	1312	19.70	25.00	3.70	10.6	.57	141				
25	25	ROYS		3	1315	1326	17.30	24.70	5.00	4.24	1.19	1733				
25	25	ROYS		3	1321	1349	18.00	23.90	5.90	5.3	1.49	622				
25	25	ROYS		3	1346	1400	19.60	23.90	4.30	8.48	.76	124				
25	25	ROYS		3	1412	1415	20.90	23.90	3.00	10.6	.37	171				
RUN: 1																
26	26	HINDMAN		6	1226	1243	26.00	26.00	0.00	1.4	.15	28				
26	26	HINDMAN		6	1226	1243	26.00	26.00	0.00	1.4	.11	27				
26	26	HINDMAN		6	1226	1243	26.00	26.00	0.00	1.4	.07	50				
26	26	HINDMAN		6	1226	1243	26.00	26.00	0.00	1.4	.016	0				
RUN: 2																
26	26	HINDMAN		6	1314	1331	26.00	26.00	0.00	1.4	.15	62				
26	26	HINDMAN		6	1314	1331	26.00	26.00	0.00	1.4	.11	31				
26	26	HINDMAN		6	1314	1331	26.00	26.00	0.00	1.4	.07	12				
26	26	HINDMAN		6	1314	1331	26.00	26.00	0.00	1.4	.016	0				
RUN: 1																
27	27	OHTAKE		19	1321	1321				1.83						
27	27	OHTAKE		19	1321	1321				1.83						
27	27	OHTAKE		19	1321	1321				1.83						

100% NOMINAL FROM
DARK PICTURE

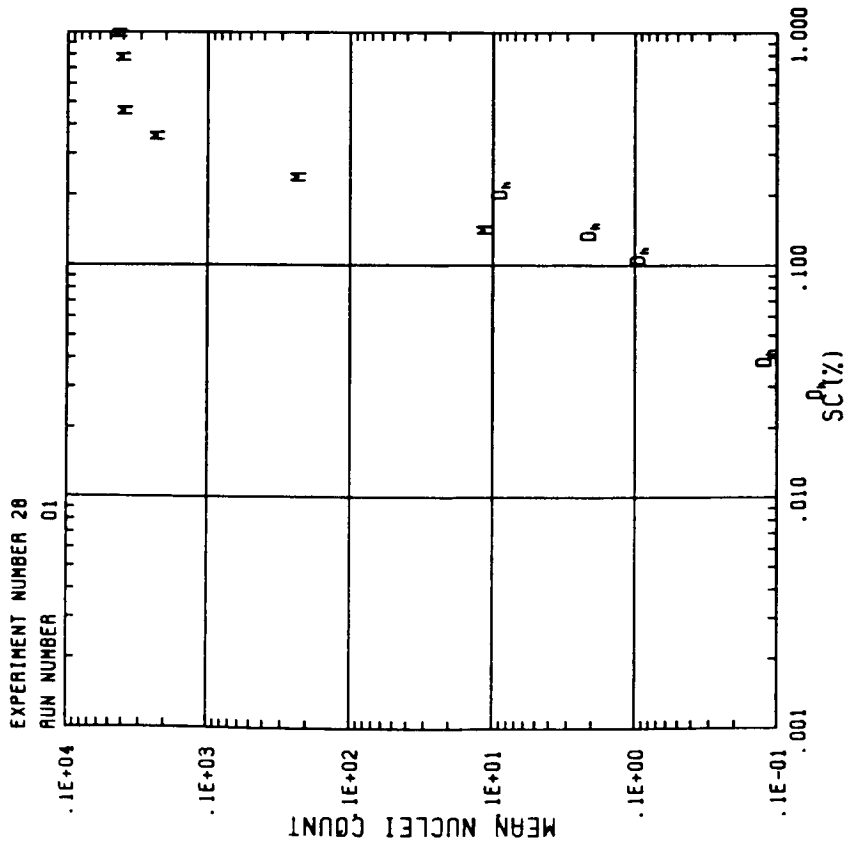
3.25

1.83

INSTRUMENT COMPARISON



INSTRUMENT COMPARISON



SDC/CFD RESULTS

HAZE CHAMBER RESULTS

80/10/15

EXPERIMENT # 2A

PURPOSE

INSOLUBLE MONODISPERSE AEROSOL TO COMPARE PLATEAUX.

DESCRIPTION OF EXPERIMENT

DATE TIME
 START END
15 OCT, 1980 1535 1642

NUCLEI TYPE

AGI

GENERATION METHOD

THERMAL, N2 CARRIER, TA BOAT.

SIZE DISTRIBUTION SHAPING

EC, 4357V, 6 L/MIN. AEROSOL FLOW.

REMARKS

QUASIPERIODIC FLUCTUATIONS +/- 10%, APPROX. 7 MIN. PERIOD.

WEATHER SYNOPSIS

NONE

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS		OUTSIDE CONDITIONS					
AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG WS (M/S)	AVG WD (DEG)
25.0	22.0	847					

PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

EXPERIMENT #: 28

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	SAMPLE END	COLD	HOT	PLATT TEMP	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
RUN: 1																
2	2	ROGERS	DAVE	22	1600	1603	22.40	24.50	2.10			.17		78		
2	2	ROGERS	DAVE	22	1602	1605	22.60	24.80	2.20			.31		186		
RUN: 1																
5	5	KITCHEN		12	1525	1525	15.50	21.10	5.60			1.4	637			
5	5	KITCHEN		12	1526	1526	17.60	21.10	3.50			.5	110			
5	5	KITCHEN		12	1527	1527	17.60	21.10	3.50			.85	111			
5	5	KITCHEN		12	1528	1528	16.60	21.10	4.50			.85	147			
5	5	KITCHEN		12	1529	1529	15.80	21.10	5.30			1.3	370			
5	5	KITCHEN		12	1535	1535	15.80	21.10	5.30			1.2	353			
5	5	KITCHEN		12	1548	1548	14.30	21.10	6.80				956			
RUN: 1																
7	7	LEAITCH		14	1546	1547		8.78			.283	.31		213		HOT TEMPERATURE IS ACTUALLY DIFFERENCE BETWEEN TUBE TEMP. AND AMBIENT INITIAL RELATIVE HUMIDITY IS 71%
7	7	LEAITCH		14	1548	1549		9.55			.283	.27		107		
7	7	LEAITCH		14	1553	1554		7.45			.283	.17		2.6		
7	7	LEAITCH		14	1604	1605		8.85			.283	.11		24.4		
7	7	LEAITCH		14	1622	1623		8.15			.283	.22				
RUN: 1																
8	8	ME		17	1535	1535	23.30	26.30	4.00			.65	245			
8	8	ME		17	1536	1536	25.70	26.30	4.60			.89	445			
8	8	ME		17	1537	1537	25.70	26.30	4.60			1.1	1363			
8	8	ME		17	1538	1538	25.70	26.30	4.60			1.1	1302			
8	8	ME		17	1539	1539	25.70	26.30	4.60			1.1	1392			
8	8	ME		17	1535	1535	19.10	26.30	7.20			2.2	1500			
RUN: 1																
9	9	GAGIN		5	1547	1547	21.70	26.70	5.00			1.02	360			
9	9	GAGIN		5	1548	1548	23.20	27.00	3.80			.17	40			
9	9	GAGIN		5	1549	1549	22.50	27.30	2.80				20			
RUN: 2																
9	9	GAGIN		5	1558	1558	21.70	26.80	5.10			1.05	540			
9	9	GAGIN		5	1600	1600	23.00	27.30	4.30			.33	50			
9	9	GAGIN		5	1607	1607	24.40	27.20	2.80			.11	20			
9	9	GAGIN		5	1607	1607	25.10	27.60	2.50				20			
RUN: 3																
9	9	GAGIN		5	1610	1610	21.40	27.20	5.80			1.35	360			
9	9	GAGIN		5	1612	1612	22.60	27.60	4.60			.85	300			
9	9	GAGIN		5	1614	1614	23.30	26.90	3.60			.5	140			
RUN: 4																
9	9	GAGIN		5	1623	1623	22.60	27.00	4.40			.76	450			
9	9	GAGIN		5	1625	1625	23.30	27.30	4.00			.62	40			
9	9	GAGIN		5	1628	1628	23.80	27.20	3.40			.46	30			
9	9	GAGIN		5	1630	1630	25.20	27.30	2.10			.17	20			
RUN: 1																
10	10	LALA		13	1535	1535	21.48	24.22	2.74			.3	122.7			

PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

EXPERIMENT #: 28

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	COLD	PLATE TEMP (C)	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
RUN: 2															
10	10	LALA		13	1537	1542	21.04	24.20	3.16		.05	280.3			
10	10	LALA		13	1538	1543	20.64	21.17	3.56		.07	350.5			
10	10	LALA		13	1539	1544	20.96	24.14	3.56		.07	355.5			
10	10	LALA		13	1540	1545	19.67	24.09	4.44		.09	349.3			
10	10	LALA		13	1541	1546	19.37	24.07	4.44		.09	316.3			
10	10	LALA		13	1542	1547	19.09	24.05	4.44		.09	316.3			
10	10	LALA		13	1543	1548	17.06	22.96	6.03		1.52	387.1			
RUN: 2															
10	10	LALA		13	1545	1550	21.42	24.16	2.74		.03	128.5			
10	10	LALA		13	1546	1551	20.02	21.18	3.16		.05	323.3			
10	10	LALA		13	1547	1552	20.64	24.17	3.53		.05	344.5			
10	10	LALA		13	1548	1553	20.31	24.16	3.53		.07	382.1			
10	10	LALA		13	1549	1554	19.99	24.15	4.01		.08	404.6			
10	10	LALA		13	1550	1555	19.71	24.14	4.44		.09	403.1			
10	10	LALA		13	1551	1556	19.43	24.12	4.44		.09	348.8			
10	10	LALA		13	1552	1557	18.05	24.08	6.03		1.52	383.8			
10	10	LALA		13	1553	1558	17.14	24.08	6.03		1.52	311.1			
RUN: 1															
13	13	AYESES		2	1536	1537	25.43	27.90	2.74		.25	77			
13	13	AYESES		2	1537	1538	24.23	27.90	3.16		.25	393			
13	13	AYESES		2	1538	1539	23.73	27.90	3.53		.25	420			
13	13	AYESES		2	1539	1540	23.47	27.90	4.01		.25	457			
13	13	AYESES		2	1540	1541	23.21	27.90	4.44		.25	570			
13	13	AYESES		2	1541	1542	21.94	27.90	5.55		.25	600			
RUN: 2															
13	13	AYESES		2	1547	1548	21.80	27.40	5.60		1.25	540			
13	13	AYESES		2	1548	1549	22.69	27.30	5.55		1.25	750			
13	13	AYESES		2	1549	1550	22.66	27.30	5.55		1.25	450			
13	13	AYESES		2	1550	1551	22.61	27.20	5.55		1.25	332			
13	13	AYESES		2	1551	1552	22.47	27.20	5.55		1.25	326			
RUN: 1															
14	14	HUDSON		9	1535	1537	18.74	25.97	6.70		.21	934			
14	14	HUDSON		9	1535	1537	19.11	24.43	6.01		.21	922			
14	14	HUDSON		9	1535	1537	19.87	24.43	5.33		.21	999			
14	14	HUDSON		9	1535	1537	20.65	23.91	4.44		.21	128			
14	14	HUDSON		9	1535	1537	21.35	22.86	3.53		.21	1056			
RUN: 1															
15	15	KOCKMOND		30	1537	1538	18.36	25.97	6.70		.86	348			MAIN FLOW ABT. 1820.
15	15	KOCKMOND		30	1545	1546	19.11	24.43	6.01		1.15	466			AEROSOL FLUCTUATIONS
15	15	KOCKMOND		30	1549	1550	19.87	24.43	5.33		1.15	352			
15	15	KOCKMOND		30	1552	1553	20.65	23.91	4.44		.86	352			
15	15	KOCKMOND		30	1600	1601	20.63	23.91	4.44		.86	321			
15	15	KOCKMOND		30	1606	1607	21.35	22.86	3.53		1.07	329			
15	15	KOCKMOND		30	1610	1611	21.35	22.86	3.53		1.07	329			
RUN: 2															
15	15	ROGERS	FRED	23	1628	1629	18.36	25.97	6.70		1.86	362			SECOND RUN WITH HIGHER MAIN FLOW.
15	15	ROGERS	FRED	23	1632	1633	18.74	24.43	6.01		1.49	332			ABOUT 2280.
15	15	ROGERS	FRED	23	1636	1637	19.11	24.43	5.33		1.15	332			

EXPERIMENT #: 28

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/09/26

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	PLATE TEMP		COLD	DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
							HOT (C)	(C)								
15	15	ROGERS, FRED		23	1640	1641	19.49	24.13	4.64		.87	306				
RUN: 1																
16	16	HUDSON		9	1535	1535	21.30	26.00	4.70	.0024	.9	411				
16	16	HUDSON		9	1535	1535	21.65	26.40	2.70	.0024	.26	399				
16	16	HUDSON		9	1535	1535	21.30	26.00	4.70	.0018	.7	415				
16	16	HUDSON		9	1535	1535	21.90	26.80	3.75	.0018	.26	408				
16	16	HUDSON		9	1535	1535	21.65	26.40	2.70	.0018	.26	408				
16	16	HUDSON		9	1535	1535	21.30	26.00	4.70	.0018	.57	398				
16	16	HUDSON		9	1535	1535	22.10	26.80	2.70	.0018	.26	389				
RUN: 2																
16	16	HUDSON		9	1604	1604	21.30	26.00	4.70	.0024	.9	396				
16	16	HUDSON		9	1604	1604	22.40	26.40	2.70	.0024	.57	402				
16	16	HUDSON		9	1604	1604	21.30	26.00	4.70	.003	.57	365				
16	16	HUDSON		9	1604	1604	21.65	26.40	3.75	.003	.57	349				
16	16	HUDSON		9	1604	1604	22.10	26.80	2.70	.003	.26					
RUN: 1																
18	18	HUDSON		9	1532	1535	21.50	27.90	6.40	.0024	1.65	396				
18	18	HUDSON		9	1535	1548	22.90	26.65	3.75	.0018	.26	405				
18	18	HUDSON		9	1545	1557	23.35	26.20	2.70	.0018	.26	158				
18	18	HUDSON		9	1604	1608	23.65	26.90	3.25	.0024	.4	624				
18	18	HUDSON		9	1608	1633	23.20	26.50	3.30	.003	.4					
RUN: 2																
21	21	TRUEBLOOD		26	1537	1538	19.88	25.00	5.12		1.08	454				
21	21	TRUEBLOOD		26	1540	1541	20.48	25.00	4.52		.87	423				
21	21	TRUEBLOOD		26	1546	1547	21.94	25.00	3.48		.37	408				
21	21	TRUEBLOOD		26	1550	1551	21.92	25.00	3.06		.25	241				
21	21	TRUEBLOOD		26	1554	1555	23.07	25.00	1.93		.19	112				
21	21	TRUEBLOOD		26	1601	1602	23.00	25.00	2.00		.16					
RUN: 1																
21	21	TRUEBLOOD		26	1614	1614	20.02	25.00	4.98		.92	390				
21	21	TRUEBLOOD		26	1617	1618	20.69	25.00	4.31		.73	385				
21	21	TRUEBLOOD		26	1620	1622	21.46	25.00	3.52		.46	359				
21	21	TRUEBLOOD		26	1625	1626	21.92	25.00	3.08		.32	161				
21	21	TRUEBLOOD		26	1635	1636	23.50	25.00	2.50		.25	13				
21	21	TRUEBLOOD		26	1636	1637	23.00	25.00	2.00		.16					
RUN: 2																
24	24	SEAPOLAY		25	1604	1604	25.40	27.90	5.0	.77	.25	0				
24	24	SEAPOLAY		25	1605	1605	25.70	28.70	3.50	.77	.26	20				
24	24	SEAPOLAY		25	1638	1638	26.00	30.60	4.60	.77	.26	480				
24	24	SEAPOLAY		25	1638	1638	26.00	30.60	4.60	.77	.26					
RUN: 1																
25	25	BOYS		3	1606	1616	21.10	23.50	2.70	10.6	.22	141				
25	25	BOYS		3	1620	1630	19.90	23.50	4.90	10.6	.26	199				
25	25	BOYS		3	1630	1630	18.60	23.50	7.90	10.6	.16	340				
25	25	BOYS		3	1642	1647	16.10	23.30	7.20	10.6	.22	2315				
RUN: 2																
26	26	HINDMAN		6	1602	1617	26.00	26.00	0.00	1.4	.15	0				

VOL SAMP X 10**5

EXPERIMENT #: 28

NUCLEI MEASUREMENTS

PROCESSING DATE: 81/09/26

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST K	REMARKS
							COLD	HOT (C)							
26	26	HINDMAN		6	1602	1617	26.00	26.00	0.00	1.4	0	0			
26	26	HINDMAN		6	1602	1617	26.00	26.00	0.00	1.4	0	0			
26	26	HINDMAN		6	1602	1617	26.00	26.00	0.00	1.4	0	0			
RUN: 2															
26	26	HINDMAN		6	1618	1632	26.00	26.00	0.00	1.2	2	2			
26	26	HINDMAN		6	1618	1632	26.00	26.00	0.00	1.2	19	19			
26	26	HINDMAN		6	1618	1632	26.00	26.00	0.00	1.2	.12	.12			
26	26	HINDMAN		6	1618	1632	26.00	26.00	0.00	1.2	.058	.058			
RUN: 1															
27	27	OHTAKE		19	1544	1545	26.00	26.00	0.00	5.5	2.84	2.84		95% 10% R.M. NOMINAL	
27	27	OHTAKE		19	1544	1545	26.00	26.00	0.00	5.5	2.34	2.34			

80/10/15

EXPERIMENT # 29

PURPOSE

COMPARE INSTRUMENTS WITH HYDROPHOBIC AEROSOL.

DESCRIPTION OF EXPERIMENT

DATE 15 OCT. 1980 1644 1750
TIME START END

NUCLEI TYPE
PARAFFIN (HOUSEHOLD TYPE).

GENERATION METHOD

N2 JET INTO LIQUID PARAFFIN SURFACE.

SIZE DISTRIBUTION SHAPING

NA

REMARKS

VERY BROAD DISTRIBUTION.

WEATHER SYNOPSIS

NONE

MEAN WEATHER CONDITIONS DURING EXPERIMENT

ROOM CONDITIONS		OUTSIDE CONDITIONS					
AVG TEMP (C)	AVG RH (%)	AVG PRESS (MB)	AVG TEMP (C)	AVG RH (%)	AVG PRESS (HR)	AVG WS (M/S)	AVG WD (DEG)

PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

EXPERIMENT #: 29

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	SAMPLE END	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OBS N	EST R	REMARKS
							COLD	HOT (C)							
RUN: 1															
2	2	POLITOVICH		20	1731	1732	23.60	25.70	2.50	.17		169	235		
RUN: 1															
7	7	LEAITCH		14	1656	1657		9.90	.283	.48		28			
7	7	LEAITCH		14	1706	1707		9.85	.283	.42		43			
7	7	LEAITCH		14	1716	1717		9.60	.283	.42		50			
7	7	LEAITCH		14	1726	1727		9.15	.283	.36		40			
RUN: 1															
9	9	GAGIN		5	1725		21.80	27.20	5.40	1.17	0	0			
9	9	GAGIN		5	1727		23.20	26.90	3.70	.53	0	0			
9	9	GAGIN		5	1729		23.90	27.00	3.10	.3	0	0			
9	9	GAGIN		5	1732		25.10	27.20	2.10	.17	0	0			
RUN: 1															
10	10	LALA		13	1707		22.05	25.26	3.21	.3	36.4				
10	10	LALA		13	1708		21.94	25.23	3.18	.4	21.5				
10	10	LALA		13	1709		21.46	25.19	3.56	.5	26.9				
10	10	LALA		13	1710		20.50	25.15	4.00	.9	10.1				
10	10	LALA		13	1711		20.50	25.10	3.09	.7	18.2				
10	10	LALA		13	1712		20.55	25.04	4.47	.9	25.5				
10	10	LALA		13	1713		19.65	24.88	4.03	.7	25.5				
10	10	LALA		13	1714		19.72	24.89	6.07	1.52	18.5				
10	10	LALA		13	1716		17.72	24.69	6.97	.7	11.3				
NO COUNTS ABOVE NOISE LEVEL															
RUN: 1															
13	13	AYERS		2	1712	1719									
NO PARTICLES IN RANGE 0.25 TO 1.25μ															
RUN: 1															
15	15	ROGERS.	FRED	23	1709	1710	18.36	25.05	6.70	1.46	110				
15	15	ROGERS.	FRED	23	1713	1714	18.74	24.44	5.01	1.45	93				
15	15	ROGERS.	FRED	23	1717	1718	19.51	24.44	5.33	1.87	100				
15	15	ROGERS.	FRED	23	1721	1722	19.49	24.44	4.64	.63	115				
15	15	ROGERS.	FRED	23	1726	1727	19.87	23.81	3.30	.265	114				
15	15	ROGERS.	FRED	23	1730	1731	20.23	23.50	2.56	.14	121				
15	15	ROGERS.	FRED	23	1734	1735	20.63	24.18	1.89	.14	106				
15	15	ROGERS.	FRED	23	1738	1739	20.99	24.88	2.18	.14	106				
MAIN FLOW ABT 2280. MAY BE SEETING ONLY LARGE WAX PARTICLES.															
RUN: 1															
18	18	HUDSON		9	1701	1703	23.20	26.45	3.25	1.45	152				
18	18	HUDSON		9	1709	1711	21.85	27.80	5.95	.0018	1.78				
18	18	HUDSON		9	1754	1756	23.60	25.85	2.25	.0018	.03				
RUN: 1															
24	24	SERPOLAY		25	1705	1708	26.30	28.40	2.10	.77	340				
24	24	SERPOLAY		25	1712	1715	26.30	28.80	2.50	.25	340				
24	24	SERPOLAY		25	1720	1722	26.30	29.30	3.00	.77	340				
24	24	SERPOLAY		25	1725	1730	26.20	28.10	1.90	.14	310				
24	24	SERPOLAY		25	1735	1739	26.20	27.70	1.50	.09	150				
RUN: 1															
25	25	BORYS		3	1714	1716	16.00	22.90	6.90	4.24	1132	1132			VOL SAMP X 10**5

PROCESSING DATE: 81/09/26

NUCLEI MEASUREMENTS

EXPERIMENT #: 29

MACH NO.	DUCT NO.	NAME	OBSERVER	NO.	SAMPLE START	TIME END	PLATE TEMP		DELTA	VOLUME SAMPLED (L)	SUPER SAT (%)	NUCLEI COUNT	OHS N	EST K	REMARKS
							COLD	HOT (C)							
25	25	RORYS		3	1726	1728	17.80	22.80	5.00	10.6	1.08	410	410		
RUN: 1															
26	26	HINDMAN		6	1716	1733	26.00	26.00	0.00	1.4	.15	203	203		+/-3.0
26	26	HINDMAN		6	1716	1733	26.00	26.00	0.00	1.4	.11	44	44		+/-0.6
26	26	HINDMAN		6	1716	1733	26.00	26.00	0.00	1.4	.041	12	12		+/-0.16
26	26	HINDMAN		6	1716	1733	26.00	26.00	0.00	1.4	.027	.038	.038		+/-0.0066
26	26	HINDMAN		6	1716	1733	26.00	26.00	0.00	1.4	.016	0	0		0
RUN: 1															
27	27	OHTAKE		19	1710	1710				6.7		0	0		75%
27	27	OHTAKE		19	1710	1710				6.7		0	0		118% NOMINAL

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16. ABSTRACT <p>Accurate knowledge of cloud condensation nuclei size distributions is paramount to improving our understanding of cloud microphysical phenomena. The problems associated with obtaining this knowledge are large and varied; therefore, many technique development efforts to resolve these problems are in progress around the globe. The objective of the Third International Cloud Condensation Nuclei Workshop was to bring a large number of the development efforts together to insure an exchange of results and progress being made in the hope that the attainment of satisfactory solutions could be accelerated. At this stage it appears as if the workshop was very successful.</p> <p>The complexity and magnitude of the workshop were such that it could be successfully accomplished only under the combined sponsorship of the National Aeronautics and Space Administration and the National Science Foundation. The workshop was supported by contract NAS8-33820; this document constitutes the final report under the contract.</p>					
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