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PLUME CONTAMINATION EFFECTS PREDICTION. THE CONTAM COMPUTER PROGRAM, VERSION II

R. J. Hoffman, et al

McDonnell Douglas Astronautics Company

Prepared for:

Air Force Rocket Propulsion Laboratory

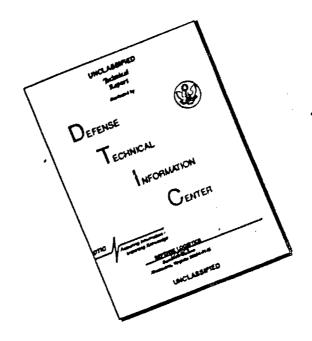
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The effect of rocket exhaust plume impingement on sensitive vehicle surfaces is an area of continuing concern in the design of spacecraft, missiles and reentry vehicle systems. Specifically, the contamination and subsequent degradation of functional surfaces, such as solar cells, thermal control coatings, optical lenses, optical view ports, and highly reflective surfaces, have resulted in compromises of mission effectiveness. The objective of this study was to develop a single computer code capable of predicting the production.

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transport, and deposition of engine and plume contaminants, and the change in absorptivity, emissivity, reflectivity, and transmissivity of a functional spacecraft surface, such as thermal control coatings and optical view ports and lenses, resulting from plume contaminant deposition or mechanical abrasion (sand blasting). Both bipropellants and monopropellants have been treated. Surface chemical reaction with a deposited plume contaminant layer was not treated. Analytical models and computer subprograms have been developed and integrated to form the CONTAM computer program. Complete User's manuals for each of the computer subprograms as well as the CONTAM program are included in this report, along with details of the analysis and numerical methods. A sample case illustrating the CONTAM program's capability is presented.

FOREWORD

This report was prepared for the Air Force Rocket Propulsion Laboratory (AFRPL), United States Air Force, Edwards, California, under Contracts F04611-70-C-0076 and F04611-72-C-0037, by the McDonnell Douglas Astronautics Company (MDAC), Huntington Beach, California. The Air Force program monitors for this study were Dr. L. P. Quinn (RTSP) and James R. Nunn, Capt., USAF (RTSP). The study was performed during the period August 1970 to April 1973, and the final report submitted to AFRPL for approval on May 15, 1973. The results of the initial contract were previously reported in AFRPL-TR-71-109, December 1971.

The MDAC study manager for this project was Mr. R. J. Hoffman, Supervisor, Plume Effects, Aero/thermodynamics and Nuclear Effects Department. Mr. W. T. Webber acted as Principal Investigator for the study and Tesk Leader for the bipropellant contamination production task. Other members of the Plume Effects Group included Mr. R. G. Oeding, Task Leader, Contaminant Transport and Chemical Kinetics; Mr. F. Cramer, Task Leader, Monopropellant Contamination; Dr. G. Yanow, Task Leader, Final Report and Programming; and Mr. S. S. Cherry, Task Leader, Contamination Experiment, Dr. D. W. English, Propulsion Department was the Task Leader for the Surface Effects Portion of the study. Capt. J. R. Nunn, USAF provided valuable information concerning system application of the CONTAM Computer Program and continuing support for which we are grateful.

This technical report has been reviewed and is approved.

ABSTRACT

The effect of rocket exhaust plume impingement on sensitive vehicle surfaces is an area of continuing concern in the design of spacecraft, missiles, and reentry vehicle systems. Specifically, the contamination and subsequent degradation of functional surfaces, such as solar cells, thermal control coatings, optical lenses, optical view ports, and highly reflective surfaces, have resulted in compromises of mission effectiveness. The objective of this study was to develop a single computer code capable of predicting the production, transport, and deposition of engine and plume contaminants, and the change in absorptivity, emissivity, reflectivity, and transmissivity of a functional spacecraft surface, such as thermal control coatings and optical view ports and lenses, resulting from plume contaminant deposition or mechanical abrasion (sand blasting). Both bipropellants and monopropellants have been treated. Surface chemical reaction with a deposited plume contaminant layer was not treated. Analytical models and computer subprograms have been developed and integrated to form the CONTAM computer program. Complete User's manuals for each of the computer subprograms as well as the CONTAM program are included in this report, along with details of the analysis and numerical methods. A sample case illustrating the CONTAM program's capability is presented.

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Section I INTRODUCTION

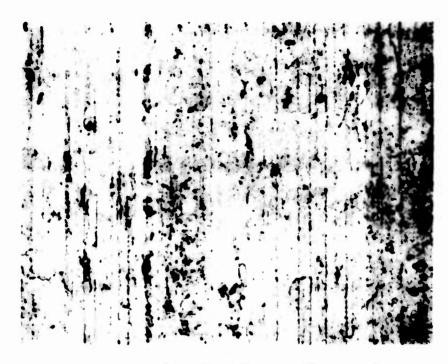
The effect of rocket exhaust plume impingement on sensitive vehicle surfaces is an area of continuing concern in the engineering design of space-craft, missiles, boosters, and RV systems. Specifically, the contamination and subsequent degradation of functional surfaces, such as solar cells, thermal control coatings, optical lenses, optical view ports, highly reflective (mirrored) surfaces, and sealants, have resulted in compromises of mission effectiveness.

To illustrate the deposition of contaminants problem, several photographic examples of contamination occurring as the result of actual bipropellant engine firings will be presented. These examples were taken from a series of contamination experiments conducted by MDAC under the MOL program.

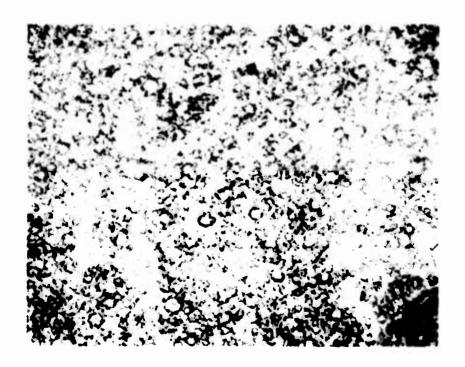
Figure 1 compares a control surface (no impingement) with a surface which has been exposed to normal impingement by the exhaust plume of a liquid bipropellant engine (MMH-NTO). Evidence of surface damage is apparent, and was postulated to be caused by condensed droplets in the core of the plume flow.

During a vacuum chamber subscale thrustor test by MDAC at AEDC, a 1-lb thrust Marquardt MMH-NTO rocket engine was fired horizontally so that the exhaust products would impinge upon a vertically oriented test panel containing several surface specimens. Surface specimens included thermal control coatings, polished metal, and specialized glass lenses.

During pulse-mode operation of the motor, copious quantities of brownish, viscous liquid were observed about the nozzle lip and upon the lower external surface of the motor. This liquid exhibited considerable activity, bubbling while suspended from the motor lower external surface apparently due to some boiling and/or decomposition phenomenon. An impingement pattern of sorts was visible upon the test panel that appeared symmetrical but did not agree well (qualitatively) with theoretical predictions of the gas-phase impingement region. The region within the symmetric impingement pattern was coated with viscous liquid and/or solid material that increased in quantity with the number of pulses to which the panel was exposed; the coloration of this material was difficult to identify, but was definitely darker than the panel. Above and below the symmetric impingement region liquids were splattered in very large quantities, particularly near and below the rocket motor where semisolid formations of brownish color were noted; deposition of this variety seemed randomly distributed and was observed fore and aft of the nozzle exit plane. The amount of liquid generated by the rocket motor



a. CONTROL SAMPLE SHIELDED FROM DIRECT IMPINGEMENT



b. SAMPLE EXPOSED TO NORMAL EXHAUST IMPINGEMENT

Figure 1. Liquid Bipropellent Plume Impingement Effects (22-lb Thrustor MMH-NTO, O/F = 1.65)

varied inversely with pulse duration, with maximum amount generated during the 16-msec "minimum impulse" pulses. Thermal control coatings exposed to these exhaust products were visibly coated and suffered losses in reflectance. Similarly, transparent samples suffered transmittance losses. After certain periods of liquid buildup upon the panel, brownish liquids ran down the panel surface. During long-duration firings in excess of several hundred seconds, a well-defined symmetric impingement pattern was noted upon the panel. It was difficult to discriminate between liquid and solid formations at this point. The symmetric impingement region gained further definition by virtue of a continuous ridge of solid and/or viscous liquid deposits at the symmetric impingement region boundaries. Posttest microphotography revealed that glass surfaces were coated with micron-sized droplets, even though the incidence angle of exhaust products was (theoretically) very small or nonexistent. Deposits upon the panel and surface samples displayed phase instability at standard temperature and pressure (STP) conditions, changing from solid to liquid to solid when disturbed physically or environmentally. When the chamber was repressurized to facility ambient conditions, much of the material deposited upon the panel became less viscous and ran off the panel.

Figures 2 through 4 show some of the deposits observed after various duty cycles.

This report will present the results of a 27-month study for the Air Force Rocket Propulsion Laboratory to develop an analytical model and computer program system for the prediction of spacecraft functional surface contamination effects caused by interactions with liquid bipropellant rocket exhaust plumes. Emphasis has been placed on development of computer codes to describe the complex two-phase combustion gas-dynamic processes occurring in a bipropellant combustor and the thermodynamic and kinetic nonequilibrium processes occurring during a two-phase nozzle and plume expansion. The operation of monopropellant hydrazine (N2H4) combustors has also been included in this study.

This report is divided into six discrete parts: the main body of the report and five appendixes. In the main body, emphasis is placed on describing the operating characteristics of the integrated Plume Contamination Effects Prediction Computer Program, CONTAM. A description of the program, User's Manual, and sample case run illustrating the ability of CONTAM to predict contaminant production, transport, and condensation are presented.

Each of the subprograms of <u>CONTAM</u> are described in detail in a separate appendix as follows:

Appendix A TCC Transient Combustion Chamber Dynamics

Computer Program (a bipropellant contam-

inant production model)

Appendix B MULTRAN Multiphase Nozzle and Plume Transport Computer Program (a multiphase nozzle

and plume flow field characterization model)

Figure 2. Deposition Effects of Rocket Exhaust Impingement after Continuous Long-Duration Firing

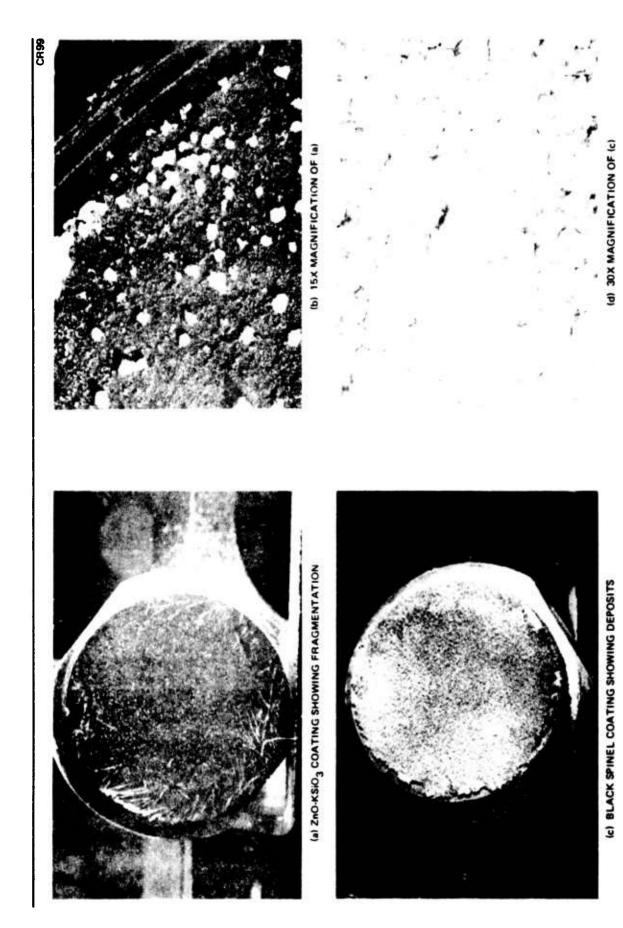


Figure 3. Effert of Thrustor Exhaust Continuous Long-Duration Firing-ZnO-KSiO₃ and Spinel Coatings



(a) HORIZON SENSOR LENS SHOWING DEPOSITS



(b) 80X MAGNIFICATION OF (a)

Figure 4. Effect of Thrustor Exhaust Continuous Long-Duration Firing on Glass Sample

Appendix C KINCON Nonequilibrium Chemical Kinetics and

Condensation Computer Program (a multi-

phase reacting gas streamtube model)

Appendix D SURFACE Deposition and Surface Effects Computer

Program (a plume impingement, deposition, abrasion, and surface contamination effects

model)

Appendix E N2H4 Monopropellant Combustion Chamber

Dynamics Program

In addition to the sample case in the main body of the report, each appendix contains a sample case illustrating additional capabilities of the particular subprogram. Detailed operating information for each subprogram is contained in the User Manual section of each appendix.

This report has been loose-leaf bound to facilitate updating of the various User's Manuals, either by MDAC or other Government and industry users. The authors would greatly appreciate comments, corrections, additions, and suggestions for inclusion in future updates to be distributed to all users. Please send comments to:

R. J. Hoffman A3-833-BBBO-28 McDonnell Douglas Astronautics Company 5301 Bolsa Avenue Huntington Beach, California 92647

Qualified persons may obtain copies of the <u>CONTAM</u> computer program by a addressing a request to:

AFRPL/DOE Edwards AFB Edwards, California 93523

Section II OBJECTIVES AND SCOPE

1. OBJECTIVES

The objective of the study was to develop a single computer code capable of predicting the production, transport, and deposition of engine and plume contaminants and the change in absorptivity, emissivity, reflectivity, and transmissivity of a functional spacecraft surface, such as thermal control coatings and optical view ports and lenses, resulting from plume contaminant deposition or mechanical abrasion (sand blasting). Surface chemical reaction with a deposited plume contaminant layer is not treated.

The study was divided into five main areas:

- (1) Improvement of predictive technology for the characterization of reactive, multiphase rocket nozzle and exhaust plume flows containing propellant contaminants and nonequilibrium combustion products, including condensables.
- (2) Continued development of an analytical model to predict the production of contaminants in bipropellant rocket-engine combustion chambers.
- (3) Development of a semiempirical model to predict changes in surface properties of functional spacecraft surfaces (resulting from deposition or abrasion).
- (4) Integration and coupling of existing computer programs and newly developed computer programs to achieve a systems engineering design tool for the prediction of contaminant effects on spacecraft surfaces.
- (5) Verification of the contamination prediction model by comparison with experimental data.

2. SCOPE

The study was restricted to the development of predictive methods for the production, transport, and deposition of contaminants from hydrazine monopropellant and hydrazine-family fuels in combination with nitrogen tetroxide and to changes in thermal and optical surface properties of common thermal-control paints and optical lenses. The model development has considered RCS engines in the 5- to 600-lb thrust range; the validity of the model for much larger engines has not been assessed.

SECTION III MODEL DESCRIPTION

Section IV describes the <u>CONTAM</u> plume contamination effects prediction computer program system developed during this study. The analytical models associated with each link of the <u>CONTAM</u> program are discussed in detail in the appropriate appendix. In this section, a summary description of the analytical models employed in the <u>CONTAM</u> system is given.

The general objectives of the study were to construct a single analytical model capable of predicting the effects of bipropellant and monopropellant plume impingement contamination on optical and thermal spacecraft surfaces based only on a knowledge of available engine operating conditions, engine/ spacecraft configuration geometries, and spacecraft orbital parameters. To this end, it was necessary to construct a model for the production of contaminants in bipropellant and monopropellant combustion chambers (unburned propellants ejected through nozzle throat); transport of these contaminants by the expanding gases in the nozzle and exhaust plume; chemical nonequilibrium composition of plume species; condensation of plume species in the expanding plume; abrasion damage and deposition resulting from plume impingement; and, finally, the changes in thermal and optical surface properties, absorptivity, emissivity, transmissivity, and reflectivity resulting from contaminant deposition and/or abrasion damage. In addition, the model considers the effect of engine duty cycle and spacecraft radiant energy transfer on the rate of contaminant deposition over an entire mission profile.

The feasibility of constructing a valid model, considering all of the above aspects, relied heavily upon the existence of several models and computer codes which could be used as a basis for construction of the overall contamination effects prediction model. Several new portions of the model and computer subprograms were developed. Figure 5 schematically illustrates the computation flow logic and the related computer codes. Details of each computer code can be found in the appropriate appendix.

1. COMBUSTION CHAMBER CONTAMINANT PRODUCTION (See Appendixes A and E for further details)

Unburned propellant and intermediate products of combustion (liquid phase) ejected from the combustion chamber are considered first as a source of contaminants. Referring to Figure 5, the Transient Combustion Chamber (TCC) Dynamics Program, developed at MDAC, is used to calculate contaminant production from bipropellant thrust chambers. The contaminant production from monopropellant hydrazine thrust chambers is calculated using a modified version of the program developed at UARL by A.S. Kesten, et al (References 1 and 2).

TCC PROGRAM

Contaminant material is produced by the combustor of a bipropellant rocket engine when unburned propellant vapor passes through the throat, when partially burned propellant droplets pass through the throat or when fuel and oxidizer droplets strike a cold chamber wall to form a liquid film of hydrazine nitrate which is moved downstream by chamber gas shear forces. When the unburned propellant or intermediate reaction products are ejected from a rocket engine, they may be transported in the plume and deposited on nearby sensitive spacecraft surfaces, changing their thermal or optical properties.

The sequence of combustion related events in the rocket engine combustion chamber is calculated by numerically integrating the differential and algebraic equations which describe the basic processes of the feed system, injector, and combustion chamber. Figure A-1 in Appendix A is a drawing of the rocket system.

a. Feed Systems

The feed systems are approximated with single lumped parameters representing the inertial and resistive aspects of the feed system, the rate of acceleration of flow being proportional to the amount that the instantaneous pressure drop exceeds the instantaneous pressure losses in the system. The opening and closing of the valves are modeled by varying the feed system resistance as a function of time. Flow reversals or initial start conditions, which result in partially or fully gas-filled feed lines, are simulated by varying both the resistance and inertia of the feed system as functions of time.

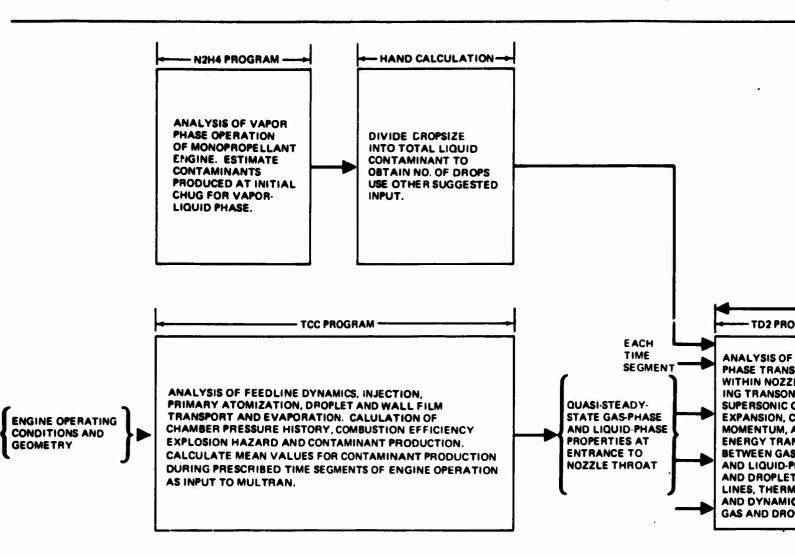
b. Atomization

The atomization process is calculated for one of several modes, depending on the chamber pressure. If the injected propellant is sufficiently supersaturated, the stream is presumed to flash-atomize. The flash-atomization process resembles the gas-atomization process, with the gas being supplied by the explosive growth of bubbles in the supersaturated stream. The flash atomization process gives relatively fine droplets, on the order of 40 microns in diameter.

When the chamber pressure is sufficiently high that the injected propellant streams do not flash, the atomization occurs by the impingement of the fuel and oxidizer streams. The median droplet diameter is obtained from an equation based on the orifice diameters, injection velocities, relative momentum of the streams, and physical properties of the propellants.

When only one stream is being injected during a start transient, there can be no impingement and droplet formation is calculated based on singlestream breakup.

The injected propellant moves from the injection point to the impingement point along the direction of injection. After impingement, the



INPUT DATA

- FEEDLINE CHARACTERISTICS
- INJECTOR AND COMBUSTOR
 GEOMETRY
- PROPELLANT TANK PRESSURES
 AND TEMPERATURES
- PROPELLANT VALVE TIMING AND RAMP CHARACTERISTICS
- PULSE WIDTH AND DUTY
 CYCLE
- PROPELLANT PROPERITIES

• COMBUSTOR WALL TEMPERATURE

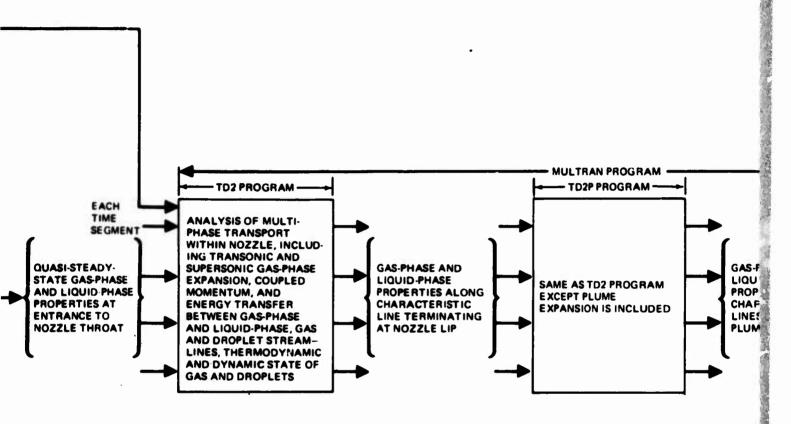
TCC DUTPUT (TIME DEPENDENT)

- CHAMBER PRESSURE AND TEMPERATURE
- AMOUNT OF UNBURNED PRO-PELLANTS
- AMOUNT OF LIQUID PROPELLANTS EJECTED FROM CHAMBER
- SIZE AND VELOCITY DISTRIBUTION OF EJECTED PROPELLANT DROPLET
- AMOUNT OF LIQUID DEPOSITED ON CHAMBER WALL
- AMOUNT OF LIQUID WALL FILM EJECTED FROM CHAMBER

TD2 INPUT (AVERAGED OVER EACH TIME SEGMENT)

- CHAMBER PRESSURE AND TEMPERATURE
- DISTRIBUTION OF SIZE AND NUMBER OF DROPLETS ENTER-ING THROAT

• ETC



TD2 INPUT (AVERAGED OVER EACH TIME SEGMENT)

- CHAMBER PRESSURE AND TEMPERATURE
- DISTRIBUTION OF SIZE AND NUMBER OF DROPLETS ENTER-ING THROAT

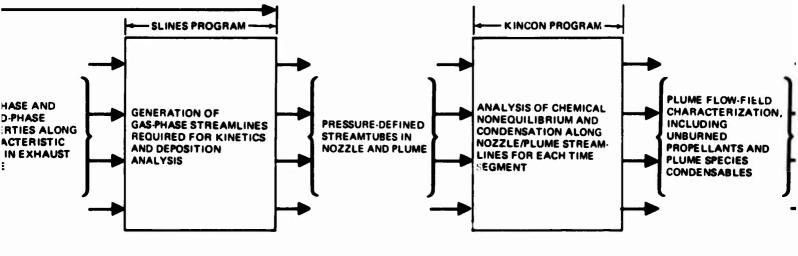
TD2P INPUT (FOR EACH TIME SEGMENT)

- STATIC GAS TEMPERATURE, PRES-SURE, VELOCITY, AND FLOW DIRECTION
- LIQUID DROPLET TEMPERATURE, DENSITY, VELOCITY, SIZE DIS-TRIBUTION, AND FLOW DIRECTION

TD2P O

- STATIC SURE, DIREC
- LIQUIC DENSI*
 TRIBU

B



JTPUT (FOR EACH TIME SEGMENT)

₹.,

GAS TEMPERATURE, PRES-/ELOCITY, AND FLOW 'ION

DROPLET TEMPERATURE, Y, VELOCITY, SIZE DIS-ION, AND FLOW DIRECTION KINCON INPUT (FOR EACH TIME SEGMENT AND EACH STREAM-TUBE)

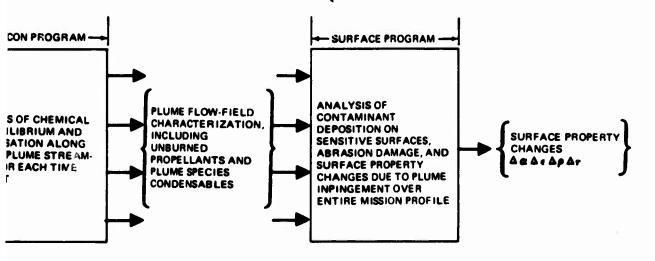
- AXIAL DISTANCE VS. PRESSURE ALQNG EACH STREAMTUBE
- INITIAL COMPOSITION, TEMPER-ATURE, PRESSURE, VELOCITY

SURFACE INPUT

- PLUME STATIC PRESSURE TEMPERATURE, VELOCITY FLOW DIRECTION
- UNBURNED PROPELLANT DROPLET TEMPERATURE, VELOCITY, SIZE DISTRIBU
- NONEQUILIBRIUM PLUME
- CONDENSED PLUME SPECI

Figure 5. Schematic Diag

1



SURFACE INPUT

- PLUME STATIC PRESSURE, TEMPERATURE, VELOCITY AND FLOW DIRECTION
- UNBURNED PROPELLANT LIQUID DROPLET TEMPERATURE, DENSITY, VELOCITY, SIZE DISTRIBUTION
- NONEQUILIBRIUM PLUME COMPOSTION
- CONDENSED PLUME SPECIES

Figure 5. Schematic Diagram of Analytical Model Elements and Related Computer Programs

stream moves in the direction of the resultant angle. The stream moves in this new direction until its atomization is complete, after which its two-dimensional trajectory is determined by the aerodynamic drag forces.

c. Chamber Calculations

The vapors or gases that fill the combustion chamber are derived from several sources: vapor from flashing propellant streams; material evaporated from propellant droplets; evaporation of material deposited on the combustion chamber walls and from the ignitor if one is used. Fuel and oxidizer vapor from these sources are axially cumulated in the chamber at each time interval, while the amounts calculated to flow through the nozzle are subtracted. This gives current values for the chamber-gas mass and stoichiometry, and the axial addition rate of mass. These are used to calculate the pressure, temperature, molecular weight, and velocity distribution in the chamber. The simplifying assumption is made that at any instant the pressure is constant throughout the chamber, and the gas is well mixed. Hypergolic ignition is based upon global vapor-phase chemical kinetics, and extinguishment is based upon a quenching distance correlation.

d. Wall Calculations

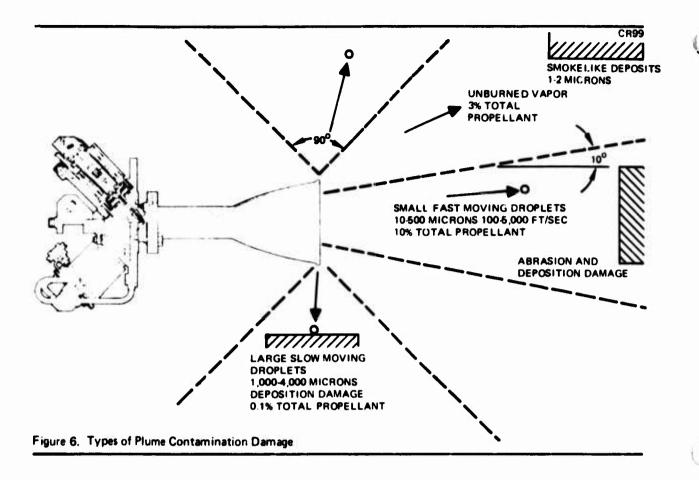
When a computed propellant droplet moves radially to the location of the combustion chamber wall, its fuel or oxidizer mass is added to the axial distribution of fuel or oxidizer previously deposited on the chamber wall. In calculating the axial and tangential velocities of the liquid material on the wall, each axial element of fluid is treated as an inertial free body acted upon by shear from the wall and gas, spin-induced forces, mass and momentum increments from the impinging droplets, from evaporation, from entrainment, and by convection at the upstream and downstream boundaries of the element. Evaporation is calculated based upon heat transfer from the gas during the firing, and by heat transfer from the wall coupled to vacuum evaporation during the "off" periods.

The heat transfer and entrainment associated with each axial segment of wall is calculated using the correlations of Gator, with corrections being made for mass transfer and rippling. The shear stress and heat transfer coefficients are correlated with local Reynolds numbers which are reevaluated at each axial segment of the chamber each time interval.

For ease of visualization, many of the calculations may be presented as computer plots or computer produced motion pictures.

e. Comparison with Experimental Results

Contamination from a conventional bipropellant RCS engine takes one or more of three forms: reacted or unreacted propellant vapor; incompletely burned droplets expelled through the throat; and unburned propellant that impinges upon the chamber wall and is eventually ejected from the nozzle lip. Figure 6 shows typical values for the amounts, directions, sizes, and velocities of the material making up these three forms for one particular small engine and one particular duty cycle.



The vapors of fuel, oxidizer, or combustion products emitted during preignition, ignition, steady state, or postcutoff dribble periods will form plumes that can impinge upon various surfaces with the possibility of deposition and in-situ reaction. Contamination from this source usually takes the form of a hazy deposit of smokelike particles (fairly uniform in size, 1 to 2 microns).

The fuel and oxidizer droplets, which are too large to burn completely in the chamber and which are centrally directed, will pass through the nozzle throat. These particles will be accelerated by aerodynamic forces both upstream and downstream of the throat, and can attain quite high velocities, which gives this class of particles the capability of doing considerable damage by abrasion.

The third form of contamination is the propellant that impinges upon the chamber wall and is then dragged downstream under the influence of shear forces from the combustion product gases. If this wall-film material is able to move to the nozzle lip without being thermally destroyed, it will be thrown off as large droplets in directions roughly normal to the axis of the chamber. This material is generally dark colored and shows the effects of thermal decomposition.

Figure 4 shows a horizon sensor lens exposed to wall-film contamination from an RCS engine operated in a vacuum chamber.

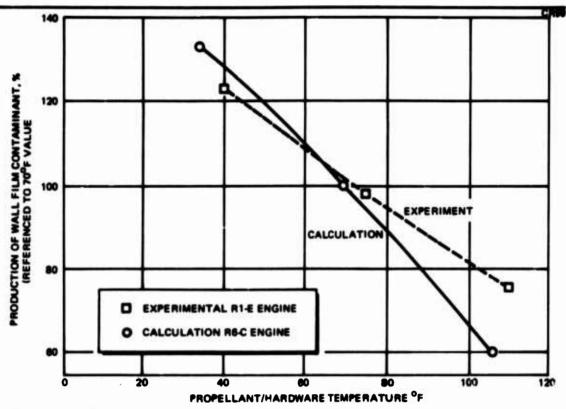


Figure 7. Comparison of Experimental and Calculated Wall-Film Contaminant Production as Function of Propellant/Hardware Temperature

Quantitative experimental data on contaminant production are still rather scarce. However, two recent papers have been published describing contaminant production from pulse-mode firings of the 22-pound Marquardt R1-E engine (References 3 and 4). This engine is very similar in design, but larger than the 5-pound Marquardt R6-C engine, which has been thoroughly parametrically analyzed, using the <u>TCC</u> program. Many aspects of the experimental firings of the R1-E engine agree with the trends calculated for the R6-C engine.

Martinkovic found that contaminant production was a function of injector temperature. His measurements for the temperature trend of the R1-E engine show good agreement with our calculations for the R6-C engine as shown in Figure 7. The absolute values for contaminant expelled as wall-film were also in good agreement. The Martinkovic 22-pound R1-E engine experimentally produced 0.772 mg of wall-film per 17 msec pulse at 75°F; i.e., about 1 part per thousand of total injected propellant ended up as wall-film contaminant. Our calculations for the 5-pound R6-C engine indicated that 0.33 mg of wall-film would be produced for each 17 msec pulse at 70°F; i.e., about 2 parts per thousand of total injected propellant. This is in agreement with the well-known trend toward increased contaminant production with decreased engine size.

Other indications of the accuracy of the <u>TCC</u> program are to be found in its excellent predictions of low-frequency combustion instability, start transients, popping amplitudes, spiking tendency, and pulse-mode specific impulse or total impulse versus pulse-width. These are discussed in some detail in Appendix A.

N2 H4: Monopropellant Combustion Chamber Dynamics Program (See Appendix E for further details)

Contaminant production data is generated for monopropellant engines by the program N2H4, based on the transient and steady-state programs originally developed by Kesten, Smith, and Smith of UARL (References 1 and 2) and modified by F.B. Cramer of MDAC. In monopropellant engines, "normal" operation will produce primarily NH3, N2, H2, and N2H4 vapors. However, during cold starts or "chugging", droplets of liquid hydrazine will also be formed as a contaminant.

The original UARL program only treated the smooth vapor phase decomposition problem. MDAC modifications have extended the program to estimate "hard start" characteristics as an alternative option. In particular, the peak pressure, time to the pressure peak, mass of liquid hydrazine ejected, and the masses of the vapor constituents are printed out. This program will not calculate beyond the start transient when calculating a cold (hard) start; however, even in its present form, this computer program gives information when cannot be obtained any other way.

The smooth monopropellant decomposition calculations are carried out using a slug flow model. That is, the vapor (vaporization is assumed to occur instantaneously) formed upon contact with the catalyst bed is modeled one dimensionally.

The MDAC N2H4 monopropellant "hard start" subprogram is based in large part on curve fits of experimental data obtained from various engine firings. At present no monopropellant program will effectively follow a cold start after the initial pressure surge. The current MDAC program does provide working information about this initial pressure peak and then shuts down with a printout of the estimated start transient operational characteristics.

2. CONTAMINANT TRANSPORT (See Appendix B for further details).

Having defined the average amount of liquid phase contaminant ejected from the combustor for each pulse segment (see Subsection III. 1) the two-phase nozzle and plume flow is then computed for transient and steady state pulse segments, using method-of-characteristics computer programs described by Nickerson and Kliegel in (Reference 5) and modified by Gabbert and Hoffman (Reference 6). In Figure 5, these programs are identified as TD2 and TD2P. Variations in chamber pressure, chamber temperature, droplet velocity, and droplet-size distribution produce considerably different two-phase 14 ow fields for each of the pulse segments (preignition, ignition, steady state, and tailoff) and therefore require a unique analysis for each pulse segment.

Because several contaminant sources originate upstream of the nozzle exit plane (for example, unburned propellant and nonequilibrium condensed fuel nitrate species), it is necessary to obtain a complete characterization of the multiphase flow at the nozzle exit, including droplet/particle

distributions (size, velocity, temperature, and species) as well as an axisymmetric distribution of the gas-phase flow. The extreme radial compression of droplet/particle laden regions indicates the need for accurate information concerning the combustor and nozzle transport of condensables as input to the plume analysis.

Starting at the convergent nozzle sections, up to 10 droplet groups are considered as an approximation to the distribution of condensed phase material produced in the combustion chamber. The concentrations, distribution, and trajectories of each droplet/particle group are considered at each mesh point in the axisymmetric method-of-characteristics flow analysis throughout the entire nozzle and plume. Fully coupled momentum exchange (drag) between the gas and droplet/particle phase is considered, including rarefication effects. The results (output) of this program set provide the initial conditions (input) for the impingement model and subsequently, the surface effects analysis.

While the transport model will provide information about the dynamic condition and flux of species arriving in the vicinity of a functional surface submerged in an exhaust plume, the kinetic/condensation model and the deposition model are required to provide information regarding the chemical composition and amount of plume exhaust material actually deposited on the submerged surface.

3. CHEMICAL KINETICS AND CONDENSATION IN THE NOZZLE AND PLUME FLOW FIELD (See Appendix C for further details)

In addition to unburned propellant droplets, many liquid-bipropellant exhausts contain condensed phases as an important contaminant source. primary condensables in bipropellant plumes are thought to be H2O and nitrate salts of the fuel. One of the major study objectives was to review existing data on plume condensables and to model the mechanism of condensation analytically in rocket nozzles and exhaust plumes. A thermodynamic nonequilibrium nucleation and condensation model has been developed and is discussed in detail in Appendix C. The MDAC Streamtube Chemical Kinetics and Condensation Computer Program, identified in Figure 5 as KINCON, has provided the framework for development of this portion of the model. A preliminary study, using the combined chemical kinetics and condensation model, was performed to size condensation characteristics in the nozzle and plume, corresponding to typical engine operating regimes (both transient and steady state). The relative effect of condensation as a contaminant source, relative to combustion chamber sources, is yet to be determined, although it is thought to be an important factor in contamination of surfaces beyond the central core of the plume where heavy, unburned propellant droplets seem to dominate.

The modeling of this phase of the contaminant-production problem requires (1) the chemical kinetic analysis of the expanding exhaust gases and (2) a realistic analysis of the condensation process.

a. Chemical Kinetics Model

The chemical kinetic processes in the nozzle and plume are calculated along streamlines utilizing the MDAC KINCON computer program. The KINCON program possesses several unique features which make it well suited for analyzing the nozzle/plume chemical kinetics. These features include (1) a fully implicit numerical integration scheme that permits the rapid integration of the full set of kinetic equations (up to 40 species and 150 reactions) with complete numerical stability; (2) capability to treat the addition (or subtraction) of mass, momentum, and energy to the streamtube by specifying the specific rate as a function of streamtube distance; (1) reaction-rate screening capability, which identifies reactions and species that are unimportant and need not be considered in the calculation of a specific species concentration or fluid property in any particular application.

Principal assumptions inherent in the use of the streamtube kinetics model include the following: (1) the flow is one-dimensional, steady, and inviscid: (2) each component of the gas mixture is a perfect gas; and (3) internal degrees of freedom of each component are in equilibrium.

b. Condensation Model

It is well known that condensation of a rapidly expanding supersonic flow does not occur at the point in the flow where the gas equilibrium temperature reaches the saturated vapor temperature of the particular species in question. Instead, condensation is delayed and eventually occurs as a "condensation shock" or condensation zone downstream of the equilibrium condensation point. Although this phenomenon is not thoroughly understood, it may be caused by a number of factors, including (1) lack of nucleation material on which condensables may form and (2) inability of the surrounding gas phase to readily remove heat from the condensing material.

To treat condensation effects in rapidly expanding gases, a kinetic model of the condensation process utilizing the classical liquid drop theory was adopted. The condensation phenomenon, as described by this model, occurs as a result of two distinct processes: (1) nucleation and (2) droplet growth.

As saturated vapor conditions are reached in a rapid expansion, sufficient surface area will not usually exist for the condensation required to maintain equilibrium ($P_v = P_{vs}$), and a supersaturated condition results ($P_v > P_{vs}$). The nucleation process (spontaneous self-nucleation) occurs in the expanding supersaturated vapor and involves the clustering of vapor molecules to give rise to very small nuclei (radius of 10 to 100 Å). Only nuclei reaching the critical drop radius r^* can exist and grow. The critical drop size is determined from thermodynamic equilibrium considerations and represents the size at which the drop has an equal probability of either evaporating or growing.

Figure 8 illustrates the effects of condensation on the flow static pressure and temperature. Following the saturation point, the vapor continues to expand along the frozen gas isentrope until a suitable number of

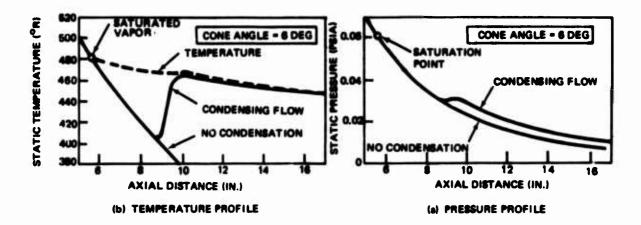


Figure 8. Static Pressure and Temperature Profiles for Condensing Nozzle Flow (Sample Case 1:90% H_2 , 10% N_2 Mixture, P_0 = 1.25 Psia, T_0 = 1,000 $^{\circ}$ R, Cone Angle = 6 Deg)

nuclei are formed and the droplet growth process begins. At this point, the effects of condensation are observed. Both the static pressure and temperature increase rapidly with the temperature approaching the saturated-vapor temperature. The expansion then continues along a different isentrope corresponding to a new gas mixture.

4. DEPOSITION, ABRASION, AND SURFACE EFFECTS (See Appendix D for further details)

An analytical model for the prediction of plume contaminant deposition, surface abrasion due to liquid and solid particle impingement, and changes in thermal and optical surface properties due to deposition or abrasion has been completed.

Development of a surface effects model was based heavily upon experimental data relating plume species deposition and mechanical abrasion characteristics to changes in α and ϵ , in the case of thermal surfaces; and changes in transmissivity and reflectivity, in the case of optical surfaces. Such data are scarce for realistic plume-deposition products, such as MMH-nitrate, although recent experiments have provided some data.

The first step in developing a model to predict surface property changes, based on a computed amount of abrasion or deposition, is to examine the possible interactions of plume material with spacecraft surfaces. The model developed accounts for plume-induced changes in α and ϵ on thermal control surfaces, such as heat-rejection radiators and optical surfaces.

Figure 9 is a sketch of the possible interactions with both coated and uncoated portions of a radiator surface.

a. Solar Absorptivity

The absorptivity of the system is primarily determined by the characteristics of the external surface upon which the external radiation falls. The average or net absorptivity α_{net} of the radiator can be taken as the mean of the absorptivity of each type of absorptive surface α_i times the area of each type Ai. Generally, α_i is a simple term, easily determined or calculated, but in some instances, such as a transparent deposit, terms related to the thickness and internal parameters of the deposit become important.

b. Hemispherical Emissivity

When contaminant deposits are thin, it is assumed that they offer little resistance to heat flux through the layer. The emissivity of the surface is assumed to be that of the contaminant layer. For thick layers of deposits, the impedance to heat flow through the layer is also modeled.

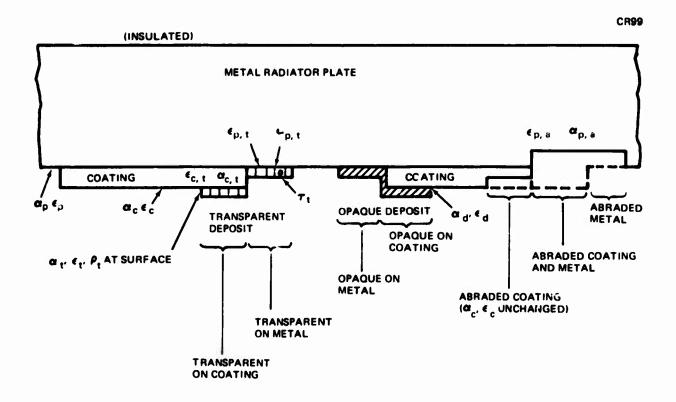


Figure 9. Contaminant Effects Physical Model-Heat Rejection Radiators

c. Surface Abrasion

Abrasion can occur on either or both of the coated or uncoated portions of the radiator surface. Abrasion of the radiator coating affects the heat flux in a step-function fashion. If only the thermal-control coating is abraded, the effect is simply that of decreasing the original coating thickness without affecting the absorptivity or the emissivity. If the abrasion proceeds far enough, it will penetrate through the coating and expose an area of the metal plate substrate. The abrasion of the metal surface, both that originally present and that exposed by removal of the coating, will alter its α_p and ϵ_p significantly. The abraded area of the metal will consist of two parts, the part that was originally bare and now abraded, plus all that was exposed when the coating is abraded away. It is assumed that in any area where the abrasion is sufficient to remove the coating, the flow field will attack the metal at once.

d. Material Deposition

A deposit of material from the plume, randomly localed on the exterior of the spacecraft radiator, acts simply like an additional coating through which the heat must be transferred. The deposit may be transparent (crystalline, glassy, or liquid), or it may be opaque due to either its basic nature or to particle sizes.

Opaque deposits affect the heat flux in a manner identical to the thermal-control coating. The situation can become more complex if the deposit forms a transparent film. Such films are not completely transparent at all wavelengths, and therefore a complex interaction occurs.

Radiant energy impinging from the environment is partly reflected, partly absorbed, and partly transmitted into the film at the outer surface in accordance with the usual ρ , α , and τ coefficients. As the energy passes through the thickness of the film, more of it is absorbed. At the bottom surface with the opaque paint or metal, the energy is either absorbed or reflected. That portion of the radiation that is reflected from the substrate then passes outward through the film, and again, part is absorbed. When the energy again reaches the outer surface, the part that strikes the surface at less than the critical angle is radiated away; but the portion that strikes the surface at an angle equal to, or greater than, the critical angle cannot escape and is eventually absorbed. A similar process also occurs for the emission of energy from and through the transparent layer.

SECTION IV THE CONTAM COMPUTER PROGRAM

1. GENERAL DESCRIPTION

This section describes the integrated computer program, CONTAM, which has been developed to provide an engineering design tool for the prediction of plume contaminant effects on sensitive spacecraft surfaces arising from direct plume impingement. Figure 10 illustrates the component subprograms of the <u>CONTAM</u> program. The <u>CONTAM</u> program is capable of independently running any of the subprograms or of running the entire analysis sequentially. Except fr. TCC or N2H4, each program is dependent upon the output of previous programs. It is suggested that CONTAM be run sequentially, with an examination of each subprogram's output before the subsequent subprogram is called since improper data inputs can result in excessive computer run time. When run independently, the capabilities of each of the subprograms may be extended to solve problems associated with: combustion dynamics; nozzle and plume multiphase flow field characterization; nonequilibrium streamtube chemical kinetics and condensation; and impingement, deposition, abrasion, and surface property changes - not necessarily associated with plume contamination.

The subprogram SURFACE has not been fully integrated into CONTAM at the present time. This program is based for the most part on very large tables of experimental data, and in the current version of the program system, the memory requirements of this particular subroutine are considerably greater than any of the other programs. Consequently, while all the rest of CONTAM has now been overlayed via "Cell Loader", SURFACE has been lest in the normal "SCOPE" loader overlay structure.

Each of the major subprograms of CONTAM are described in detail in separate appendixes as follows:

Appendix A TCC Transient Combustion Chamber Dynamics

Computer Program (a bipropellant con-

taminant production model)

Appendix B MULTRAN Multiphase Nozzle and Plume Transport

Computer Program (a multiphase nozzle and plume flow field characterization

model)

Appendix C KINCON Nonequilibrian Chemical Kinetics and

Condensation Computer Program (a multi-

phase reacting gas streamtube model)



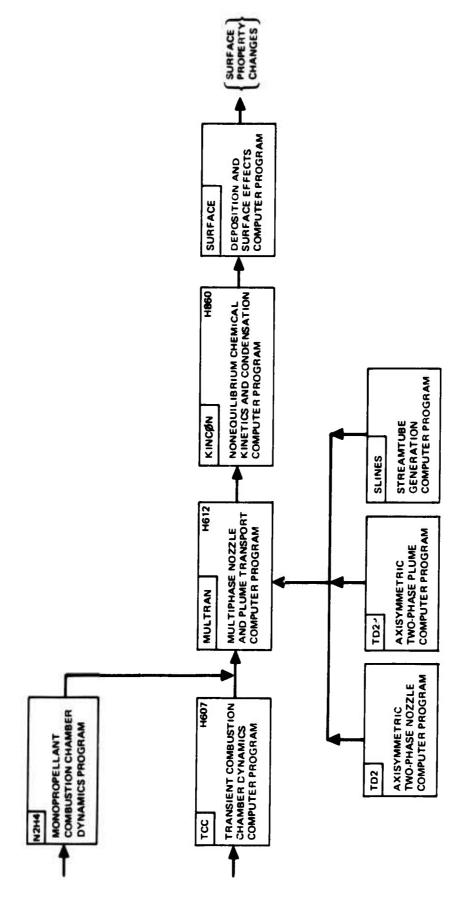


Figure 10. Schematic of Plume Contamination Effects Prediction Computer Program, CONTAM

Appendix D SURFACE

Deposition and Surface Effects Computer Program (a plume impingement, deposition, abrasion, surface contamination effects model)

Appendix E N2H4

Monopropellant Combustion Chamber
Dynamics Program

This section concentrates on describing the operation of the integrated or controlling program <u>CONTAM</u>, and the interfacing of the various subprograms. The reader is referred to the appropriate appendix for detailed information concerning the operation of the individual subprograms.

The <u>CONTAM</u> program is written in FORTRAN IV. The primary version of the system includes the option of a monopropellant engine, subprogram N2H4, and requires 150,000 words of memory. However, a secondary version which only has a dummy N2H4 program, i.e., to be used with bipropellant engines only, can be run in only 120,000 words of memory. The cell loader directives for each of these editions are illustrated in this report. The program <u>SURFACE</u> has not been incorporated into <u>CONTAM</u> proper at this time for reasons previously explained in the report, and in its present form <u>SURFACE</u> requires 240,000 words of memory to execute. All aspects of the programming system have been written to run on a CDC 6000 series computer.

a. Combustion Chamber Contaminant Production

Unburned propellant and intermediate products of combustion (gas and liquid phase) ejected from the combustion chamber are considered first as a source of contaminants. Referring to Figure 11, the Transient Combustion Chamber Dynamics (TCC) subprogram or the monopropellant combustion chamber dynamic subprogram (N2H4) are used to generate contaminant production data. The results of the TCC or N2H4 subprograms are time dependent and require interface manipulation for subsequent modeling and analyses since the transport model treats the flow as steady state. After examination of the production of contaminants during the entire transient pulse, representative "time slices" are chosen so that gas and liquid properties may be averaged over the time intervals for use as input to the MULTRAN subprogram. The required output from TCC includes: chamber pressure, chamber temperature, droplet size distribution, droplet velocity, and mass flux of gas and droplets.

b. Contaminant Transport

Having defined the average amount of gas and liquid phase ejected from the combustor for each pulse segment, the two-phase nozzle and plume flow is then computed for each steady state pulse segment, using method-of-characteristics computer subprogram <u>MULTRAN</u> (subprograms TD2, TD2P, and SLINES are included in <u>MULTRAN</u>).

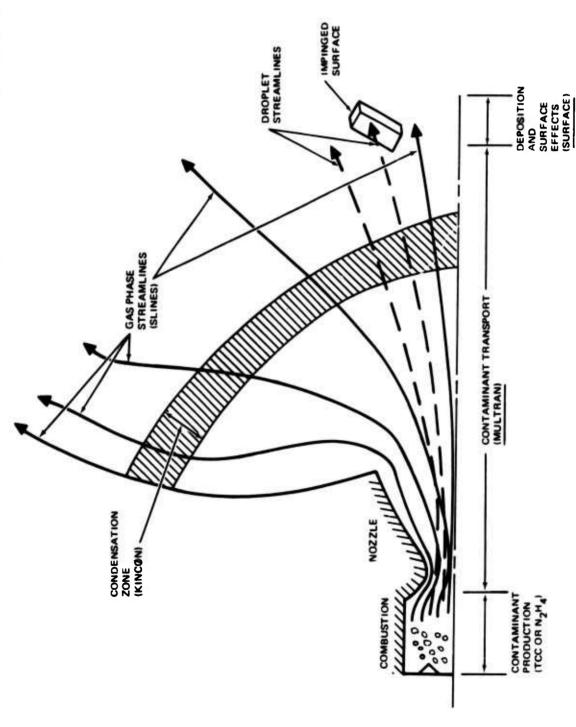


Figure 11. Schematic of Contamination Processes and Related Subprograms

Computer subprogram <u>SLINES</u> was developed to provide the necessary interface between the <u>TD2</u>, <u>TD2P</u> and the <u>KINCON</u> subprograms. Basically, <u>SLINES</u> interpolates between points on each characteristic line to provide exhaust gas properties for points on a streamline. A streamline is defined as that line which runs through the throat, nozzle, and plume, bounding a given constant percentage of the mass flow between it and the nozzle axis.

c. Chemical Kinetics and Condensation

In addition to unburned propellant droplets, many liquid-bipropellant exhausts contain condensed phase products of combustion as an important contaminant source. The formation of condensables in liquid propellant exhausts has been analytically modeled. The <u>KINCON</u> subprogram predicts the nozzle and plume condensation effects utilizing a classical nucleation and droplet growth model. It also provides the gas-phase chemical kinetics analysis along streamlines.

d. Deposition and Surface Property Effects

The flux of contaminants approaching a surface submerged in a bipropellant plume, as determined above, provides the starting point for the analysis of liquid and solid deposition on impinged surfaces. A model has been developed to account for the accommodation of momentum and energy upon impact of liquid and solid particles and to predict the amount and state (thin film, thick film, droplets, crystals, etc.) of the deposited materials. Damage and changes in surface properties due to mechanical abrasion and/or deposition are also treated in this subprogram, <u>SURFACE</u>. The surface property changes considered are absorptivity, emissivity, reflectivity, and transmissivity.

2. PROGRAM DESCRIPTION

This section describes the structure and logic of the Plume Contaminant Effects Prediction Computer Program, <u>CONTAM</u>. Particular emphasis is placed on the description of the main program, the cell loader structure, and the data interface between the various subprograms. Detailed descriptions of the subprograms may be found in the appropriate appendix.

The <u>CONTAM</u> program is structured so that any one subprogram may be run independently or any number of the subprograms may be run sequentially. Only the main program and required subprograms as defined by cell loader directive, reside in computer core during operation.

The main program (EXEC) was coded to perform the required selection of the various subprograms. It initializes certain logical control variables, accepts control variables through input, and provides overall logic control for the program. It also provides overlay communication.

3. PROGRAM STRUCTURE

The makeup of CONTAM is illustrated by Figure 12, a list of the various subprograms of the overall system.

4. PROGRAM DATA INTERFACES

The resultant data from the various subprograms reside on magnetic tape or disc file for use by subsequent subprograms.

The data interface has been designed so that the input and output data to a particular subprogram are preserved subsequent to the running of that subprogram, while at the same time, the number of lagical file units used are minimized by reusing logical files. This means that while the user may run each program independently but sequentially, if the data output from a particular subprogram is not acceptable for some reason, the input data file to that subprogram has been preserved so that the user has a "restart" capability without having to restart from the initial subprogram of the sequence.

a. TCC Data

Data output from <u>TCC</u> is written on to logical file 9 (in addition to the NAMELIST'S RESRT1 and RESRT2) for subsequent use in the cases of multiple pulses or reprintouts of the results. "Movie" information is stored on TAPE 13. In addition, <u>TCC</u> uses files 1, 16, and 48 internally. File 1 contains the variable data used by the plotting routine in the subroutine GRAPHS; files 16 and 48 are required for the system plotting package. A quick visual inspection of the final printout of <u>TCC</u> will allow the user to choose the correct inputs for MULTRAN.

b. N2H4 Data

When using the monopropellant option, the needed input for <u>MULTRAN</u> is obtained by quick visual inspection of the printed output from <u>N2H4</u>.

c. TDMAIN

TDMAIN is the driver program for TD2 and TD2P. It also defines the overlay structure for COMMON block storage used by both of these programs.

The input logical file number for the <u>TD2</u> program is 10. The <u>TD2</u> subprogram provides data output on two logical files; 8 and 12. Logical file 8 contains data required to be input to the <u>TD2P</u> subprogram. Logical file 12 provides input data for the <u>SLINES</u> subprogram and the deposition and surface effects subprogram, SURFACE.

Subprogram <u>TD2P</u> receives its input data from logical file 8. <u>TD2P</u> provides output data on logical files 9 and 12. File 9 contains radiation and

Figure 12. CONTAM Structure

force field data. File 12 contains the contaminant properties data. The subroutine which writes on file 12 is common to both <u>TD2</u> and <u>TD2P</u>. It is located in the main overlay level so that it is accessible to both subprogram overlay levels.

d. SLINES Data

The <u>SLINES</u> subprogram reads data from logical file 12 which has been generated by subprograms <u>TD2</u> and <u>TD2P</u>. It generates the streamline data and writes this data on logical file 8 for use by the KINCON subprogram.

e. KINCON

The KINCON subprogram uses five logical file units. The file unit numbers are 1, 4, 8, 11, and 12. Files 1 and 11 are scratch files used internally by the subprogram. File I contains the initial conditions and area ratio table to be used in the condensation calculation. File 11 contains reaction tables. Files 4 and 8 are input files. File 4 is an optional input which contains a list of thermal properties in JANAF format. Since logical file 4 is an optional input, if it is not used as such, it will default to an internally used file by the subprogram. If the logical file 4 input is not exercised, the thermal properties data must be input by punched cards (see Appendix C). The subprogram will then write JANAF thermal properties on logical file 4 from the punched card input. Obviously, with the appropriate control cards, file 4 may then be saved for subsequent use. File 8 contains the streamline properties required by the KINCON subprogram. Logical file 12 is the output tape of the subprogram and contains the multiphase and kinetic results to be used by the deposition and surface effects subprogram, SURFACE.

5. PROGRAM USER'S MANUAL

a. Input to CONTAM

Punched card input is required. Logical file 4 input is optional, but the option must be specified in the card input.

The punched card inputs required are of three types; NAMELISTS, some nonstandard format, and variable formats. Subsection 5. b describes the general nature of the data input through the various NAMELISTS and indicates the subprograms to which they apply. For a detailed description of the contents of the various NAMELISTS, the user should refer to the appropriate Appendix for each subprogram as listed in subsection of this section. A detailed description of the variable format inputs will also be found in the appropriate appendix. Section 5. d discusses the nonstandard format data and the required stacking (organization) of the punched cards.

Section V presents a sample case which includes a data listing, the resulting output, and a "day file" from the CDC 6500 computer system. The "day file" is included to illustrate the required system control cards and their proper sequence.

In addition to the aforementioned inputs, there are the data interfaces which are required by the various subprograms and discussed in Subsection 4 of this section. These interface data should be considered as inputs when running the subprograms independently.

b. Program NAMELISTS

Table I presents a list of the names of each <u>NAMELIST</u> and the subprograms to which they correspond. To determine the details and parameters contained in each <u>NAMELIST</u>, the user should refer to the appropriate Appendix as listed in Subsection 1 of this section. <u>NAMELISTS</u> required by <u>CONTAM</u> (the main program) are described in the Subsection 5. c.

c. <u>CONTAM NAMELISTS</u>

Two NAMELISTS are required by the main program, <u>CONTAM</u>, to provide control of the overall program. They are:

- (1) NAMELIST/NCASE/
- (2) NAMELIST/IPATH/

Table I NAMELIST AND SUBPROGRAM

NAMELIST Name	Subprogram
NCASE	CONTAM(EXEC)
IPATH	CONTAM(EXEC)
PROCED	TCC
PRTDIR	TCC
RESRT1	TCC
RESRT2	TCC
STEADY	STDST
ENGINE	N2H4
CATLST	N2H4
CONTRL	N2H4
DATA	TD2
DATAP	TD2P
SID	XMGKS(SLINES)
THERMO	TTAPE(KINCON)
PROPEL	INPUT(KINCON)
There are eight NAM to Appendix D	MELIST sets in SURFACE, refer

There is only one parameter in the <u>NAMELIST</u>/NCASE/. It is ICASE. ICASE is an integer variable which tells the program how many pulse segments are to result from the transient pulse output of the TCC subprogram. Each pulse segment is considered a "case." Each subsequent subprogram in the sequence must operate on each pulse segment (case), one segment at a time. The default value is ICASE = 1, which is recommended for most applications.

NAMELIST/IPATH/ contains ten variables. However, only two of the variables are required per run (sequential mode). The variable list for NAMELIST/IPATH is:

- TC
- BIPROP
- SSCP
- NOZZLE
- PLUME
- SLINES
- MGKS
- SURFAC
- KMODE
- NSL

Each variable except KMODE and NSL is logical. For example:

TC = T, BIPROP = T,

The above example indicates that the subprogram to be run will be TCC. If the data input were changed to

TC=T, BIPROP=F,

the program run would be N2H4. When running the MULTRAN portion of CONTAM, the user may desire to run sequentially during a single run the programs TD2, TD2P, and SLINES. In this instance, the data input would read

NOZZLE=T, SLINES=T.

In the case the subprogram KINCON is to be run, KMODE and NSL may also be required.

The variable KMODE is used only when the <u>KINCON</u> subprogram is run independently. If used, it is input as the integer variable "one." It indicates that certain options will be exercised in the operation of the <u>KINCON</u> subprogram. These options are described in Appendix C.

The variable NSL is required only when the <u>KINCON</u> subprogram is run independently. NSL indicates the number of streamlines that will be analyzed by the <u>KINCON</u> subprogram. If subprogram <u>SLINES</u> is run in

conjunction with the KINCON subprogram, any NSL value input here will be overridden by the value of NSL input for the SLINES subprogram.

d. Nonstandard Format Inputs

All of the nonstandard format punched card inputs are used by the KINCON subprogram. The names of these inputs are

- Thermodynamic Data (Optional)
- Title Card
- Species Cards
- Reaction Cards

The format for the data on these cards is detailed in Appendix C.

Section V SAMPLE CASE

This section presents a data listing and printed output for a sample case run in the sequential mode on the CDC 6500 computer system.

1. DEFINITION OF SAMPLE CASE

The sample case is chosen to illustrate the application of the successive subprograms of **CONTAM** for the prediction of the contamination effects resulting from a typical firing of a real rocket engine. The engine chosen for these computations is the Marquardt R-6C. The R-6C is a commercially available NTO-MMH rocket engine having a nominal thrust of 5 pounds and designed for pulse-mode operation. It is an excellent choice for experimental vacuum-chamber studies of contaminant production because of its small size, which makes it easier to maintain vacuum in the test chamber, and its short pulse capability which can be used to aggravate the production of contaminants. The effect of the small engine size, however, must be considered in designing the experiment. The R-6C is being used in an experimental contaminant effects study currently underway at NASA Lewis Research Center under the direction of Dr. Herman Mark. It is hoped that the computed values for contaminant production, transport, deposition, and surface effects can be experimentally verified by comparison with experiments such as those being run at NASA Lewis Research Center.

A pulse width of 17 milliseconds was chosen for these computations. This duration corresponds to a minimum impulse bit firing. Minimum impulse bit firings are particularly important for production of contaminant material, and are commonly employed for limit-cycle pointing corrections on vehicles which must maintain a prescribed attitude.

In our calculations, the engine is fired with its walls initially clean and with its dribble volumes both initially empty, but with the lines full of propellant behind the valves. This initial condition would be easier to match experimentally than any prescribed axial accumulation of fuel and oxidizer on the wall or partially filled dribble volumes. The chamber, injector, and tankage were initially set to room temperature values, again for ease of experimental comparison. The line lengths, line diameters, tank pressures, and other installation and operational variables were chosen to agree with the NASA Lewis vacuum chamber installation.

2. CONTAMINANT PRODUCTION - THE TCC PROGRAM

The <u>TCC</u> (transient combustion chamber) program calculates contaminant production by digital integration of the time-dependent engine processes, i.e., propellant flow, atomization, and combustion of the

injected droplets. The calculated two-dimensional trajectories for the burning droplets determine how much propellant is deposited on the wall, and how much passes through the throat unburned. The trajectories for the ejected unburned droplets are calculated up to the throat by the TCC subprogram, and in the nozzle and plume by the MULTRAN subprogram. The unburned propellant which is deposited on the combustion chamber wall is subjected to burnoff and axial flow from the action of the hot, fast-moving chamber gases. The amount of this wall film material which survives and passes through the throat is calculated by the TCC subprogram. Experimental firings of small pulsing engines show that much of this material accumulates on the nozzle lip during each firing, and is blown off during the succeeding start transient. The droplet size, initial direction of flight, and subsequent trajectory of this material may be defined statistically from experimental firings, but is outside of the present scope of the CONTAM program.

a. TCC Input-Sample Case (R-6C Engine)

Input data for the <u>TCC</u> program are broken down into several large blocks of related data. The block headings are: General Instructions, Time-Average Performance Values, Droplet Trajectory Plot, Flow Rate Overrides, Operating Conditions, Ignition Description, Fuel Feed System, Oxidizer Feed System, Atomization Parameters, Fuel Properties, Oxidizer Properties, Product Properties, Thrust Coefficient Table, Adduct Properties, Contaminant Viscosity, Valve Timing, Multi-Ring Injector, and Combustion Chamber Profile.

Loading such an extensive array of input information can never be easy; however, considerable effort has been made to make it as easy as possible. Wherever possible the input units chosen are those which are obtainable directly from a blueprint, engine test, literature reference, or standard engine performance computer program.

Certain of the data blocks, i.e., Fuel Properties and Oxidizer Properties, remain fixed for particular materials and can be passed on from one data set to another uncharged. The Product Property Deck for a particular propellant combination and pressure can also be reused when appropriate. Having such subdecks on hand for the common propellant constituents and combinations greatly facilitates loading the input for the TCC subprogram. When subdecks are created with the injector and chamber characteristics for particular engines, these can be used for a variety of operating conditions or for a variety of feed-system configurations, i.e., line lengths and diameters, restrictor areas, etc.

(1) Origin of Engine and Propellant Data

The known physical properties of nitrogen tetroxide and monomethylhydrazine were taken from the Battelle "Liquid Propellant Handbook" (Reference 7) and the Aerojet publication "Performance and Properties of Liquid Propellants" (Reference 8). The burning-rate coefficient for monomethylhydrazine was estimated to be halfway between the experimental values given for hydrazine and UDMH by Dykema and Greene (Reference 9). The burning-rate coefficient for NTO was calculated using Godsaves' equation. The equilibrium combustion gas properties of chamber temperature,

mean molecular weight, gamma, and vacuum thrust coefficient for an expansion area ratio of 40 were calculated, using the MDAC thermochemistry program H099 and the JANNAF (Reference 10) values for heats of formation. The chamber dimensions, injector parameters, and valve ramp durations come from the manufacturer, while the feed system values were supplied by NASA Lewis Research Center.

The ignition parameters, activation energy, and frequency factor multiplied by heat of reaction are the experimental values measured for NTO and MMH by Seamans, Vanpee, and Agosta (Reference 11). The fuel and oxidizer droplet size correction factors are taken from Rocketdyne Report R-8455 by L. J. Zajac (Reference 12). The fuel and oxidizer fan lengths are taken to be the same multiple of orifice diameter as for another motor where this value could be taken from NACA TN 3835 (Reference 13), and then experimentally verified by measuring chugging frequency and chugging amplitude. The distance to single-stream breakup is taken to be the same multiple of orifice diameter as shown in the photographs of NACA TN 3835. The flash cone angle was taken from photographs of flashing streams in the thesis of Brown (Reference 14). The drop size distribution is from NACA TN 4222 (Reference 15). The fraction of propellant hitting the wall which is presumed to stick is suggested by film-coolant injection effectiveness values published by B. L. McFarland (Reference 16). (Using a value of 1.0 for the fraction sticking when the film coolant is injected tangentially and a value of 0.5 for nontangential injections appears to give good agreement with experiment.) Experimental chamber wall temperature histories for small motors have also been published by McFarland (Reference 17).

The decomposition temperature for the MMH nitrate was taken from the paper of Perlee, Christos, Miron, and James (Reference 18). The values used for the density, vapor specific heat, latent heat and viscosity of the MMH nitrate and its solutions, and the accommodation coefficients for MMH and NTO are estimated values since no experimental values have ever been published.

The details of the fuel and oxidizer feed systems were unknown, while the flow rates and overall pressure drops were known. For this reason equivalent feed system orifice diameters were obtained using the flow override option. These values were obtained in an earlier computer run and are incorporated into the data input used for this run.

(2) Input Data for TCC Subprogram (Marquardt R6C Engine)

Title information is input via a variable format on the first card of the data deck. At present a title 58 characters long may be input.

The first card in the data for the TCC program is a variable format card which is used to introduce the hollerith information used as a printed heading for the TCC data (see Table II).

(IHO, 10X, * MARQUARDT R-6C ENGINE, SAMPLE CASE FOR CONTAM *)

This card begins in card column 1 and extends to card column 70 inclusive. The hollerith title is inserted between the asterisks.

Table II
INPUT DATA FOR MARQUARDT R-6C ENGINE TCC

DDD1 Array Number	Variable	Value	Comments/Source
	GENERAL IN	STRUCTIONS	
9	Punch for restart flag (1.0 or 0.0)	1.0	Punch a card deck giving final distribution of propellant mass on the wall i.e. RESRT1 & RESRT2
10	Work from tape flag (1.0 or 0.0)	0.0	Calculate chamber processes instead of taking values from a prewritten output tape
11	K BUG flag	0.0	Do not print a large set of diagnostic values at the K BUGTH iteration
12	No movie tape flag (1.0 or 0.0)	0.0	Write an output tape to be used to make chamber movies
201	Stop time	0.03 sec	Terminate calculations at a model time of 0.030 seconds
202	Time interval	0.0001 sec	Use an integrating time interval of 0.0001 (based upon previous experience
203	Print one out of	10	A propellant disposition summary is printed for every tenth time interval
204	Plot one out of	30	A wall-film thickness profile is plotted for every 30th time interval
205	Film vertical exaggeration	20	The wall-film thickness is exaggerated by a factor of 20 for the combustion chamber movie

Table II
INPUT DATA FOR MARQUARDT R-6C ENGINE TCC (Continued)

DDD1 Array Number	Variable	Value	Comments/Source
206	Start movie iteration	150	Motion picture frames will be produced starting with the 150th iteration
207	End movie iteration	150	Motion picture frames will be produced ending with the 150th iteration
208	Data review only flag	0.0	The program will com- pute the chamber proc- esses instead of stopping after printing out the input data set and values derived from the flow overrides
	TIME-AVERAGE PEI	RFORMANCE	VALUES
313	Start time 1	0.0 sec	The first time slice for average values starts at 0.0 sec
314	Finish time 1	0.015	The first time slice for average values ends at 0.015 sec
315	Start time 2	0.015 sec	The second time slice for average values starts at 0.015 sec
316	Finish time 2	0.017 sec	The second time slice for average values ends at 0.017 sec
317	Start time 3	0.017 sec	The third time slice for average values starts at 0.017 sec
318	Finish time 3	0,030 sec	The third time slice for average values ends at 0.030 sec

Table II
INPUT DATA FOR MARQUARDT R-6C ENGINE TCC (Continued)

DDD1 Array Number	Variable	Value	Comments/Source
	DROPLET TRA	JECTORY PL	.OT
209	Fuel trajectory group	3.0	A trajectory will be plotted for the third fuel size group
210	Oxidizer trajectory group	0.0	Only one droplet can be plotted per run. Either fuel or oxidizer depending upon whether 209 or 210 is given a non zero value
211	Trajectory start time	0.015/sec	The fuel droplet tra- jectory will be plotted for the fuel droplet injected when the model time is 15 milliseconds
212	Injector ring	1.0	The fuel droplet tra- jectory is for a droplet originating from the first ring
	FLOW RAT	E OVERRIDES	5
213	Fuel flow rate	0.0 lb/sec	No fuel feed system values are being calculated. (Flow rate at which experimental pressure drops are known)
214	Total pressure drop	0.0 psi	Fuel-side orifice diam- eter is not being calculated (fuel tank pressure — chamber pressure)
215	Valve pressure drop	0.0 psi	Fuel valve port area is not being calculated

Table II
INPUT DATA FOR MARQUARDT R-6C ENGINE TCC (Continued)

DDD1 Array Number	Variable	Value	Comments/Source
216	Injector pressure drop	0, 0 psi	Injector orifice discharge coefficients are not being calculated
61	Percent of total fuel flow which passes through ring 1	0.0	Injector orifice discharge coefficients are not being calculated
62	Percent of total fuel flow which passes through ring 2	0.0	Injector orifice discharge coefficients are not being calculated
63	Percent of total fuel flow which passes through ring 3	0.0	Injector orifice discharge coefficients are not being calculated
217	Oxidizer flow rate	0.0 lb/sec	No oxidizer feed system values are being calculated
218	Total pressure drop	0.0 psi	Oxidizer-side orifice diameter is not being calculated
219	Valve pressure drop	0.0 psi	Oxidizer valve port area is not being calculated
220	Injector pressure drop	0.0 psi	Injector orifice dis- charge coefficients are not being calculated
85	Percent of total oxidizer flow which passes through ring l	0.0	Injector orifice dis- charge coefficients are not being calculated
86	Percent of total oxidizer which passes through ring 2	0.0	Injector orifice dis- charge coefficients are not being calculated
87	Percent of total oxidizer which passes through ring 3	0.0	Injector orifice dis- charge coefficients are not being calculated

Table II
INPUT DATA FOR MARQUARDT R-6C ENGINE TCC (Continued)

DDD1 Array Number	Variable	Value	Comments/Source
	OPERATIN	G CONDITIONS	
13	Fuel tank pressure	180 psia	NASA Lewis
14	Fuel tank temperature	294°K	Approximate room temperature
15	Fuel tank psi/sec	0.0	Fuel tank pressure is not rising or falling
16	External pressure	1 x 10 ⁻⁶ psia	Estimated vacuum environment
17	Oxidizer tank pressure	165. psia	NASA Lewis
18	Oxidizer tank temperature	294°K	Approximate room temperature
19	Oxidizer tank psi/sec	0.0	Oxidizer tank pressure is not rising or falling
21	Injector initial temperature	294.°K	Approximate room temperature
22	Throat initial temperature	294°K	Approximate room temperature
25	Injector maximum temperature	350°K	Estimated
26	Half-rise time	10.0 sec	Estimated
27	Injector minimum temperature	294.°K	Approximate room temperature
28	Half-fall time	100.0 sec	Estimated
29	Throat maximum temperature	1,400°K	Estimated, based on values for similar engines

Table II
INPUT DATA FOR MARQUARDT R-6C ENGINE TCC (Continued)

DDD1 Array Number	Variable	Value	Comments/Source
30	Half-rise time	10 sec	Estimated-see McFarland for experi- mental curves of similar engines (Reference 17)
31	Throat minimum temperature	294°K	Approximate room temperature
32	Half-fall time	100.0 sec	Estimated
	IGNITION	DESCRIPTION	
33	Assigned ignition delay	0.0 sec	Not used when chemi- cal kinetic values are specified
34	Igniter port location	0.0 in. from injector	No igniter used
35	Igniter fuel flow rate	0.0 lb/sec	No igniter used
36	Igniter oxidizer flow rate	0.0 lb/sec	No igniter used
37	Activation energy	5,200 cal/ mole	Seamans, Vanpee, and Agosta (Reference 11)
38	Frequency factor x heat of reaction	3.4 x 10 ¹⁴ (cc/mole sec) x (cal/mole)	Seamans, Vanpee, and Agosta
39	Perfect mixing flag	0.0	Use normal ignition calculations
40	No axial mixing flag	0.0	Use normal ignition calculations
	FUEL F	EED SYSTEM	
41	Fuel line length	480 in.	NASA Lewis Research Center
42	Fuel line diameter	0.189151 in.	NASA Lewis Research Center

Table II
INPUT DATA FOR MARQUARDT R-6C ENGINE TCC (Continued)

DDD1 Array Number	Variable	Value	Comments/Source
43	Fuel restrictor diameter	0.01672% in.	Calculated from pressure drop and flow rate values
44	Fuel venturi diameter	0.0 in.	No venturi used
45	Fuel valve port area	0.0281 in. ²	No valve restriction when open — line area used
46	Fuel check valve flag	0.0	Reverse flow in feed system is possible
47	Fuel valve opening ramp duration	0.001 sec	Manufacturer
48	Fuel valve closing ramp duration	0.001 sec	Manufacturer
57	Fuel initial void volume	0.00113 in. ³	Set equal to dribble volume for initially empty condition
59	Fuel transition volume	0.0 in. ³	No hot fuel in injector
60	Fuel dribble volume	0.00113 in. ³	Manufacturer
	OXIDIZER I	FEED SYSTEM	
65	Oxidizer line length	480 in.	NASA Lewis Research Center
66	Oxidizer line diameter	0.189151 in.	NASA Lewis Research Center
67	Oxidizer restrictor diameter	0.020997 in.	Calculated from pressure drop and flow rate values
68	Oxidizer venturi diameter	0.0	No venturi used
69	Oxidizer valve port port area	0.0281 in. ²	No valve restriction when open — line area used

Table II
INPUT DATA FOR MARQUARDT R-6C ENGINE TCC (Continued)

DDD1 Array Number	Variable	Value	Comments/Source
70	Oxidizer check valve flag	0.0	Reverse flow in oxidizer feed system is possible
71	Oxidizer valve open- ing ramp duration	0.001 sec	Manufacturer
72	Oxidizer valve closing ramp duration	0.001 sec	Manufacturer
81	Oxidizer initial void volume	0.000580 in. ³	Set equal to dribble volume for initially empty condition
83	Oxidizer transition volume	0.0 in. ³	No hot oxidizer in injector
84	Oxidizer dribble volume	0.000580 in. ³	Manufacturer
89	Fuel drop factor	0.50	Droplet size correction factor for injector orifices which are nonturbulent, not flowing full or noncircular. L. J. Zajac's value for laminar flow (Reference 12)
90	Oxidizer drop factor	0.50	Droplet size correction factor for injector orifices which are non turbulent, not flowing full or norcircular. L.J. Zajac's value for laminar flow
91	Fuel fan length with 90-deg impingement, balanced momentum	3.0 orifice diameters	Estimated from NACA TN 3835. (Reference 13). No experimental data for orifices this small

Table II
INPUT DATA FOR MARQUARDT R-6C ENGINE TCC (Continued)

DDD1 Array Number	Variable	Value	Comments/Source
92	Oxidizer fan length with 90-deg impinge- ment, balanced momentum	3.0 orifice diameters	Estimated from NACA TN3835. No experimental data for orifices this small.
93	Hold at triple point flag	0.0	Assume flashing propellant equilibrate with chamber pressure even though freezing occurs
94	No initial dribble flag	1.0	The quill-type Marquardt injector is not likely to dribble during start before the dribble volume fills completely
95	Flash cone angle	30 deg	Apex angle of flashing liquid spray taken from photographs of R. Brown (Reference 14)
96	Single-stream breakup distance given in orifice diameters	10	Estimated from photographs of single- stream breakup. See NACA TN3835 (Refer- ence 13)
97-101	Drop size distribution table	0.198, 0.759 1.0, 1.23, 2.3045	Taken from experimental values of NACA TN 4222 (Reference 15)
102	No wall breakup flag	0.0	Streams are assumed to atomize on wall impact
103	Drop rebound velocity ratio	1.0	Droplets which bounce off wall are assumed perfectly elastic

Table II
INPUT DATA FOR MARQUARDT R-6C ENGINE TCC (Continued)

DDD1 Array Number	Variable	Value	Comments/Source
104	Fraction sticking	0.5	One-half the streams and droplets impacting with the cold wall are assumed to bounce and the remainder stick. B. L. McFarland data (Reference 16)
105	No fuel flash flag	0.0	The fuel stream is permitted to flash atomize when the correlations indicate that it should
106	No oxidizer flash flag	0.0	The oxidizer stream is permitted to flash atomize when the correlations indicate that it should
107	No entrainment flag	0.0	The material deposited on the chamber walls is permitted to entrain when the correlations indicate that it should
108	Delete droplet means flag	0.0	The D ₃₀ , D ₃₁ , and D ₃₂ will be calculated for the chamber droplet population at each time interval
109	Fuel normal boiling point	360°K	Aerojet compilation (Reference 8)
110	Fuel freezing point	222°K	Aerojet compilation
111	Fuel critical temperature	594°K	Aerojet compilation
112	Fuel critical pressure	l, 195 psia	Aerojet compilation

Table II
INPUT DATA FOR MARQUARDT R-6C ENGINE TCC (Continued)

DDD1 Array Number	Variable	Value	Comments/Source
113	Fuel vapor specific heat at film temperature	0.995 cal/ gram °K	Value for propylene specific heat at 1500°K extrapolated from NBS C461. Structure similar to MMH. No data for MMH
114	Fuel liquid specific heat at 300°K	0.69 cal/ gram °K	Aerojet compilation
116	Fuel vapor molecular weight	46.074 grams/ gram mole	Aerojet compilation
117	Fuel latent heat of vaporization (at normal bioling point)	210 cal/gram	Aerojet compilation
118	Fuel latent heat of fusion	67.5 cal/gram	Battelle handbook (Reference 7)
119	Fuel liquid thermal conductivity	0.000545 cal/ cm °K	Aerojet compilation
120	Fuel accommodation coefficient	1.0	Strawman value, no data available
121	Reference temperature for fuel properties	300°K	Experimental values available at this temperature
122	Fuel density at reference temperature	0.88 gram/cc	Aerojet compilation
123	Fuel viscosity at reference temperature	0.0104 poise	Aerojet compilation
124	Fuel surface tension at reference temperature	47 dynes/cm	Battelle handbook
125	Fuel burning rate coefficient	0.0325 cm ² /	Dykema and Greene (Reference 9)

Table II
INPUT DATA FOR MARQUARDT R-6C ENGINE TCC (Continued)

DDD1 Array Number	Variable	Value	Comments/Source
126	Fuel monopropellant intercept (A)	0.0 cm/sec	Strand burning tests fitted r = A + B P _C with r in cm/sec; P _C in psia
127	Fuel monopropellant coefficient (B)	0.0 cm/ sec psi ⁿ	MMH does not burn in liquid strand tests
128	Fuel monopropellant exponent (n)	0.0	
	OXIDIZER	PROPERTIES	
129	Oxidizer normal boiling point	294 °K	Aerojet compilation
130	Oxidizer freezing point	262 °K	Aerojet compilation
131	Oxidizer critical temperature	431 °K	Aerojet compilation
132	Oxidizer critical pressure	l,470 psia	Aerojet compilation
133	Oxidizer vapor specific heat at film temperature	0.298 cal/ gram °K	JANNAF tables for NO ₂ (Reference 10)
134	Oxidizer liquid specific heat at 300°K	0.36 cal/ gram °K	Aerojet compilation
136	Oxidizer vapor molecular weight	46.008 gram/ gram mole	Vapor mostly NO ₂ at high temperature or low pressure
137	Oxidizer latent heat of vaporization (at normal bioling point)	99.0 cal/gram	Aerojet compilation
138	Oxidizer latent heat of fusion	39.2 cal/gram	Battelle handbook

Table II
INPUT DATA FOR MARQUARDT R-6C ENGINE TCC (Continued)

DDD1 Array Number	Variable	Value	Comments/Source
139	Oxidizer liquid thermal conductivity	0.000306 cal/ cm °K	Aerojet compilation
140	Oxidizer accommodation coefficient	1.0	Strawman value, no data available
141	Reference tempera- ture for oxidizer properties	300°K	Experimental values available at this temperature
142	Oxidizer density at reference temperature	1.45 gram/cc	Aerojet compilation
143	Oxidizer viscosity	0.00446 poise	Aerojet consp.lation
144	Oxidizer surface tension	28 dyne/cm	Battelle handbook
145	Oxidizer burning rate coefficient	0.027 cm ² / sec	Calculated from Godsaves' equation
146	Oxidizer mono- propellant intercept (A)	0.0 cm/sec	Strand burning rate fitted to: $r = A + B p_c^n$ with r in cm/sec and P_c in psia
147	Oxidizer mono- propellant coefficient (B)	0.0 cm/sec psia	NTO does not burn in liquid strand tests
148	Oxidizer mono- propellant exponent (n)	0.0	NTO does not burn in liquid strand tests
	PRODUCT I EQUILIBRIUM GA	PROPERTIES S TEMPERATUI	RE
	Equilibrium combustion gas temperature at fuel fractions 0.0, 0.1, 0.2, 1.0	300; 2103 3084; 3397; 3061; 2368; 1705; 1433; 1344; 1266; 1190; in ^o K	From standard equili- brium thermochemistry calculations

Table II
INPUT DATA FOR MARQUARDT R-6C ENGINE TCC (Continued)

ODD1 Array Number	Variable	Value	Comments/Source
	EQUILIBRIUM GAS	MOLECULAR WE	CIGHT
161-171	Equilibrium combustion gas mean molecular weight at fuel fractions 0.0, 0.1, 0.2, 1.0	46.008, 28.79, 26.41, 23.39, 19.88, 16.75, 14.41, 13.91, 14.00, 14.10, 14.29	From standard equi- librium thermo- chemistry calculations
	EQUILIBRI	UM GAS GAMMA	
173-183	Equilibrium combustion gas gamma at fuel fractions 0.0, 0.1, 0.2, 1.0	1. 120, 1. 252, 1. 220, 1. 217, 1. 235, 1. 268, 1. 309, 1. 299, 1. 270, 1. 247, 1. 228	From standard equi- librium thermo- chemistry calculations
		EFFICIENT TABLI HRUST COEFFICI	
221-231	Vacuum thrust coefficients for the correct nozzle expansion area ratio of the motor, and fuel fractions of 0.0, 0.1, 0.2, 1.0	1.9240, 1.8028, 1.9082, 1.9617, 1.8470, 1.8122, 1.8680, 1.9331, 1.9294, 1.9224, 1.8959	obtained from thermo chemical calculations assuming equilibrium
232	Nozzle expansion area ratio	40	Manufacturer
	ADDUC	r properties	
185	Contaminant mixture density	l gram/cc	Strawman value, no experimental data
186	Contaminant vapor specific heat	l cal/gram °K	Strawman value, no experimental data
187	Contaminant latent heat of vaporization	100 cal/gram	Strawman value, no experimental data

Table II
INPUT DATA FOR MARQUARDT R-6C ENGINE TCC (Continued)

DDD1 Array Number	Variable	Value	Comments/Source
188	Contaminant decomposition temperature	500°K	Perlee, Christos, Miron, and James (Reference 18)
	CONTAMIN	ANT VISCOSITY	
189-199	Contaminant mix- ture viscosity at fuel fractions 0.0, 0.1, 0.2 - 1.0	0.00446, 0.024, 0.043, 0.068, 0.081, 0.100, 0.082, 0.064, 0.046, 0.029, 0.0104 poise	Experimental values for pure fuel and pure oxidizer. Strawman values for MMH nitrate mixtures
	VALV	E TIMING	
233	Fuel valve 0.0 sec Time of first ropening time opening		Time of first motion opening
234	Oxidizer valve opening time	0.0 sec	Time of first motion opening
235	Fuel valve closing time	0.017 sec	Time of first motion closing
236	Oxidizer valve closing time	0.017 sec	Time of first motion closing
237-264	Valve timing for 2nd, 3rd, 4th, 5th, 6th, 7th, and 8th pulses		Not being used
	FIR	ING INJECTOR ST RING L HOLES	
49	Fuel hole diameter	0.0158 in.	Manufacturer
50	Fuel hole length	0.0625 in.	Estimated from engine drawing
51	Axial location of fuel hole	0.0 in.	Approximately flush with injector face

Table II
INPUT DATA FOR MARQUARDT R-6C ENGINE TCC (Continued)

DDD1 Array Number	Variable	Value	Comments/Source
52	Radial location of fuel hole	0.045 in.	Estimated from engine drawing
53	Fuel radial injection angle	-45 deg	Manufacturer
54	Discharge coefficient	1.0	Assumed value
55	Number of fuel holes in this ring	1.0	Manufacturer
56	Transverse angle	0.0	Manufacturer
	OXII	DIZER HOLES	
73	Oxidizer hole diameter	0.0186 in.	Manufacturer
74	Oxidizer hole length	0.0625 in.	Estimated from engine drawing
7 5	Axial location of oxidizer hole	0.0 in.	Approximately flush with injector face
76	Radial location of oxidizer hole	-0.045 in.	Estimated from engine drawing
77	Oxidizer radial injection angle	45 deg	Manufacturer
78	Discharge coefficient	1.0	Assumed value
79	Number of oxi- dizer holes this ring	1.0	Manufacturer
80	Transverse angle	0.0	Manufacturer

Table II
INPUT DATA FOR MARQUARDT R-6C ENGINE TCC (Continued)

DDD1 Array Number	Variable	Value	Comments/Source			
	SECOND RING					
265-272		Second fuel ring values equivalent to 49-56 for first fuel ring (This engine has no second ring)				
273-280	Second oxidizer ring values equivalent to 73-80 for first oxidizer ring (This engine has no second ring)					
		THIRD RING				
281-288	Third fuel ring values equivalent to 49-56 for first fuel ring (This engine has no third ring)					
289-296	Third oxidizer ring values equivalent to 73-80 for first oxidizer ring (This engine has no third ring)					
	COMBUSTI	ON CHAMBER PROF	ILE			
297	Axial location of injector face	0.0 in.				
298	Diameter corresponding to above station	0.41735 in.	Scaled from print of the chamber			
299-310	Six pairs of stations and diameters, describing the chamber contour	0.25, 0.41735, 0.5, 0.41735, 0.75, 0.41735, 0.95872, 0.41735, 0.97363, 0.30607, 1.01436, 0.22461	Scaled from print of the chamber			
311	Axial location of throat	1.07 in.				
312	Throat diameter	0.19479 in.				

The input data are introduced through two namelists, Namelist PROCED and Namelist PRTDIR. Namelist PROCED consists of a 319-word array DDD1. These values contain the detailed description of the motor, propellant, operating conditions, etc. Namelist PRTDIR consists of an 80-word array DDD2. These values are nonzero or zero to indicate which time-varying parameters are to be plotted as graphic output.

The input values to Namelist PROCED are printed out by the program along with headings in logical groupings which have been found to be convenient. The input values will be described in the order of this print, rather than in ascending order, as this will be a more logical presentation.

Namelist PRTDIR directs the program to produce graphics of the time-varying parameters which are desired. The default value for the members at this array is zero, and graphics are produced only for members which are given nonzero values. The graphics which can be produced are:

DDD2 array

Number Variable to be Plotted 1. Chamber pressure 2. Fuel valve trace (fraction of wide-open port area) 3. Oxidizer valve trace (fraction of wide-open port area) Fuel flow rate leaving tank 4. 5. Oxidizer flow rate leaving tank Total propellant flow rate leaving tanks 6. 7. Fuel injection rate Oxidizer injection rate 8. Fuel (streams, fans, and droplets) mass in chamber 9. Oxidizer (streams, fans, and droplets) mass in chamber 10. Propellant (streams, fans, and droplets) mass in chamber 11. Fuel mass on chamber wall 12. Oxidizer mass on chamber wall 13. 14. Propellant mass on chamber wall Gas mass in chamber 15. Mass fraction of the chamber gas which is fuel derived 16.

17.	Chamber temperature (gas phase)
18.	Fuel evaporation rate
19.	Oxidizer evaporation rate
20.	Propellant evaporation rate
21.	Gas outflow rate
22.	Fuel droplet outflow rate
23.	Oxidizer droplet outflow rate
24.	Total droplet outflow rate
25.	Fuel film outflow rate
26.	Oxidizer film outflow rate
27.	Total film outflow rate
28.	Mass fraction of total outflow which is gas phase
29.	All chamber contents - total explosion pressure capability
30.	Mass of unatomized fuel streams and fans in chamber
31.	Mass of unatomized oxidizer streams and fans in chamber
32.	Mass of fuel droplets in chamber
33.	Mass of oxidizer droplets in chamber
34.	Gas and droplets only - total explosion pressure capability
35.	Chamber wall adduct only — chamber explosion overpressure capability
36.	D30 of all fuel droplets in chamber
37.	D31 of all fuel droplets in chamber
38.	D32 of all fuel droplets in chamber
39.	D30 of all oxidizer droplets in chamber
40.	D31 of all oxidizer droplets in chamber
41.	D32 of all oxidizer droplets in chamber
42.	Fuel injector void volume
43.	Oxidizer injector void volume
44.	Thrust

b. TCC Output-Sample Case (R-6C Engine)

The computer output is in the form of printout and computer graphics done on a Stromberg SD-4060 microfilm recorder operating in a SD-4020 emulation mode. Many of the output values are presented both in the form of printout and in the form of plots of variables versus time. It is generally easier to use the plots to follow the course of events in the chamber, but it is easier to use the printout to obtain exact numerical values, exact timing relationships, etc.

Since the complete set of graphics is expensive to produce in terms of computer time, it is best initially to request only the graphics which are really needed, plus chamber pressure, fuel injection rate and oxidizer injection rate (the three graphics which give the greatest amount of general information about the firing). If physical tape 9 and tape 13 are mounted before the computer calculations are done, they will record enough time-varying data that any additional graphics, additional time-slice average performance values, or motion picture frames that are desired can be recovered later from the tapes, without any additional combustion chamber calculations being required.

The information which is typically desired about the principal portions of a rocket engine firing is the duration, total impulse, fuel and oxidizer consumption and total contaminant production during the start transient; the same parameters for the cutoff transient; and the thrust level, fuel and oxidizer flow rates and contaminant production rate during the steady-state portion. It is obvious that for firings with sufficiently similar initial conditions such information can be used to estimate the mean performance level and cumulative contamination effects of longer or shorter pulses or of trains of pulses. For example, the total impulse of a pulse is simply the lumped impulse of the start transient plus the lumped impulse of the cutoff transient plus the steady-state duration multiplied by the steady-state thrust level. Total pulse specific impulse is obtained as the ratio of total pulse impulse to total pulse propellant consumption.

This segmental breakdown information cannot easily be obtained in a single pass through the computer, because it is not yet possible to describe "steady-state" in mathematical terms which will permit the computer to make satisfactory interval selections by itself (In typical pulsemode firings, mathematical steady-state is never reached; however, the major parameters are sufficiently slow-varying that "steady-state" assumptions are useful engineering approximations. The procedure used is to perform the chamber calculations, recording data on tape 9 and tape 13. Next, inspect the graphics which are produced describing chamber pressure, propellant injection rates, wall accumulation, etc., and choose a time for the end of the start transient. Tape 9 can then be remounted, and the program instructed to work-from-tape in calculating interval values for the designated start transient, for the steady-state portion of the firing, and for the cutoff portion. Additional graphics, wall-film profiles, or combustion chamber "snap shots" or movies may be obtained in the same way. (Because of the details of the programming the only droplet trajectory which can be obtained from tape, is the single one specified in the original computer run, other trajectories can be estimated from the chamber "snapshots" however.)

The computer printout includes a recapitulation of the input values and the derived fuel and oxidizer properties versus temperature, followed by the values of the major system variables computed at each computing time interval. At each tenth time interval, there is a summary print which gives the disposition of the total mass of each propellant constituent injected up till that time, i.e., how much has been ejected in the gas phase, as droplets, or as wall film and how much is being retained as gas, as droplets, or as wall film.

Statements are printed to give the times when the oxidizer manifold fills, when the fuel manifold fills, when ignition occurs, and when extinguishment occurs. There are several error diagnostics programmed, but none are illustrated in the sample case.

A fairly detailed summary is given at the end of the computation, which includes mean values for the entire pulse. Entire pulse values are given for mixture ratio, C-Star, and specific impulse for three possible mass bases: propellant mass out of the tank, propellant mass through the injector, and propellant mass through the nozzle. The final summary also gives the propellant droplet size distribution for injected and ejected fuel and oxidizer, the fraction of injected propellant which is expelled or retained in seven different categories. The axial and radial mean velocities of the expelled droplets, and the axial distribution of unburned propellant deposited on the walls.

Approximately the same data can be obtained for any specified timeslice by using the data on tape 9 in a program rerun from tape. The timeslice calculations also give values to use as input to the MULTRAN Subprogram of CONTAM.

(1) Computer Printout

Printouts are reproduced illustrating the recapitulated input data for the R6-C engine, the curve-fitted values for fuel and oxidizer physical properties, the derived feed system values, the chamber description at the start of the run and exerpts from the voluminous time-varying description of the firing. Prints specifying manifold priming, ignition, extinguishment, and periodic propellant disposition summaries are illustrated.

A time-averaged interval print for the time-slice between 15 and 17 milliseconds is illustrated. Thrust, chamber pressure, propellant flow rate, mixture ratio, C-Star, I_{sp}, contaminant production, expelled droplet mean diameters, axial and radial velocity components are given. The final printout gives input values for the MULTRAN subprogram, which follows TCC in the CONTAM ensemble. Chamber pressure, chamber temperature, gas constant, specific heat, gamma, gas viscosity, and the densities, amounts and droplet size distribution are given in units compatible with MULTRAN.

(2) Computer Graphics

Only a small fraction of the computer graphics which can be called for are illustrated here. There are four general classes of graphics, the droplet trajectory plots, the time-varying parameter plots, the wall-film thickness plots, and frames from chamber movies.

The droplet trajectory plot shows the path followed by the fuel droplet of the third size group (median fuel droplet) injected 15 milliseconds after start from the first ring (the only ring in this motor). This graphic is drawn so as to fill the frame, and hence will show details of even small motions of the droplet in either the x or y direction. Because of this expansion to fill the frame, there is usually distortion because the x and y scales are generally not expanded to the same extent. Plots of x vs. t, y vs. t and y vs. x are produced, but only the y vs. x plot is illustrated here.

The time-varying parameter plots illustrated here are chamber pressure (psia), fuel injection rate (lb/sec), oxidizer injection rate (lb/sec), gas outflow rate (lb/sec), outflow rate for incompletely burned fuel droplets (lb/sec), outflow rate for incompletely burned oxidizer droplets (lb/sec), outflow rate for fuel film on the wall (lb/sec), and outflow rate for oxidizer film film on the wall (lb/sec).

Wall-film thickness plots are shown for T = 6.0 milliseconds and T = 9.0 milliseconds. As in the case of the droplet trajectory plots, these are expanded to fill the frame, and thus are distorted by having a very large vertical exaggeration. This makes it possible to see details of the film which would not be visible otherwise.

The final type of graphic is a "motion picture frame," showing the outline of the chamber, the location of the droplet groups in the chamber and the (exaggerated) deposit of propellant film on the chamber wall. "Stills" or motion pictures of this kind are very valuable for aiding in the visualization of the very complex sequence of events which take place during the start and cutoff transients.

3. CONTAMINANT TRANSPORT-THE MULTRAN PROGRAM

The programs composing MULTRAN, TD2, TD2P, and SLINES are used to calculate the two-phase transonic and supersonic flow. The reader is referred to Appendix B for a detailed discussion of the computation methods.

The printout "Input Values for MULTRAN" of TCC supplies the input data for MULTRAN in the correct units, and this information has been used to generate the NAMELIST/DATA/, as illustrated. Using the results of the RC-o engine, a calculation has been made for the fuel droplets. The given value for the ratio of droplet mass to gas mass has been taken directly from TCC, i.e. 0.123. However, some discussion is required in relation to the choice of droplet size used in the calculation.

THE INPUT DATA FUR THIS CASE ARE AS FOLLOWS

PROBLEM DESCHIPTION

MARGUARDT WEAC ENGINE. SAMPLE CASE FOR CONTAM

GENERAL INSTRUCTIONS

FUNCH FOR RESTART BORK FROM TAPE KAUG 191 C.	\$100 TIME TIME INTERVAL PHINT ONE OF THE POLICY CASSOCIATION (202) .000100 (203) 10.000000 (204)	FILM VERT, EMAE, START MOVIE ITER, LMD MOVIE ITER, DATA STYTEM CAIV (20%) 28.606666 (20%) 150.000000 (20%) 150.000000 (20%) 0.000000	TIME-AVERAGE PERFORMANCE VALUES	START TIME 1 FILTSH TIME 1 START TIME 2 FILTSH TIME 2 (313) 0.00500 (314) .015000 (315) .015000 (315)
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DROPLET TRAJECTORY PLOT

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0.000000 0.0000.0 INJECT MAK, TEMP MALF-RISE TIME INJECT MIN, TEMP MAIF-FALL TIME (25) 350-000000 (24) 10-000000 (27) 294-700000 (28) 100-000000 FUEL TANK PRESS FUEL TANK TEMP FUEL TANK PSI/SEC EXTERNAL PRESSURF (13) 180.000000 (14) 200.00000 (15) 0.000000 (16) .000000 INJECT INIT TEMP THROAT INIT TEMP (21) 204.000000 (22) 244.000000 (23) 0.000000 (24) OPERATING CONDITIONS



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PRODUCT PROPERTIES

TABULAR PROPERTY DATA IS FOR FUEL FRACTIONS OF 0.0. 0.1. 0.2 - - - 1.0

EQUILIBRIUM GAS TEMPERATURE

:160		(150)	TEMP. 1 TEMP. 2 TEMP. 5 300.0000 (151) 3064.00000	(151)	1EMP. 3	(152)	TEMP. 4 (152) 3397.00000
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		2	FOUTLIBRIUM GAS MOL.	S MOL.	. #E16#1		
3	#0L. #T. 1 46.005000 (162)	(162)	MOL. WT. 2 28.793000 (163)	(163)	HOL. MT. 3	÷:	#0L. #7. 4 23.300000
(165)	HOL. HT. 5 14.880000 (104)	÷	MOL. WT. 6 18.750009 (167)	(167)	14.410000	(10)	#DL. #7. R
:	MOL. WT. 9 14.666666 (170)	(170)	40L. #T. 10 14.100000 (171)	(171)	HOL. WT. 11	(172)	8.9066.0
		ŭ	EGUILIBRIUM GAS GAMMA	18 GAH			
(173)	1.120000	(373)	SAMA 2 1.250000	(175)	GAMMA 3	(176)	6444 4. 1.217000
(177)	GAMMA 5 1.235000	(174)	6APHA 4 1.288000	(174)	1.3090n0	(140)	6444 A
3	GAMMA 4 1-270000 (182)	(182)	GAMMA 10 1.247000 (183)	(163)	54MMA 11	(184)	0.00000.0
			THRUST COEFFICIENT TABLE	TCIENT	TABLE		
		EOU	EQUILIBRIUM THRUST COFFFICIENT	1ST CO	FFICIENT		
(152)	CF VAC 1	(223)	CF VAC 2 1.802600 (223)	(223)	CF VAC 3	(\$54)	CF VAC 4 1.001700
(225)	CF VAC 5	(526)	CF VAC &	(727)	1.964000	(224)	C+ V4C 6 1.933100
(\$24)	CF VAC 9	(830)	CF VAC 10 1.022400	(231)	1.895900	EXP.	4HEA RATIO 40.000000

ADDUCT PROPERTIES

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FUEL (233)	VALVE DPEN 0.000000	0x10 (254)	VALVE DRF%	FJEL (235)	FUEL VALVE NPEN. CHIN VALVE NPEN. FUEL VALVE CLOSF CHID VALVE CLOSF (335) 0.0000000 (235) 0.017000 (235) 0.017000	0x10	VALVE CLUSE .017000
			SECOND PULSE TIMING	SF 71.	97.1		
FUEL (237)	VALVE DPF% 0.000000	0x10 (238)	** ***********************************	FUEL (239)	FUEL VALVE MPER. DWID VALVE MPER. FUEL VALVE CLOSK. MXID VALVE CLOSK (237) 0.000000 (238) 0.000000 (239) n.ncoon (240) 0.ncngng	0x10	VALVE CLUSF
			THIRD PULSE TIVING	SF 71.	9 1 1		
FUEL (241)	VALVE DPEN 0.00000	0x19 (5#5)	VALVE DPEN 0.000000	FUEL (243)	FUEL VALVE OPEN OXIO VALVE OPEN FUEL VALVE CLOSE OXIO VALVE CLOSE (241) 0.000000 (243) 0.0000000 (243) 0.0000000	7×10	0.000000 0.000000
			FRURTH PULSE TIMING	SE 71	11.6		
FUEL (245)	FUEL VALVE OPEN (245) 0.000000	0x10 (540)	VALVE OFFN 0.000000	FUFL (247)	DRIO VALVE OPEN FUEL VALVE CINSE TRIT VALVE CLUSE (244) 0.000000 (247) 0.000000 (244) 0.100000	7KT (24A)	VALVE CLOSF 6.130000
			FIFTH PULSE TIMING	.SE 71	1.6		
FUEL (249)	VALVE DPEN 0.000000	0x10 (250)	VALVE DREN	FUFL (251)	FUEL VALVE OPEN OXID VALVE OPEN FUEL VALVE CLOSE OXID VALVE CLOSE (244) 0.000000 (252) 0.000000 (253) 0.000000	0x10 (252)	VALVE CLUSE A.OCORDO
			SIXTH PULSE TIMING	SE 71	971,		
FUEL (253)	VAL -E OPEN 0.00000	0 × 10	VALVE OPEN	FUEL (255)	FUEL VALVE OPEN CYIO VALVE NPEN FUEL VALVE CLOSE OXIO VALVE CLOSE (2553) 0.000000 (254) 0.0000000 (255) n.nononn (256) n.goronn	0x10 (256)	VALVE CLP4E 0.00F071
			SEVENTH BULSE TIMING	SE 71	11.6		
FUEL (257)	VALVE DPEN 0.930000	OXID (PSP)	0-00000 0	F.,EL (259)	FUEL VALVE PPEN DYID VALVE PPEN FUEL VALVE CIGSE FXTO VALVE CIFSE (257) 0.1100000 (258) 0.1000000 (259) 7.100000	Cac	7-130003

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MULTI-RING INJECTOR

FIRST WING FUEL HOLES MOLE DIAMETER HOLE LENGTH ANIAL LOCATION RADIAL 'OCATION (201) 0.000000 (201) 0.000000 (201)

OF HOLES TRANSVERSE ANGLE

RADIAL INJ. ANGLE DISCHARGE COEFF. NUMBER (20%) 0.000000 (264) 0.00000 (267) MOLE DIAMETER HOLE LENGTH AXIAL LOCATION RADIAL LOCATION (200) 0.000000 (201) 0.000000 (202) 0.000000

OXIDIZER HOLES

RADIAL INJ. ANGLE DIBCHARGE COEFF. NUMBER OF HOLES TRANSVERSE ANGLE (20%) 0.000000 (20%) 0.000000

68

COMBUSTION CHAMBER PROFILE

IMJECTOR LOCATION INJECTOR DIAMETER AXIAL LOC. 2 CHAMBER DIAM. 2 (207) 0.00000 (200) .417350 (200) .250000 (300) .417350 (301) .250000 (300) .417350 (301) .250000 (302) .417350 (303) .750000 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .417350 (303) .

IMPUT UNITS ARE INCHES. PSIA. SECONDS AND DEGREES KELVIN. PROPELLANT PROPERTIES ARE IN GRANS/CC. POISE. DYNE/CM.

REDUCED TEMP.	TEMPERATURE	LIO. ENTHALPY	VAP. FATHALPY		TOTAL SECTION OF THE PROPERTY	ALBOORIA	BURFACE TENBION
.30000	178.200000	122.958000	145.004000	.000001		.200413	74.710000
.325000	193.050000	133.204500	354.784750	.000024	. ******	.127107	72.494632
.350000	207.900000	143,451,00	374.560510	.000321	. *****	.083897	500000
.375000	222,750000	221-107500	300,336250	.002316	. 952733	.056426	64.573200
******	237.60000	231.444000	404.112000	.011917	.938751	.039100	40.845285
.425000	252.450000	241.690500	418.847750	.046397	.924770	.027716	57.319306
. *50000	207.30000	251.937000	431.663500	.144632	.910108	.020032	53,430744
.475000	282.150100	262-183500	448.439250	**250**	.696806	.014735	50.44400
. 500000	297.00000	272.430000	261.215000	.97030~	. 662A2S	.011011	47.404344
.\$25000	311.85000	282.676500	477.000750	2.0A0199	. 868843	.018346	44.461380
.550000	326.70000	292.923000	492.700500	4-110665	.654861	80**00*	41.855208
.575000	341.550000	303.169500	507.542250	7.540A74	.64087	.00.07	30,163300
	354.400000	313.416000	522.318000	12.990598	.826898	.003400	36.641504
25000	371.250000	323.002500	537.003750	21.203041	.012916	.003433	34.225506
	366,130000	333.404000	551.444500	33.064106	.798934	.003034	31.930919
.475000	400.950000	344,155500	544.445250	49.557001	.784953	*00200*	20.753691
.70000	415,830000	354.492000	561.421000	71.754147	.770971	.002402	27.689749
.725000	430.450000	364.648500	500.196750	100.749887	.75.484	.002151	25.735095
.750000	445,500000	374.895000	010.072500	137.066477	.743007	.001433	23.885800
.775000	460.350000	385.141500	425.748250	184,149820	.729026	.001743	22.138009
. 800000	475,200000	3.5.366000	640.524000	240.036511	.715044	.001577	20.487937
.425000	400.050000	405.634500	655.200750	309.045044		.001432	16.443615
	504.401001	415.861000	470.075500	390.008007	.678221	.001303	16,150055
.875000	510.750000	426.127500	664.851250	484.656476	.654035	.001189	13.277417
. • • • • • •	534.600000	436,374000	499.427000	564.008405	. 624086	.001088	10.011953
. 425000	540.451000	446.620500	714.402750	718.050165	.564626	.0000	
. • \$0000	564.300000	456.867000	729.178500	860.315075	.538231	.000010	3.463114
. 15000	574.150000	467.113500	743.454250	1016.789321	.470167	*****	1.024387
1.000000	594.000000	477.360000	758.750090	1195.00000	.292811	.000340	. 000000

DERIVED GXID PROPERTIES	MOPL BTIES						
REDUCED TEMP.	TEMPERATURE	LID. ENTHALPY	VAP. ENTHALPY	VAPOR PRESSURE LIG. DENSITY	LIG. DENSITY	VISCOSITY	BURFACE TENSION
.30000	124.300000	46.548000	104.959400	. 000000	1.865309	.365550	76.696121
.325000	140.075000	50.427000	198.170350	000000	1.639178	.232006	72.473674
.35000	150.050000	54.300000	201.381300	.000001	1.812957	.152296	68.428405
.375000	101.625000	58.165000	204.592250	.000012	1.786736	.102920	64.554710
00000	172,400000	#2.0##n-0	207.403200	.000103	1.760515	.071334	60.847856
. 425000	163,175390	000840-54	211.014150	.000451	1.734294	.050553	57.302983
	193,950000	69.822000	214.225100	.003200.	1.709073	.036538	53.015301
.475000	204.725000	73,701900	217.436050	.012872	1.661652	.026877	50.660093
.500000	215.500000	77.580200	220.047003	.043714	1.655631	.020084	47.592715
.525000	225.275000	91.459100	223.857950	.129195	1.629410	.015223	44.648592
.550000	237.050300	A5.338.00	227.008900	.339845	1.603189	.011988	41.843223
.575000	247.625000	69.217000	230.279850	45000V.	1.574968	000000	39.172179
00000	258.690000	93.00000	233.490800	1.774303	1.550747	.007130	36.631102
.625000	264.375000	136-175000	236.701750	3.014636	1.524526	.000201	34.215705
0000\$**	260.150300	140.054000	230.012700	6.014437	1.498305	.005533	31.921776
75000	294.925700	145.933000	243.123650	12.519297	1.47.084	.004013	29.745171
.70000	301.730300	147.012000	246.334600	21.500537	1.445863	.004361	27.681920
.725000	312.475000	151.691300	249.545550	35.711441	1.419642	.003923	25.727720
.750000	323.250000	155.570000	252.756500	56.85276A	1.393421	.003525	23.878960
.775000	334,025980	159.449300	255.967450	A7.511463	1.367200	.003100	22.131669
.800000	344.500000	163.328500	259.178400	130.701590	1.340974	.002877	20.482070
.625000	355,575000	167.207000	262.369550	199.066723	1.308419	.002611	10.438334
.650000	366.350000	171.096000	205.600300	200.406397	1.271923	.002377	10.145430
.875000	377.125900	174.965000	264.811250	373.918685	1.226564	.002160	13.273615
000000	367.900000	178.844900	272.022200	508.479A68	1.170403	500100	10.00000
. \$25000	398.675000	142.723000	275.233150	678.929785	1.100147	.001921	6.617793
	409.450010	186.002000	279.444100	801.405836	1.009367	.001674	3.442122
. 475000	420,225000	190,481000	201-655050	1152.636831	.661741	.001543	1.02002
1.000000	431.000000	194.360000	244.856000	1470.000000	.549131	.000712	. 000000

DERIVED FEED SYS	DERIVED FREG SYSTEM VALUES - INJECTION, RESTRICTOR, AND PORT AREAS REFER TO VENA CONTRACTA	ECTION. MESTAICT	DR. AND PORT ARE	AS BEFER TO VENA	CONTRACTA		
FUEL STDE VALUES	VALUES						
LINE AREA 2.810005E-02	INJECTION AREA 1.960608E-04	RESTRICTOR AREA 2.197748E-04	VENTURE AREA 2.810005E-02	VALVE PORT AREA 2.810000E-02	LIME AREA INJECTION AREA PESTAICTOR AREA VENTURI AREA VALVE PORT AREA LINE RESISTANCE LENGTHYAREA 2.816005E-02 1.960668E-04 2.197748E-04 2.816005E-02 2.8160605E-02 4.88848	IJ. REBISTANCE 624932	LENGTH/AREA 17400.6
ONID SIDE VALUES	VAL UES						
LINE AREA 2.810005E-02	INJECTION AREA 2.717163E=04	RESTRICTOR SHFA	VENTUDI 4864 2.810005E-02	VALVE PORT AREA P.B10000E-02	LINE AREA INJECTION AREA RESTRICTOR ANFA VENTUDIAREA VALVE PORT AREA LINE RESISTANCE LENGTWAREA 2.010005E-02 2.717103E-04 3.442616-04 2.01005E-02 2.010005E-02 2.01024	1J. RESISTANCE 325379	LENGTH/AREA 17311.8
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THE CALCULATED AUN PARAMETERS ARE PRINTED FOR EACH INTERVAL OF TIME FOLLOWING START

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¥.	FUEL INJ BATE		0.0	ONIO INJ MATE	FUEL .	FUEL FLOW RATE	0 × 10	OXID FLOW RATE	FUEL VOID VOL	OXID VOID VOL	FUEL INJ TEMP	0110 I	OKIO INJ TEMP
3.	FUEL DROP BURN		OXID.	OXID D409 BURN	FUEL	FUEL FLASH RATE	0	OFID FLASH RATE	FUEL MALL BURN	OKID MALL BURN	FUEL WALL EVAP		OKID MALL EVAP
ğ:	FUEL GAS MASS		OI N	OKID 648 MASS 6-417895E-13	FUEL 0.	FUEL DROPLETS	Oxto	OKID DROPLETS	FUEL STREAMS	OXID STREAMS	FUEL ON WALL	ONID ON MALL	174.
, if	FUEL GAS EFFLUX 0.		0:	OKID GAS EFFLUX	FUEL 0.	FUEL DROP RATE	0 x 10	OKID DROP MATE	FUEL FILM RATE	OKID FILM RATE	648 FRACTION 1.00000	VAC THRUST LBF	107 11
ğ.	FUEL DROP 052		o .	0x10 000 032	FUEL 0	FUEL INJ. VEL.	0110	OKID INJ. VEL.	MAK. EXPLOSION	MASS OUT OF TANK INTEGRAL PEODT TOTAL EMPULSE 0.	M INTEGRAL PC+D	T 101AL	30 Indus
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PE.	FUEL DROP BURN		0.0	0 10 0409 BURN	FUEL 0.	FLASH DATE	0x15	FLASH RATE	FUEL FLASH DATE OXID FLASH RATE FUEL WALL BURN	OXID WALL BURN	FUEL WALL EVAP		OXID WALL EVAP
4.	FUEL GAS MASS		OIX.	OXID GAS MASS 6-417805E-13	FUEL	FUEL DROPLETS	0×10	OKID DROPLETS	FUEL STREAMS	OKID STREAMS	FUEL ON HALL	OXID ON MALL	174.
¥ .	FUEL GAS EFFLUI	_	9:	ONID GAS EFFLUX	FUEL 0.	FUEL DROP RATE	OKID 0.	OKID DROP MATE	FUEL FILM RATE	OXID FILM MATE	649 FRACTION 1.00000	VAC THRUST LBF	5
ë.	FUEL DROP 032		01 × 10	0x10 540# 032	FUEL 0.	FUEL INJ. VEL.	0x19	DXID INJ. VEL.	1.020408E-06	HASS OUT OF TANK INTEGRAL PCODT 7.447032E-08 1.00000E-10	1.000000E-10		TOTAL EMPULSE

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PUEL HALL EVAP OXID WALL EVAP OXID DROP RATE FUEL FILM RATE OXID FILM RATE GAS FRACTION VAC TWOUST LIBER 02. FUEL GAS EFFLUX OXID GAS EFFLUX FUEL DROP RATE OXID DROP RATE FUEL FILM RATE GXID FILM RATE GAS FRACTION VAC THRUST LBF 0. 1.378449E-05 0. 1.378449E-05 0. FUEL WALL EVAP OXID MALL EVAP HASS OUT OF TANK INTEGRAL PC-OFT TOTAL IMPULSE 6.243966E-05 1.218017E-06 6. 1074 1mPULBE FUEL WAL! EVAP OXID WALL EVAP FUEL INJ. VEL. DXID INJ. VEL. HAK, EXPLOSION HASS DUT OF TANK INTEGRAL PCOOT TOTAL IMPULSE O. 7.510626E-05 2.18713 OXID INJ TEMP OXID INJ 7840 OXID INJ TEMP 204,000 OXID ON HALL DXID ON MALL OXID ON HALL OXID FILM ALTE GAS FRACTION VAC THRUST LOP 1.91820 1,91829 1.01624 DAID INJ. VEL. MAK. EXPLOSION MASS OUT OF TANK INTEGRAL PCODT 70.2416 A.24556 6.763772E-05 2.135130E-04 FUEL INJ TEMP 204,000 FUEL INJ TEMP 204.000 PUEL 19.3 TEMP FUEL ON MALL FUEL ON MALL FUEL ON MALL 54mma 1.16664 1.16044 1.16944 1.00000 DXID .ALL BURN DKID MALL BURN OXID MALL BURN OXIO AOID AOL DAID VOID VOL OXID VOID VOL OXID STAEAMS OKID STREAMS OKID STREAMS CHAMBER TEMP K MOL MT. 203,334 46,0080 MOL WT. CHANDER TENP K HOL MT. • FUEL FLASH RATE OXID FLASH RATE FUEL HALL BURN CHAMBER TEWP K 224.880 FUEL FLASH RATE OXID FLASH RATE FUEL WALL BURN. 0. 1.5139596-04 0. FUEL FILM MATE FUEL PLASH RATE DXID FLASH RATE FUEL WALL BURN 0. FUEL VOID VOL 1.304507E-04 FUEL INJ. FVEL. DXID INJ. VEL. MAX. EXPLOSION 0. 1.42452 3.407651E-02 FUEL FLOW RATE OXID FLOW RATE FUEL VOID VOL 1.233460E+02 1.352402E+02 9.00039E+05 OXID FLOW RATE FUEL VOID VOL 1.347123E-02 6.098774E-05 FUEL STREAMS FUEL STREAMS FIJEL STREAMS 0x10 FLOW RATE 1.337025E-02 ONID DROPLETS DKID DROP MATE OKID BROPLETS 1.511391E-06 FUEL FRACTION 6.727304E-03 FUEL FRACTION FUEL FRACTION FUEL FLOW RATE 1.245061E-02 FUEL GAS ETFLUX OXIO DAS EFFLUX FUEL DROP SATE 0. 7.246/516-04 0. PUEL GAS EFFLUX DXID GAS EFFLUX FUEL DROP MATE FUEL FLOW RATE 1-221308E-02 FUEL INJ. VEL. FUEL DROPLETS FUEL DROPLETS FUEL DROPLETS IGNITION IONITION IGHTTON . DXID D40P BURN CHAMBER FRESS 1.213817E-02 OXID 648 MASS 1.514001E-08 OXID DROP BURN OXID INJ RATE 1.367123E-02 UNID DROP BURN 0xID 648 MASS 1.158647E-06 OXID INJ RATE 2-4576-64 DKID INJ RATE 1.352402E-02 OXID DROP D32 6.864919E-07 CHAMBER PRESS. 0x10 080F 032 0x10 DROP 032 CHAMBER PRESS 1.30116 SEAM SAD CING FUEL BRUT SURN PUEL DROP BURN VEL DROP BURN TIME MILLIBEC 4.40000 TIME MILLIBEC 4.50000 PUEL BAS MASS TIME MILLIBEC 4.30000 PUEL INJ RATE FUEL GAS MASS FUEL DROP 032 PUEL INJ RATE FUEL GAS MASS FUEL DROP 032 PUEL INJ RATE FUEL DROP 032

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CP VAC	0X10 1HJ TENP	OXID WALL EVAP	OXID ON MALL	VAC THRUST LBF	707AL 1MPULBE 2.52[573E-85		1.91020	OXID INJ TEND 204.000	DATO WALL TVAF	OXID ON WALL	VAC THRUST LBF	3.723802E-09	CF VAC 1.01020	204.000 204.000	5.296286E-64	OXID ON MALL 3.645010E-87	VAC THRUST LBF .126561	TOTAL IMPULSE S.BEGGOTE-DS	CF VAC 1.91820
GAMMA I-16488	FUEL 195 TEMP 204.600	FUEL WALL EVAP	FUEL ON MALL	648 FRACTION V	MASS OUT OF TANK INTEGRAL PC-OT 7.027402E-05 78.073502E-04		64HH1 1-10066	FUEL INS TEMP 294.000	FUEL WALL EVAP	FUEL ON MALL	645 FRACTION V	MASS OUT OF TANK INTEGRAL PCODT 7.293591E-05 6.526217E-04	6 A L R A B A B A B A B A B A B A B A B A B A	FUEL 143 TEMP 294.080	FUEL MALL EVAP	FUEL ON MALL	645 FRACTION V	1.554187E-05 8.775473E-04	1.16643
**************************************	OXID VOID VOL	OXID MALL BURN	OKID STREAMS	DXID FILM RATE	1.027442E-05		#5_ #¥.	OKID VOID VOL	OXID WALL BURN	OXID STREAMS	CYIB FILM RATE	MASS OUT OF TAN 7.203501E-05	**************************************	OXID VOID VOL	OXID WALL BURN 0.	OXID STREAMS 1.407678E-06	OXID FILM MATE	T.SS4187E-05	#0 # # # # # # # # # # # # # # # # # #
CHAMBER TEMP K 235,722	FUEL VOID VOL 2.173379E=05	PUEL WALL BURN	FUEL STREAMS	FUEL FILM RATE	HAK. EXPLOSION 6.23707		C14MBER 72MP K 242.950	FUEL VOID VOL	FUEL MALL BURK 0.	FUEL STREAMS 5.708549E-07	FUEL FILM BATE	MAX. EXPLOSION 101.048	C * 4 * 4 * 4 * 4 * 4 * 4 * 4 * 4 * 4 *	FUEL VOID VOL	FUEL MALL BURN	FUEL STREAMS	FUEL FILM RATE	*** EXPLOSION 243-167	CLANDER TETO K 245.456
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71ME MILLIBEC	PUEL INJ RATE	FUEL DAGE BURN	FUEL 648 MASS	FUEL GAS EFFLUX OXID	FUEL DROP 032	UEL MANIFOLD FILLS	TIME MILLIBEC 4.70000	FUEL INJ RATE 5.708540E-03	FUEL 040P BURN 0.	FUEL 618 1188	FUEL GAS RFFLUX DXIC	FUEL DROF 032	TIME MILLIBEC	FUEL INJ RATE 1-238290E-02	FUEL DROP BURN	FUEL GAS MASS	FUEL GAS EFFLUX DXID	FLEL DROP 032	714E MILL18EC

	ENP OXID INJ TEMP	EVAP OXID WALL EVAP	LL 0XID ON WALL 6-21	DN VAC THRUST LOF	PC-DT TOTAL IMPULBE		CF VAC 1.91649	EMP CKID INJ TEMP	EVAP OXID WALL EVAP	E-07 0xID ON WALL E-07 0.250113E-07	DN VAC THRUST LBF 5.26911	PC=07 T07AL IMPULBE E=02 5.000355E=04	1 00011057 00010357 00001055 00001000	.	CF VAC 1.9466	EMP OKID INJ TEMP 294.000	EVAP OXID HALL EVAP	
	FUEL INJ TEMP 294,000	N FUEL WALL I	FUEL ON WALL	1.00000	INK INTEGRAL PC+DT		GANNA 1.22410	FUEL INS TE	FUEL WALL EV	FUEL ON HALL 2.283419E-07	1.00000	NK INTEGRAL !	DROPS RETAINED .00181887 .00280513 .00421999	707AL MOMENTUM 6.168038E+80 0. 5.913033E+80	64HA 1.21795	FUEL ING TEMP 204,000	FUEL WALL	
	DAID VOID VOL	OKID HALL BUR	OXID STREAMS 1.420234E-06	OKID FILM MATE	MASS BUT OF TAN 7.821542E-05		HOL WT.	CKID VOID VOL	OXID HALL BURN 0.	DXID STREAMS 1.432398E=06	OXID FILM RATE	MASS OUT OF TANK INTERRAL PCODT 8.083876E-05 1.631407E-02	648 RETAINED .00030272 .00078421 .00115093	A. 121340E-03 A. 121340E-03 D. A. A. 030E-03	HOL WT.	OKID ADID AD	CKID HALL BURN 4.4634836-04	
	FUEL VOID VOL	FUEL WALL BURN	FUEL STREAMS	FUEL FILM MATE	MAK. EXPLOSION 362.887		C1448ER TERP 4 3264.44	FUEL VOID VOL	FUEL WALL BURN 0.	FUEL STREAMS	FUEL FILM BATE	MAK. EMPLOSION ADD. 430	#ALL EXPELLED 0.0000000 0.0000000	00. MAY AND 01 00. MAY AND 01 00. WAY AND 01 00. WAY 00	CLANDER TEXP X NACTORS	FUEL VOID VOL	FUEL #ALL BURN 2.112813E-05	G.161 6485AM
•	OKID PLOM RATE 1.4202346-02	OKID PLASH MATE	OKI' DROPLETS 3.686063E-06	OKID DRCP MATE	OKID INJ. VEL. A2.3201		FUE. FRACTION . 339451	OKID FLOW MATE	CKID FLASM RATE	OKID DADPLETS	OKID DROP RATE	0xID 1%J. VEL. 83.0251	0000 0000 0000 0000 0000 0000 0000 0000 0000	SUM D 3.481708402 0. 0. 0. 0.	FUEL FRACTION	OXID FLOW RATE	DXID FLASH RATE	DKID DROPLETS
	FUEL FLOW RATE 1.213311E-02	FUEL FLASH RATE	FUEL DROPLETS 3.022455E-06	FUEL DROP MATE	FUEL 193. VFL.		100000	FUEL FLON MATE 1.190937E-02	FUEL FLASH MATE	FUEL DROPLETS 3.119245E-06	FUEL DROP RATE	FUEL INJ. VEL. 158.199	6A8 EXPELLED 0.0000000 .00044765	NUMBER OF THE STATE OF THE STAT	16%1710%	FUEL FLOW RATE 1.150819E-02	FUEL FLASH RATE	FUEL DROPLETS
	0x10 143 4ATE 1+420234E=02	OXIO DROP BURN	OXID 645 HASS 1-618072E-06	OXID 645 EFFLUX 2-010654E-03	0x10 040P 032	MILLISECONDS	CHAMBER PRESS 92.2611	OKID INJ RATE 1-432390E-02	DKID DHOP BURN 1-401036E-03	OXID GAS MASS 1.684797E-06	OXID 645 EFFLUX 1-876262E-03	33.0550	SECTED TABLE OF SECTION OF SECTIO	18.56.760 EJECTED 18.56.760 EJECTED	CEATORE PERSON	DXID INJ RATE 1.424640E-02	0xID DROP BURN 1-788193E-02	OXID 648 MASS
	FUEL INJ RATE 1.2133116-02	FUEL DROP BURN	FUEL GAS MASS	FUEL GAS EFFLUX 0.	FUEL DROP D32	ISMITE AT 5.00 m	TIME MILLISEC	FUEL 145 RATE 1-1909376-02	FUEL DROP BURN 8-458055E-03	FUEL GAS MASS 8.458055E-07	FUEL GAS EFFLUX 0.	FUEL DROP 032	FUEL GRANS OXIO GRANS TOTAL GRANS	######################################	TIME MILLIBEC 5.10000	FUEL INJ RATE 1-150819E-02	FUEL DROP BURN 5,75573E-03	PUEL GAS MASS

	OXID INJ. VEL. MAX. EXPLOSION MASS DUT OF TINK INTERNAL PEODT TOTAL IMPULSE 82.5806 503.949 8.3413612-05 2.1450062-02 1.8383726-03	FUEL FRACTION CHAMBER TEMP K HOL HT. GAMMA CP VAT 25.0725 1.21728 1.05640	DAID FLOW RATE FUEL VOID VC. DAID VOID VOL FUEL INJ TEMP DAID INJ TEMP 1.411686-02 0. 294.000	DXID FLASH RATE FUEL WALL BURN DXID WALL BURN FUEL MALL EVAP DXID WALL EVAP 0. 2.287712E-05 1.229740E-03 0. 0.	OXIO DROPLETS FUEL STREAMS OXIO STREAMS FUEL ON MALL OXIO ON MALL 3.773745E-06 0. 1.04013E-06 2.46736E07 1.04013E-06	OXID DROP RATE FUEL FILM RATE DXID FILM RATE GAS PRACTION VAC THRUST LEFF 6.440013E-04 0. 7.10033	OXID INJ. VEL. MAX. EXPLOSION MASS OUT OF TANK INTEGRAL PCODT TOTAL IMPULAE 81.8247 524.787 8.593540E-05 5.373749E-02 1.454409E-03	FUEL FRACTION CHAMBER TEMP K MOL NT. GAMMA CF VAC 1.224819 3251.11 21.8660 1.22482 1.91199	OXID FLOW BATE FUEL VOID VOL OXID VOL FUEL INJ TEMP OXID INJ TEMP 1.3972265-02 0. 294.880	OXIO'FLASH RATE FUEL WALL BURN OXID WALL BURN FUEL WALL EVAP OXIO WALL EVAP 0. 3.154402E-05 1.644519E-03 0. 0.	OXID DROPLETS FUEL STREAMS DXID STREAMS FUEL ON WALL DXID ON WALL 4.112740E=06 0.	OXID DROP RATE FUEL FILM RATE DXID FILM RATE GAS PRACTION VAC THRUST LDF 0.552996E-04 0.	FEL. MAX. EXPLOSION MASS OUT OF TANK INTEGRAL PC-051 T 540.994 8.840.951E-05 8.594151E-02	FUEL FRACTION CHAMBER TEMP K MOL NT. GAMMA CF VAC .364346 3180.79 21.1514 1.22858 1.86789	OXID FLOW RATE FUEL VOID VOL DXID VOID VOL FUEL EMJ TEMP OXID INJ TEMP 1.363376E-02 0. 294.000	OXID FLASH RATE FUEL WALL BURN DXID WALL BURN FUEL WALL EVAP OXID WALL EVAP 0. 5.364662E-05 1.73222E-05 0.	OXID DROPLETY FUEL STREAMS DXID STREAMS PUEL ON MALL DXID ON MALL 4.340170E-06 0. 1.383376E-06 2.422687E-07 7.076582E-07	ID DADP RATE FUEL FILM RATE OXID FILM RATE GAS FRACTION VAC THRUST LBF 4.972621E-04 0. 7.07616	
	OKID OROP 052 FU 52.6442	CHAMBER PRESS IS	OXID INJ RATE PU	1.362439E-02	DYID 645 MASS FU 2.510965E-06	OKID GAN EFFLUX	0X10 0R0P 032 FU	CHAMBER PRESS IG	OXID INJ RATE FU 1.397226E-02	0x10 040F BURN FU 1.063979E-02	OXID 645 MASS FU 2.213720E=04	OXID GAS EFFLUX F 1.525498E=02	0x10 DRDP 032 FU	CHAMBER PRESS 16 125.661	OXID ING RATE FU 1-SBISTAE-02	OXIO DROP BURN FU 1-167482E+02	OXID 648 MASS FU 2-176204E-06	0xID 648 EFFLUX 1.370135E-02	
	FUEL DROP D32 64.2436	THE HILLIBEC S. 2000	FUEL INJ BATE 1-110391E-02	10-361408-6 10-361408-6	FUEL 648 HA38 1-028627E-06	FUEL 6AS EFFLUX 9.367169E-03	FUEL DROP 032 67,7231	TARE ARC. S. MODO	FUEL INJ #ATE	FUEL DROP SURV	FUEL 544 MASS 1.157867E-06	FUEL GAS EFFLUX 6.251251E-03	FUEL DROP D32	TIME MILLIBEC 5.4000	FUEL ING BATE 1.035860E=02	FUEL DROP BURN 8.870173E-03	FUEL 648 MASS	FUEL GAS EFFLUX 7.200210E-03	

P. 300	. 077034		4. 052057E-40	E	18.00	1.10960	- Marie
- N W	0119 INJ RATE	FUEL FLOW RATE	OKID PLOW BATE	FUEL VOTO VOL.	1. 101010 vo.	PART PARTON	W. P. Links
PUEL DOOP BURN	OXIO DAGS BURN	FUEL FLASH BATE	OXIO PLASH BATE	FUEL MALL BURN	P. CASSAGE-OV	rue, "ath tras	0119 Hills 6745
FVEL 646 MASS	0170 640 4400 0.0521036-07	FUEL DROPLETS	0x10 0#0PLE78	FUEL BTREAMS	OK10 878EAMS	1765 OH 1861	OXID ON BALL
FUEL 648 EFFLUX 6-036679E-12	1.003709E-04	FUEL DROP RATE 6.666612E-17	3.390313E-06	FUEL FILM MATE	OKID FILE BATE	GAS PRACTION V	VAC THRUST LAF
FUEL DADP 052	OKID DAGE 032	FUEL INJ. VEL.	0110 14J. VEL.	HAY. EXPLOSION 15.6254	3.53600E-64	I.PSSS	TOTAL STREET
71ME MILLIBEC 29.9888	CHAMBER PRESS.	Jewition 0.	FUEL PRACTION 6.184988E-00	C1448FF 1F1F K 256.644	#OL #7.	GANNA 1.1994	CF VAC 1.01620
PUEL INJ RATE	ONID INJ RATE	FUEL FLOM MATE	OXID FLOW RATE	FUEL VOID VOL.	0x 10 v010 v0L 5.600006-04	PUEL INJ 7EMP 204.055	001.00 INJ 7EM
FUEL DROP BURN 9.	OXIO DROF BURN	FUEL FLASH BATE	OXID FLASH RATE	FUEL WALL BUPN	OXID HALL BURN 2,616669E-87	FUEL WALL EVAP	0KID HALL EVAP 7.6679726-04
FUEL 648 MASS 3.953560E-15	OKID 648 MASS 6.475950E-07	PUEL DROPLETS	0x10 D40PLETS	FUEL STREAMS	OXID STREAMS	FUEL ON MALL	0110 OH BALL
FUEL GAS EFFLUX OXID 5.320351E-12 7.4	0x10 6x8 EFFLUX 7.652916E-04	FUEL DROP RATE 5.165189E-17	OKID DROP RATE 2.542734E-06	FUEL FILM RATE	OXIO FILM RATE	eas PRACTION V	VAC THRUBT LBF 5.0395606-02
FUEL DROF 032	0x15 5400 552	FUEL INJ. VEL.	OKID INJ. VEL.	MAX. EXPLOSION 15.7819	NASS DUT DF TAMK INTEGRAL PCODT 3.336063E-06 1.25367	INTEGRAL PC-DT	7.10046-02
TIME MILLIBEC 30.000	CHAMBER PRESS.	IGNITION 0.	FUEL FRACTION 5.339663E-89	CHAMBER TEMP K 256,595	100 100	\$5001-1	1,010P0
PUEL INJ RATE	OXID INJ RATE	FUEL FLOW RATE	OXIO PLOM BATE	FUEL VOID VOL	OXID V010 VOL 5.800000E-04	FUEL THJ TEM-	0118 INJ 7210 204,000
	ONID DADP BURN	FUEL FLASH RATE	OXID FLASH RATE	FUEL WALL BURN	OXID WALL BURN 2.602656E-87	PUEL WALL BOAP	0x10 HALL EVAP
FUEL 648 HASS 3.666602E-15	OXID 648 HASS 6.526125E-07	FUEL DROPLETS	0x10 DROPLETS	FUEL STREAMS	OXID STREAMS	בחבר פא אקרי	0XIO DN HALL
### 648 EFFLUX 0X10 4.6486 6.12	7.60001E-04	FUEL DROP RATE 3.873832E-17	0x10 040P RATE 1.073045E-05	FUEL FILM RATE	OXID FILH BAPE	648 PHACTION 9	VAE THRUST CBF
FUEL DROP 052	0x10 0809 052 154.677	FUEL INJ. VEL.	OXID INJ. VEL.	MAX. EXPLOSION 15.5742	3.33669E-04	INTERRAL PC-DT	707AL 2MPULBE 7.1000006-02
PUEL GRAND ON 20 DEANS TOTAL DEANS	16,750 1688 06401422 00000071	6AB EXPELLED .0324960 .0640039	DROPS EXPELED .01175073 .01553774	**************************************	648 RETAINED	0.000000000000000000000000000000000000	#ALL RETAINED 0.00000000

1011 MOMENTUM 0.7007776-01 1.227816002 1.224486002 9.0000746-01
2.534634E-62 2.534634E-62 1.13887E-61 2.62626E-62
2.0724686+01 2.0724686+01 5.2524726+00 7.0044726+01 5.4858576+00
8-110820E+03 9-710820E+03 5-62282E+03 1-650740E+63
4.951507E+06 1.407994E+05 1.218997E+07 2.89591E+05
FUEL DROPS INJECTED PUEL DROPS EXECTED ONIO DROPS INJECTED OXIO DROPS EXECTED

PERPORANCE SURFARY					
ROPELLANT HASS BASES	FUEL FLOW	OKID FLOW	OKID FLOW HINTURE RATIO C STAR	C 8748	VACUUM 18P
ASS OUT OF TANK - POUNDS	.000135	.0001	1.467000	3003.320247	215.621406
ABB THROUGH INJECTOR - POUNDS	••0000•	.0001	2.003970	4041.357435	242.057205
ABS THROUGH NOIZLE - POUNDS	••0000	.0001	1.05044	4170.076803	250.20052
RESSURE INTEGRAL - PSIA-SEC.	1.2538				
OTAL VAC IMPULBE - POUND*SEC.	.0720				

		018608	1	DISPOSITION OF INJECTED PROPELLANT	PROPELLANT		
					FUEL	OKID	TOTAL
	PRACTION	EXPELLED	3	PRACTION EXPELLED AS BURNT GAS	.733766	.707953	.716546
	PRACTION	EXPELLED	7	PRACTION EXPELLED AS UNBURNT VAPOR . DODGES	.00000	.00000	.045827
	PRACTION	PRACTION EXPELLED AS DACPS	3	94049	.201020	.172620	.202285
	PRACTION	PRACTION EXPELLED AS PTER	7	PILE	.00400.	.001350	.002440
	PRACTION	RETAINED	7	PRACTION RETAINED AS BAS-VAPOR	00000	.003200	.002194
	PRACTION	PRACTION RETAINED AS DROPS	2	-	0.00000	.000214	.000143
İ	PRACTION	PRACTION REFITHED AS FILM	7		0.00000.0	-045842	.030595
	TOTAL OF	TOTAL OF ABOVE VALUES	20		1.00000	1.00000	1.00000

TOTALS ORESTER THAN 1.0 MAY INDICATE --

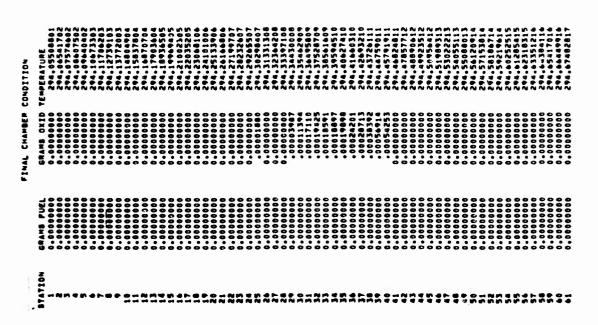
DEPLETION OF CHAMBER GAS MASS - THERE MAY BE AN APPRECIABLE MASS OF COLD VAPOR IN CHAMBER BEFORE START. ISMITER FLOWS - NOT CLASSED AS THROUGH-INJECTOR OR DUT-OF-TANK.

DEPLETION OF MALL-FILM. MHEN A FIRING IS STARTED WITH PROPELLANT-COATED MALLS.

INJECTED OXID EJECTED OXID 41.2082 46.0354 58.1664 200.32 32.22 21.0576 26.2572 36.7903 45.55 INJECTED FUEL EJECTED FUEL 56.4597 15.4097 77.0110 370.56 36.98 26.9441 33,3454 46.7350 71.52 HEAN RADIAL VELOCITY FT/SEC HEAN AXIAL VELOCITY FT/SEC HEAN DROP DIAMETER DIO HICRONS DS1 HICRONS 632 HICRONS

NOTE . THE ABOVE EJECTED DROPLET AND EJECTED FILM VALUES ARE AT THE THROAT PLANE. NOT THE EXIT PLANE.

HICRONS	100 0	BUND SIZE		- CUMULATIVE	\$87#
0.	_		0.000		.000.
20	1.3010	1486	000		-005
30	1.4771	2001	070	1001	1881
	1.6021	2000	0726	516	
20	000	0002	1001	408	101
0	1.7782	4414	7516	7777	****
70	1.045		28.0	7.5	
		144		7454	
	•	404	1 1 5		
	2.0000	5402			4367
		2482			****
	•				
	•				
	: -	5492			
150		1477	;		
	2.2041	7			
170	2.2304	1977			
100	2.2553	5	2800.	978	0750
0 - 1	2.2788	5400°	- 00 B	176	:
200	٦.	•	2000.		
210	2.3222		- 0045		
220	7.		2400°	. 0785	
200				527.	
250	2.5070			- 18	
240		:	:	. 0785	
270		-	0000	185	****
280	2.4472	5	0000	•	3
540	•	0500	0	•	.000
200	•	5	000	•	.000
9.0	•	9000	•	1.0000	.000
0 0 0	2.5051	: :			00
000	•	-			
150	•			•	000
200				•	
370		:		•	000
300	2.570	Ī		0000	000
300		9000	0000	1.0000	000
00	•	:	0000	1.0000	.000
410	•		0000	1.0000	.000
420	•	œ.	:	1.0000	6
430	2.0135	***	•	1.0000	1.0000
•	•	ĕ	.000	1,0000	.000
450	2.4532	:	8	1.0000	000
940	1	1.0000	.000	00000	4



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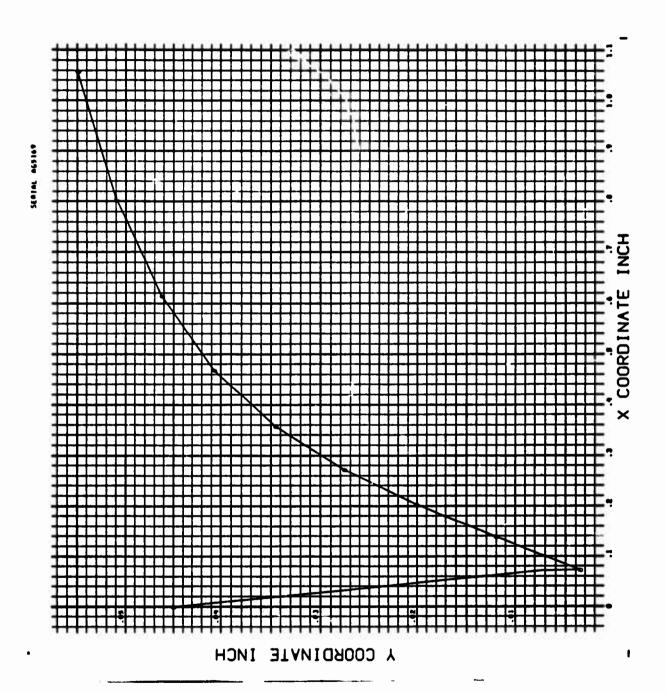
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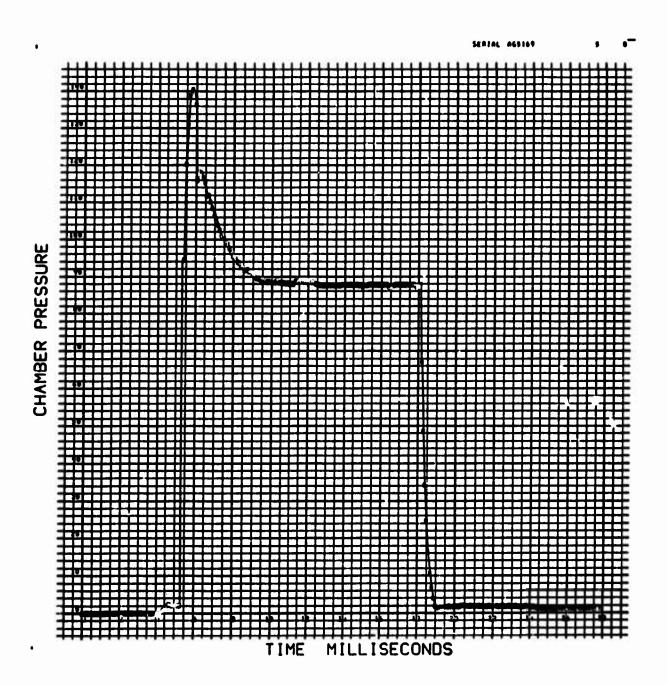
	THE-AVERAGED	INTERVAL PER	TIME-AVERAGED INTERVAL PERFORMANCE VALUES		
START TIME	.01500	.015000 BECONDS			
FINISH TIME	.017000	.017000 SECONDS			
INTERVAL LENGTH	.002000	.002000 BECONDS			
TOTAL IMPULDE	.010047	.016047 POUND+SEC	THRUBT	5.048442 POUNDS	POUNDS
CHANGER PRESSURE INTEGRAL	.175398	.175398 PSIA+BEC	CHAMBER PRESSURE	67.698917 PSIA	P814
FUEL OUT OF TANK	.000014 POUNDS	BONNOS	FUEL FLOW RATE	.00001	.004979 POUNDS/SEC
OXID OUT OF TANK	.000024 POUNDS	\$0N004	OXID FLOW RATE	.011994	.011944 PDUNDS/SEC
FUEL-OXID DUT OF TANK	.0000036 POUNDS	800004	TOTAL FLOW RATE	.010073	.018973 POUNDS/SEC
MINTURE RATIO	1.718620	1.718628 OXID/FUEL			
C-87AR	4431.8580&0 FEET/SEC	FEET/8EC			
VAC. SPECIFIC IMPULSE	266.095743 BECONDS	SECONDS			

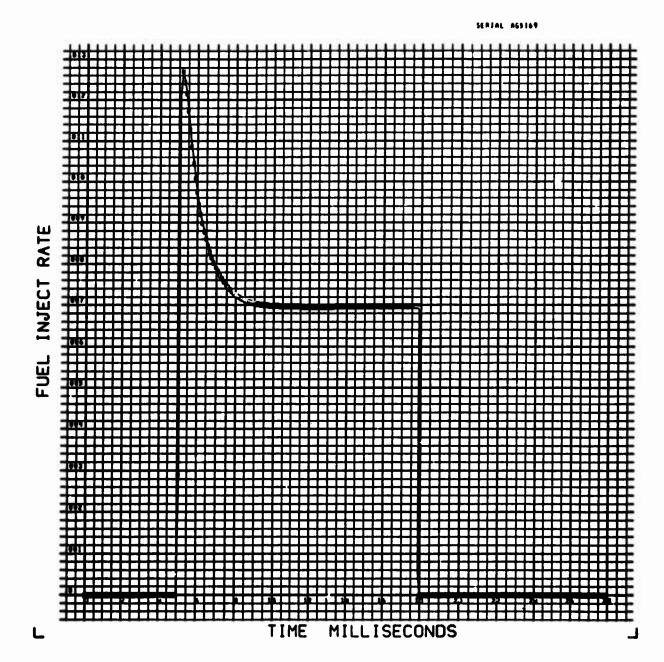
	PWOPELLANT	4		
	FUEL POUNDS	FUEL POUNDS/SEC	OXID POUNDS	OXID POUNDS/SEC
PRODUCTS OF COMBUSTION	.00001020	•00200•	.00001	.000072
UNBURNED VAPOR	0.00000000	000000	0.00000000	0.00000
UNBURNED DROPLETS	.00000372	.00185	.00000404	.002021
HALL FILM	0.000000.0	0.000000	0.00000000	0.00000

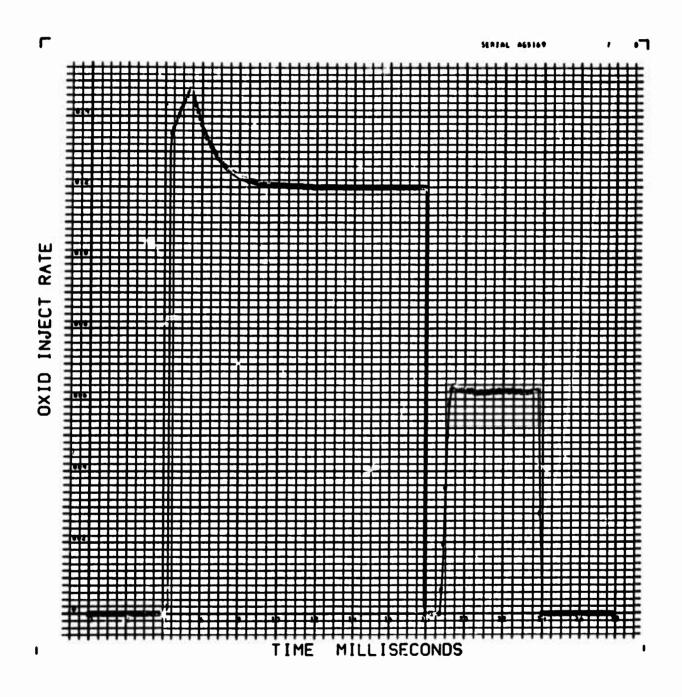
TELER AND VELOCATIVE	OXID DROPLETS	40.057004	000000000000000000000000000000000000000	-098103800	232.088485	43.342764
EXPERIEN UNCFER MEAN DIAMETER AND VELOCATE	FUEL DROPLETS	56.615033	65.639041	76.266954	391.921920	30,373362
		DSO MICHONS	D31 HICRONS	D32 HICRONS	AXIAL VELOCITY FT/SEC	RADIAL VELOCITY FT/SEC

	TUGNI	INPUT VALUES FOR HULTRAN		
CHAMBER PRESSURE	67.690017 PEIA	P814		
CHANDER TEMPERATURE	5001.745005 DEG.R.	DEG.R.		
SAB CONSTANT	2255.530752	2259.530752 FT++2/BEC++2 DEG.R		
GAS SPECIFIC HEAT	12331.005403	12331.985463 FT0+2/8EC++2 DES.A		
6A8 6AMA	1.224013			
SAS VISCOSITY	•60000	.000096 PDUNDS/FT.SEC.		
1	FUEL DROPLETS		OKID DROPLETS	18
DROPLET DENSITY	55.264500	55.264500 POUNDS/FT++3	11.3011	91.391105 POUNDB/FTB0
HASS DROPS/HASS 648	.123344		.134123	2
	EXPELLED	EXPELLED DROPLET SIZE DISTRIBUTION	20	
	FUEL N	FUEL MASS FRACTION	OXID MASS FRACTION	FRACTION
DIAM MICRON PADIUS PEET	ET INCREMENTAL	CUMULATIVE	INCREMENTAL	CUMULATIVE
10 1.6406-05	0.00000	0.00000	0.00000	0.0000
20 3.2016-05		0.00000	0.0000	0.0000
30 4,0216-05		.063294	.109228	.109220
46 6.5626-05	9.00000	.063204	0.00000	.109220
50 8.202E-05	.118126	.201419	.245770	.355007
\$0-2570.0	3 .021192	.222011	0.000000	.355007
70 1,1466-04	0.00000	.222611	000000-4	.355007
80 1.312E-04	. 286755	.50036	00000000	.355007
90 1.476E-64	0.00000	.5003.	00000000	.355007
100 1.6406-04	000000-	.5003.	0.00000	.355007
110 1.8042-04	0.00000	.5003.	0.00000	.355007
120 1,969E-04	0.00000	.50034	.644903	1.000000
130 2.1336-04	0.00000	.50934	0.000000	1.00000
140 2.2076-04	0.00000	.500366	0.00000	1.000000
150 2.461E-04	0-00000	.509366	0.00000	1.00000
160 2.625E-04	0.000000	.50936	0.00000	1.00000
170 2.789E-04	***************************************	1.00000	0.00000.0	1.000000

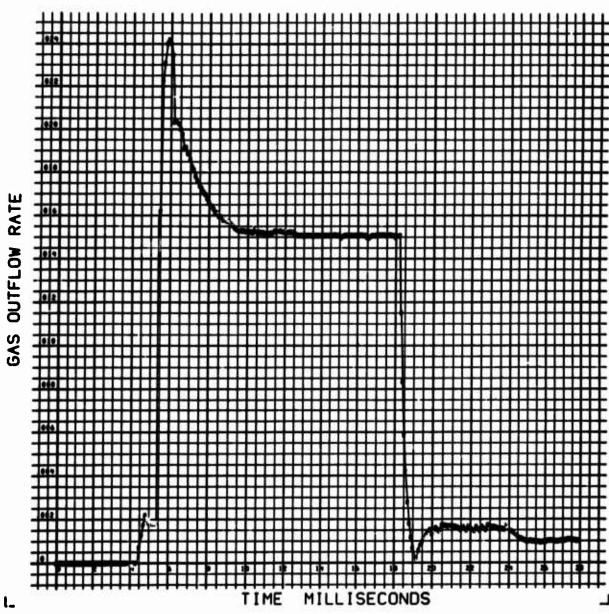


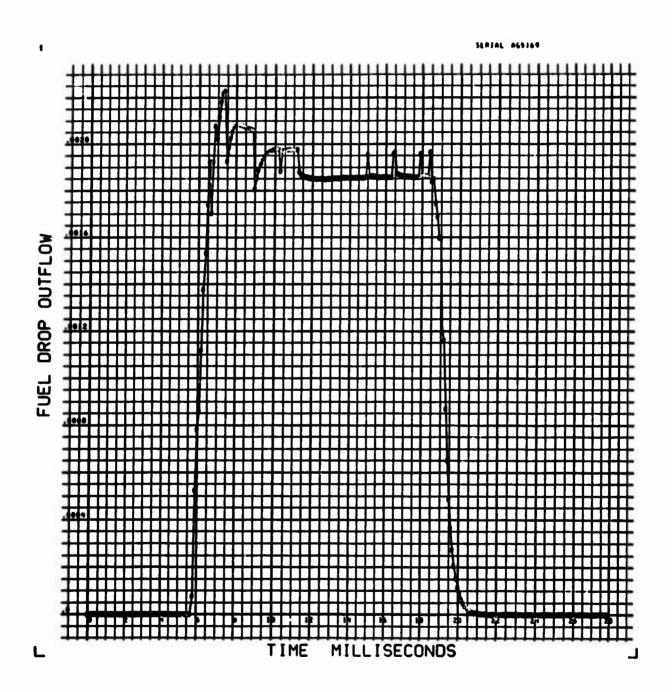


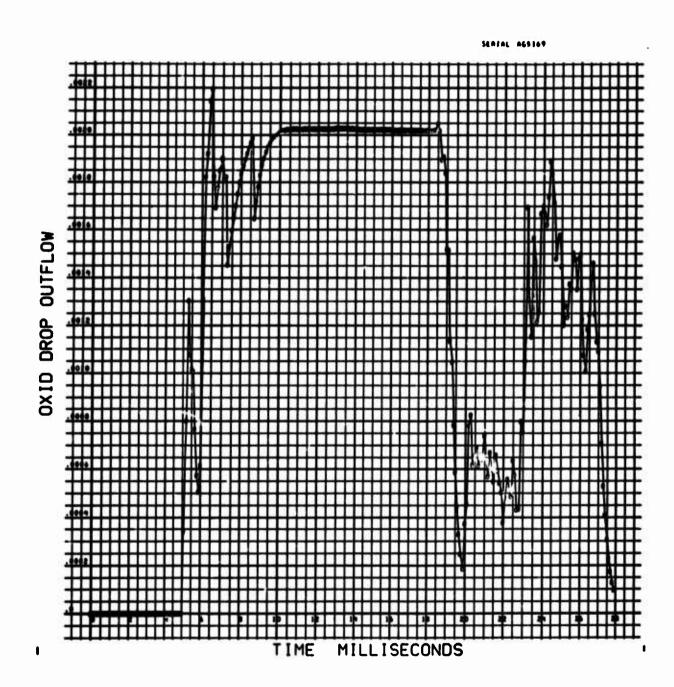


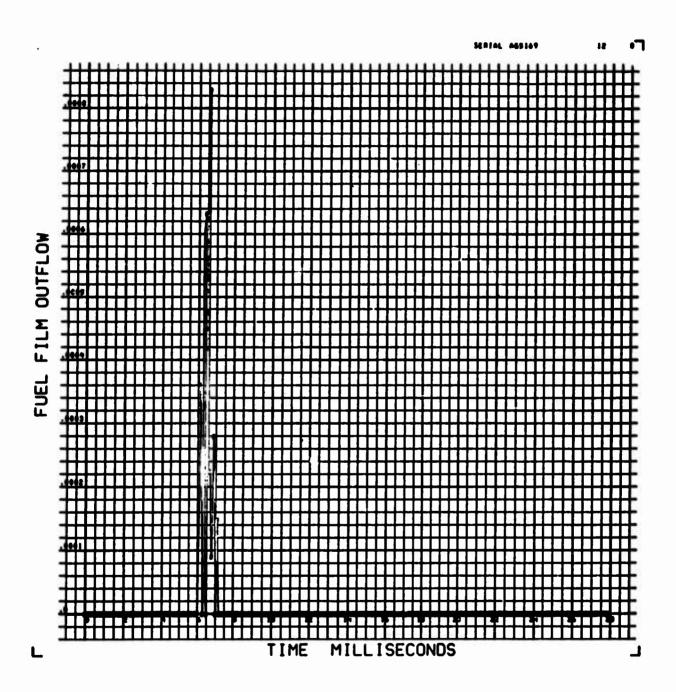


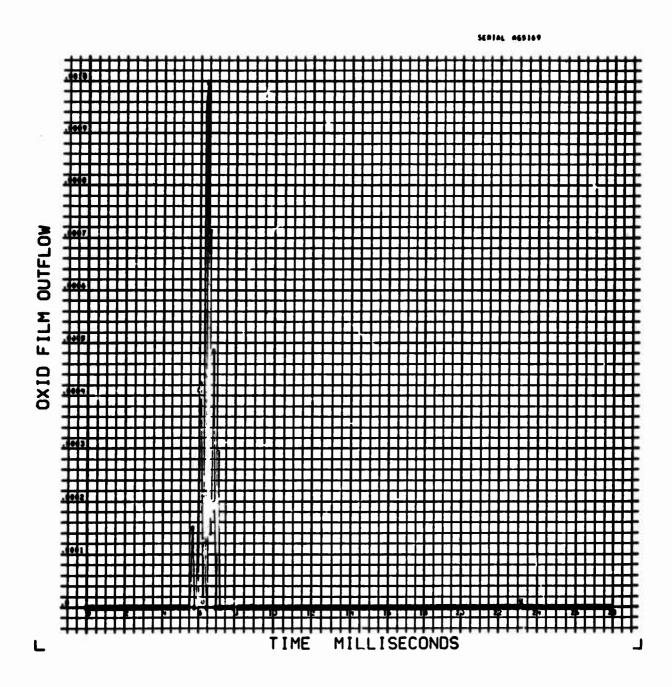


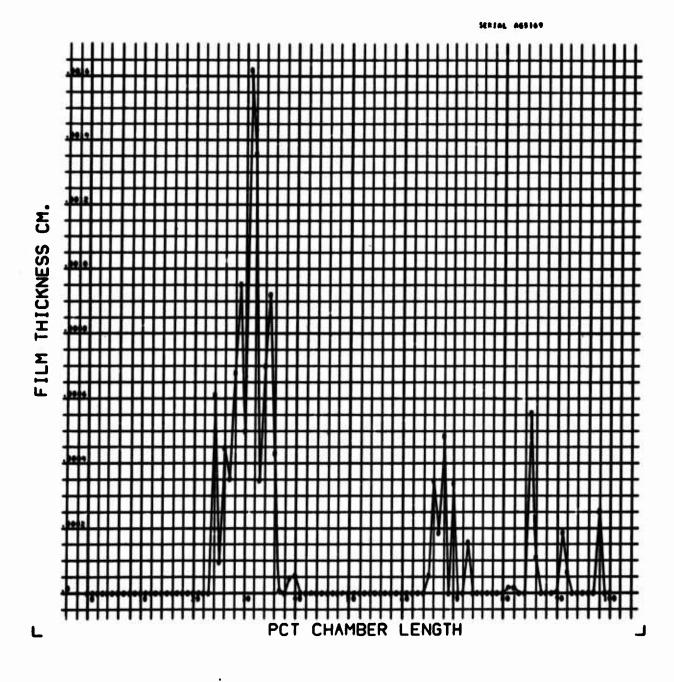


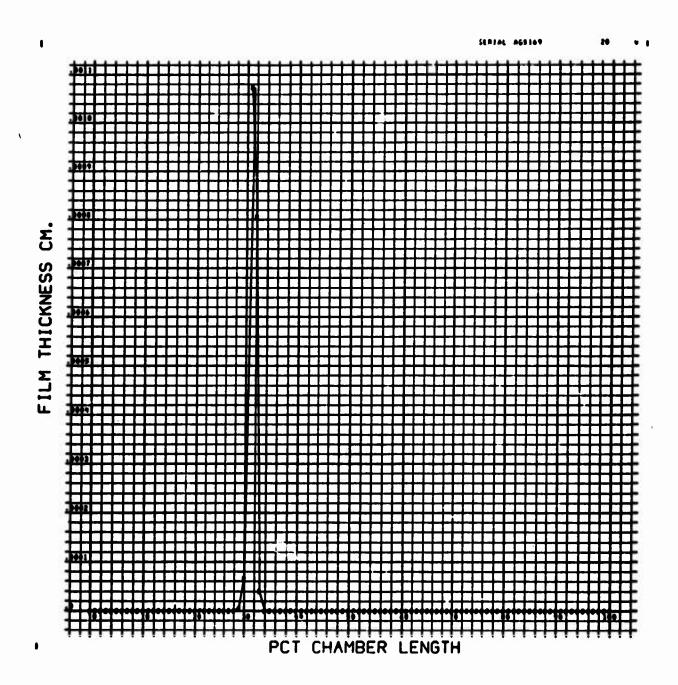


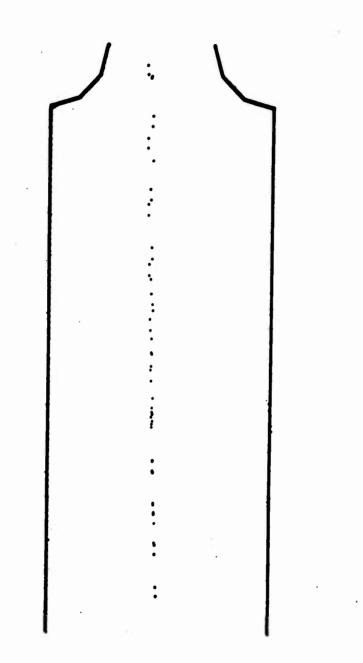












Initial computer runs were made with the total particle population as given in the "Expelled Droplet Size Distribution Table" of the <u>TCC</u> output, that is, five droplet radii ranging in value from 1.6410⁻⁵ to 2.78910⁻⁴ feet. The limiting streamlines proved to be so close to one another that to retain computation accuracy the program used an extremely fine characteristic mesh. After several minutes of computer calculation, the solution was still very close to the start line, and further determinations of the flow using all particles sizes were not made because of the great expenditure of computer resources that would be needed.

The "Expelled Droplet Mean Diameter and Velocity" table of the <u>TCC</u> printout suggested that drops with a diameter on the order of 50 microns would be highly probable in the flow of this particular rocket motor, and consequently for the example droplets of this diameter were used. It should be noted that runs with 85µ droplet diameters and droplet to gas mass flow ratios of 25 percent have been made with no numerical difficulties in MULTRAN.

The input data and the last few pages of <u>TD2</u> output are shown to illustrate the results of this part of <u>CONTAM</u>. The main function of <u>TD2</u> is to produce a start tape, tape 8, for the plume flow calculations of <u>TD2P</u>. The additional NAMELIST/DATAP/used in <u>TD2P</u> is also shown. It should be noticed that a Prandtl-Meyer expansion angle of 60 degrees about the nozzle lip has been used. The flow calculation was carried out for a distance of 40 throat radii along the axis of the nozzle. This number was chosen to keep the overall calculation within reasonable computer time limits (including <u>KINCON</u> and <u>SURFACE</u>) and to match the requirements of the data that will be needed by <u>SURFACE</u>.

As with TD2 the last few pages of <u>TD2P</u> printout are presented to illustrate the results of the plume flow field calculation. Again as with TD2 the main task of TD2P is to produce a carry-on tape, tape 12, to be used by the Streamline Generation Program, SLINES.

The card input to <u>SLINES</u> is accomplished via the short NAMELIST/SID/. For example,

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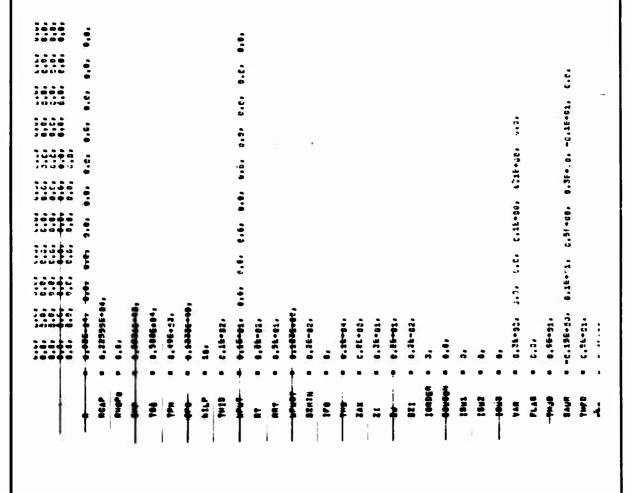
where the start of the NAMELIST begins in column two of the card. This input indicates that six streamlines will be determined for the limiting percentages of 0, 25, 50, 80, 90, and 99 percent of the total gas mass.

The output of this aspect of MULTRAN is adequately illustrated by the results for the streamlines shown. The printout is nondimensionalized or normalized in relation to the physical throat of the nozzle. Figures are given for axial, radial, and streamline distances. As will be shown below, the results of KINCON are given in terms of streamline distance, and the use of these tables will provide an avenue to interface the data points requested by SURFACE in axial and radial distance with the values given by KINCON.

A further discussion and intrepretation of the MULTRAN outputs will be presented in the SURFACE section when the determination of the later programs inputs are made.

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4. KINETICS AND CONDENSATION-THE KINCON PROGRAM

The KINCON subprogram performs the chemical-kinetic and single-species condensation calculations along gas-phase streamlines as defined by the Multiphase Nozzle and Plume Transport program, MULTRAN. When operated in the sequential mode, KINCON accepts card input data describing the initial gas-phase composition, chemical reactions with rate coefficients, and pertinent integration control parameters. The streamline definition including initial streamline conditions (pressure, temperature, and velocity) and the streamline pressure distribution are obtained from the MULTRAN program via TAPE 8.

a. Description of Input

The chemical system (12 species and 24 reactions) describing the kinetics of MMH/NTO combustion products is a commonly used set and was obtained from the <u>ODK</u> (the ICRPG One-Dimensional Kinetic Nozzle Analysis Computer Program) kinetics library (Reference C-1 and C-2 in Appendix C). The rate constants were updated whenever possible with the latest values from the literature. The initial gas-phased chemical composition was obtained from thermochemical equilibrium results corresponding to the values calculated by the <u>TCC</u> subprogram.

A listing of the <u>KINCON</u> card input data for this sample case is presented. Nothing is entered in the namelist \$ THERMO since a master tape of thermodynamic data (in JANAF format) was attached to TAPE 4. The 12 chemical species with initial mass fractions follow the title card. The chemical reactions are specified in two groups; the three body dissociation and recombination reactions appearing before card <u>END TBR REAX</u>, and the binary exchange reactions following the <u>END TBR REAX</u> card. The rate constants for the reverse reaction are input on the same card as reaction, which is input in symbolic form. Integration control parameters and miscellaneous data follow the reaction cards in namelist \$PROPEL.

The normalizing factor for the streamline coordinate was taken as the throat radius, RSTAR = 0.0972 in. Initial step size (HI), minimum step size (HMIN), and maximum step size (HMAX) are input in values of the normalized streamline coordinate. DEL is the relative error criterion and represents a measure of the truncation error in the finite difference scheme. Every tenth integration step is output as specified by ND3.

\$PROPEL RSTAR=0.0972, JPFLAG=1, DEL=0.01, ND3=10, HI=0.0001, HMAX=1.0, TCOND(1)=12*0.0, T=5881.8, AREPS=0.001, ETA=0.1, IDCOND=2, \$END

At the initial step of each streamline a table of "Dissociation Recombination Reaction Rate Ratios" is printed as is shown. The kinetic results determined along the streamlines are based on the pressure at that point, and accordingly the next item printed out is a "Pressure Table", as illustrated. Results are shown for the 25 percent mass limiting streamline.

A further discussion and interpretations of the results of KINCON will be made in the next section when the inputs for SURFACE are determined.

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	-4	17 6.	3	34 85	53	AV 62	•	AC 643	167	163 -ACH	123	120 W2K1	
	-4	28 5	2	39 BF2		* W P C 9	4	B2H0	101	4	121	121 VHSK	
		19 SF4	*	46 973	3	61 NAS	2	8202	102	274	122	¥	
	8	20 57.0	7	41 BUCL	7	CF2		8029	101	7	123	1203	
ļ	~	21 C	*	108 27	**	5 40							

8	0,2017			
C02	4,52E-02			
2	0.020			
2	0,42509741			
9	0,002			
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05	0.000764			
U	1,696-11			
02.1	0,2736			
*	0.00115			
2	1.3vE-06			
0	6,175-0+			
PEACT IONS			•	
02 . 2.0,	A83.3E-17,	401.7	P=0,3,	RFACTION
N2 = 20N,	4.9.66.17.	WBS.C.	80.0.	REACTION
10 . N . O.	487,26-15,	15130%	.0.0.0	REACTION
MS - 20Hg	A.5.0E-14,	Net.	M.0.0.	REACTION
M20 - H - OH,	401.176-17.	NaC . D.	40.08	REACTION
0 · H · HO	A#2.36+10.	2.0.V	40.04	REACTION
602 = C0 + 0,	A=4.1E-14,	40.0 a	8=3,58.	REACTION
.0 • C • 0;	A=6.0E-3:	AC BOOK	H=50.0,	REACTION
END TUR REAK				
*: + 20 = 0 + 0M	A#3.6E+11.	46.0-5	Ne+6,5, He7,15,	REACTION
NO + N = N2 + 0s	A=5.0E=13,	10101	9e75.5r	REACTION
N2 + 02 = 2+10.	A=1,0E+13,	NEO. P.	AE79,468.	REACTION
\$6 • 62 • C02 • 0.	A-1.9E-13,	*c *0 * N	Na0,7, 8854,13,	REACTION
CO + N = C + NO.	AB1.3E-1.,	weed.1. Hed.5.	4-0.5.	RFACTION
60 + 0 + C + 02	AB2,4E-13,	¥80.7	VED. 1. HEL, 09,	REALTION
CO+ H + C + 04	A-1, 2E-14,	NEC.T.	NEG-TP HE25,65.	REACTION

2 : 2

Per - H = 00 - 0H.	ABS.66-11, NUO.01 001.08,			MEACTION 10
190	***************************************	***	1019100	MEASTION 17
OH - HE M HED - M.	ASS. 41513, Neg.0, 8028,1,	Neel to	0.50,1,	REACTION 18
3+0H = H20 + 0,	Ass, 79613, N=0,3; 0=16.0;	N=0,3+	0.510.0	REACTION 19
*******	407,53612; Ned.0; 840;0;	10 7 0 01	.0.0.0	REACTION 20
hg - 02 # 2-04,	444,98623, Nº2,9, 8985,7,	No.2,9,	9005,7,	REACTION 21
NO - CO - COZ - N.	Am1,0613,	46 0 0 N	900,93,	REACTION 22
** • # • B# • #*	A-3, 4613,	40.00	0-1,36,	MEACTION 23
CO - CO + COS - C.	A-1,0613,		120,000 10.000	REACTION 24

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200000000000000000000000000000000000000		7.200000E-19	513000006010	± •	#20 0 0717	2,300000E-16 OH	3-1000006-13 CO2 - 0	0 • 3	3.0000006-11 B	M-000000000000000000000000000000000000	4. 000000013 NO • 02 NO • NO	1,9000006-13 C	00 00 00 00 00 00 00 00 00 00 00 00 00	27-300007-19	
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BISCOCIATION RECOMBINATION MEACTION RATE RATIOS

ALL REACTION MATE MATIOS INPUT AS 1.0

9 1 4 4 6	9079K	-1,06006016-01	-1,76737941E-01	** 10-96-96-96-91	-2,00092731E-01	-1,962205106-01	10-900010160-01
ME30000E4	•	6,85586798E+01	6,67200017E-01	10-90036400619	6,32247246E-01	6112981941E+81	10-3000000-016 - 30-8000000-0-1
	*	-	. 96664166-62	78-9404080541	1.000409216-01	1.624900678-01	

-8,346466866-01 -8,394218646-01 -2,364277966-01

4,84278906E-01

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-2,230784806-01 -2,307731206-01

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	1.145885656+00	4,300074326-01	-2,366663106-01
:	1.241214788-00	4,16401921E+01	-2,34678916E-01
*	- 24-B68787888-13	24-945348534912	10-90140968614-
:	1,432890968-00	3,72348296+01	-2,24620464E-01
13	1.527575528+00	3,911864796-01	-2,18792539E-61
#	\$1.000000001.2		
2	1.710633746+00	3,000013936+01	-2,09564601E-C1
2	1,414229116-00	2,90203996E+01	-1,54734302E+00
*		10-91011001012	6-384060056-03
2	1,982646978-88	2,903446716+01	-2,0471147F-01
2	2.3873999E-00	2,69121237E+01	-1,07071173E+61
*		21-404-0006-01	*************
2	2, 33628461E.00	2,22276180E+01	-1,71397207E-01
2	2,465964776-00	1,961392466+01	-1,546494276-01
	30-300000G0'3	***************************************	************
2	2,86444926E+00	1,46189896+01	-1,14794449E+81
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#	3,700000128-00	7,039129146-00	-5,276721276-00
=	4,27405906-11	4,00181208E-98	-3,14684898+80
*	*********		40-948806809-I-
*	9,766939716+88	2,48409900E+00	-9,967443895-01
2	7,342848496+00	1,417727146.00	-4,15297899E-01
		\$120-400576-00	10-92613609616-
ñ	0,335177196-00	1,044202476+00	-3,07084829E-01
=	0.756899372-90	9,22960196E-01	-2,57014426E-01
\$		10-96403040016	20-30170176179
=	9,91196991E-0C	6,795488346-01	-1,68841648E-01
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Ŧ	1,103617026+01	4 31472883E-01	-0,42400205E-02
7	1,292005586+01	3,45-47430E-01	-6,81377149E-02
:	1.417605286-01	2,72550030E-01	-4,80723983E-02
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:	1,743244835+01	1,000363816-01	-2,35504378E-02
47	1,302979406+01	1,309022556-01	-1,53380178E-02
:	1,085216896-01	1,309016736-01	-5,09144302E-03
•	1,399718746+01	1,2012606E-01	-1,51457966E-02
2	2,382314046+01	1,00312037E-01	-1,37250061E-02
=	2, 234619516-01	9,02015792E-02	-0,71172113E-03
25	2,94756432E-01	9,92441386E-02	-7,16309705E-03
2	2,887401466-01	4,394244356-02	-4,41346227E-03
I	3.265929306-01	2,941317786-02	-3,156419736-03
93	3,970105166-01	2,17384446E-02	-1,00707350E-03
2	4,13671494E+01	1,364130846-02	-1,44072430E-03

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CALCULATION
STREAMTUBE
KINETIC
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FLOW PROPERTIES		KINETIC COUPLING TERMS	
MACH NUMBER PRESSURE (PSIA)	5,918423386-61 6,85867586-81	COUPLING TERM A	-0.67909441E-01 1.28416828E-90
TEMPERATURE (DEG-R)	5,601,00006-03	INTEGRATION PARAMETERS	The state and the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the
	2.964967256.63	CORPORATION STATE STATE	1.000000000
MAT CAPACITY (BTJ/LB-DEGR)	5,273461426-01	2.0 - SUPPATION COLD	-1,00540737E=11
AAS CONST(FT2/SEC2/DES-R) 2,926445346+03	2.526445346+03	CONFIDENCE MEDIATION PRODUCTION	

CHEMICAL COMPOSITION

	1	HOLLOWA BOWN	HOLE FRACTION	#0168/00	*	Precies	-WO. PPECIED AND PARCTION HOLE FARCTION HOLEGAGE NO. SPECIES MASS FARCTION HOLE FARGTION - MOLEGAGE	HOLE FRAGILOR	************
	8	2,617000E=01	2,617040E=01 1,417122E=01 1,4830E=18 2 CO2	1,4838E-18	~		6,920000E-02 2,919528E-02 3,0927E-17	2,919923E-02	3,09276-17
	*	2,000000E=02	2,000000E=02 1,992433E+01 2,0443E+10	2,04436+18	4	7	4,250976E-01 2,986197E-01 3,1267E-18	2,986197E-01	3,12676-18
	*		£1-354465E-03	1193966612	•	1		21-1100011-00	2147266027
_	20 4	7,640000E-04	7,640000E=84 4,698654E=04 4,9198E=15	4,91986-15	•,	U	1,0900006-11 2,760003E-11 2,0094E-80	2,7690938-11	2,00046-00
	430	2,738000E=01	2,730000Em01 2,000922E-01 3,1317E-10 10	3,13176-10	10	x	1,1900806-03 2,2452646-62 2,35896-17	2,2452646-62	2,35006-17
		A APPROACH A SECRET AS A ASSESSMENT AS A APPROACH A SECRET AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH AS A APPROACH	*********	A 8448E447		•	4 490000		

KI ETIC STREAMTURE CONSITIONS AXIAL POSITION 1,90000E-03

FLOW PROPERTIES		AINETIC COUPLING TERMS	
PRESEURE (PSIA)	5.91980764E=01	COUPLING TERM DOUBLING TERM	-8,70484298E=01 1,19987304E+00
TEMPERATURE (DEG+R) DEMBITY (LB/FTS)	3.8733734E-03	INTEGRATION PARAMETERS	
BATMALPY-HO (97U/LE) GAS HOLECULAR WEIGHT	2 *** ** ** ** ** ** ** ** ** ** ** ** *	PERCENT STEP SIZE	2,00000000E=04
MEAT CAPACITY(HTU/LR-DEGR)	5,2722962CE=01	1,0 - SULMATION C(1)	•
BAS CONSTITY2/SEC2/DEG-4) BUN C(1)+H(1) (F72/SEC2)	2.52763043t-03	MARIMON MELATINE EMBOR SOVERNING EQUATION	4, 3047461 PE-63

				rutulcal contotivios	-	ž			
9	SPECIES	HASS FACTION	NO. SPECIES MASS FRACTION VOLE FRACTION MOLEC/CO.	MOLECZCC	9	SPECIES	NO, SPECIES MASS FRACTION HOLE FRACTION HOLEC/CC	HOLE FRACTION	#OLEC/CC
-	00 →	2,017270E=01	2,017270E=01 1,410051E=01 1,4050E=18	1,48566+18	~	2 002	6,915671E-02	6,515671E-02 2,012221t-02 3,0940E-17	3,09406-17
*	*	2,003513£=02-	2-003375-02- 1.094746-01 -16400E-16	01-300+01e	•	4 #2	4,2549766-61	4,2580766-81 2,0047566-81 3,53886-18	3,13006-10
ň	9 %	1,0009186-03	1,999918E+03 1,316977E+03 :13748E+16	: 13740E+16	•	30	1,336466-02	1,336446E-02 1,198726E-02 1,2971E+17	1,29716-17
^	20 /	7,6011276-04	7,4011276=04 4,472554E=04 4,9000E+15	4,90006-15	•	υ	1,8234346-11	1,823434E-11 2,986329E-11 3,1327E-88	3,1317E-08
+	2	8,7307 346 weg.		211300E-14	97	1	1,1562256-63	1,1562256-63 2,2563596-62 2,36636-17	2,36636-17
#	33 N	1,4368376-08	1,436837E-08 2,017734E-04 7,1159E-13 12 0	7,11596-13	15	0	4,242380E-04	6,242380E-04 7,674619E-04 8,8482E-19	8,0402E-15

KI'ETIC STREAMTUBE CONDITIONS AXIAL POSITION 3,90000E-03

PLOM PROPERTIES		AINETIC COUPLING TERMS	
MACH NUMBER	5,93282866E=01	COUPLING TERM A	-8.09136830E-01
PRESSURE (PSIA)	6,84434830E+01	COUPLING TERM IN	1.09693470E+00
Vacacity (PP/SEC)	2.53948948618		
TENPERATURE (DEG-4)	5.85629834E+03	INTEGRATION PARAMETERS	
DEMBITY (LB/FT3)	2.14894886E-02		
BETWELLTONE (BTL/LE)	2.40149337E+B3	CURRENT STEP BIZE	2.86000000E-04
DAS MOLECULAR METGHT	1.04993392E+01	PERCENT PRIMALPY CHANGE	-7. 93211699E=03
MEAT CAPACITY(BTU/LB-DEGR) 5,270910506-01	5,27091050E-01	1.0 - SUMMATION CCI.	-1.67972303E-11
PROSEN BENNA	1.247290906-00	THE PARTY RELACTOR ERRER	NO SO SO SO SO SO SO SO SO SO SO SO SO SO
3EG2/DEG-R)	2.529129316-03	GOVERNING EQUATION	19
BUM C(I) OH(I) (F72/SEC7)	1,132763376-87		•
	CHEN	CHENICAL COMPOSITION	

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Ī	19 PRICTION	-we, opecies mass parchies note function notecom	HOLEGYCC	ş	SAECIES	NO. SPECIES HASS FRACTION HOLE FRACTION HOLEC/CC	HOLE FRACTION	*0.66/66
~	0175556-01	2,0175956m01 1,4160386m01 1,48776+18	1,48776-18	~	2 002	6,511289E-02 2,908536E-02 3,8558E-17	2.906536E-02	3,09906-17
~	.006179E-02	2,004179E=02 1,056357E=01 2,0954E-18	2,09546+10	•	4 42	4,250976E-01	4,250974E-01 2,982987E-01 3,1348E-18	3,13406-10
Т	20m3220000 11	######################################	37496-1	•	3.	1,1083436-02	1,1083436-62 1,2011286-52 1,34666-17	1,34666-17
•	7.57e849E=84	7,57e849E=84 4,654379E=84 4,8988E+15	8900E-15	•	v	1,9984266-11	1,998426E-11 3,257888E-11 3,4228E-08	3,42206.00
	2,7230406-01	2,723896E=01 2,071476E=01 2,1219E=18 10	1219E-1	10	1	1,1713146-03	1,171314E-g3 2,284450E-g2 2,4881E-17	2,4001E-17
- 1	1. +025+3t=00	13 4 1,492543twge 2,9946e9t-96 p,Red?te-18 12 0	P8076-5	4	•	+,	4,496866-64 7,9852815-64 8,38946-19	6,36946.18

MILETIC STREAMTURE CONDITIONS AL POSITION 9, 900006-03

PLOW PROPERTIES				
00-00-00-00-00-00-00-00-00-00-00-00-00-	FLOW PROPERTIES		AINETIC TOUBLING TERMS	
G48) 9.64329376443 INTEGRATION PARALETERS 1.244334976402 COULENT SIZE EGAT 1.644896461 PERCENT SIZE FIZEBET 1.6448396461 PERCENT SIZE EZZZGGRA 9.2792642461 L.J SOURATION CETS EZZZGGRA 1.27934224640 MARINUM PERCENT CETS FFZZEGRA 1.31401966667 SOURATION CETS 1.3140196667 SOURATION CARACTERION CETS FFZZEGRA 1.31401966667	PART ACTORN PARSOCAR (PRIA)	5.943792848-01 6.946819478-01	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	-7,45537020E-01 1,01772160E-08
	PREPREATURE (URGLE)	9.04020378E-03	INTEGRATION PARALETERS	
FIZEBER 5.2/967135=01 1.0 - Scumatio, cf1) 1:2/92/92/04/92/92/92/92/92/92/92/92/92/92/92/92/92/	BATTALPTO (DTD/LD)		PRESTANTA STATES OF THE STATES	2,00000000E=U4
E.22/36G=#) 2-1/2-01/2-01/2-02	MEAT CAPACITY(STI/LE-DEGR)	5,2,0671196-01	1.0 - SOUMPTION O(1)	-1.67799140Ee11
	945 CONSTITUZ/SE12/3EG-#) SUM C(1)#M(1) (F72/SEC?)	1.5 130549246-03 1.1 11001006-07	1014164 LVICESON	0

4,5-8063E-62 2,909443E-62 3,8977E-17 4,2909746-01 2,9813186-01 3,13796-18 1,163647E+02 1,344293E+02 1,4147E+17 2,140819E-11 3,911988E-11 3,6998E-88 1,141496E-03 2,322438E-02 2,4441E-17 6,652773E+34 8,419347E+04 8,8962E+19 NO. SPETIFS MASS FRACTION HOLE FRACTION HOLEC/CC 2 005 ... • • • 1 0 , 029 SE - 16 - 37 ADE - 10 4,89866-15 1,1184E-18 1,542234F=00 7,163179E-00 :,2769E-13 2,017744E=01 1,4153-2E=01 :,4485E+18 NO. SPECIES HASS FRACTION HULE F'ACTION MOLEC/CT 2,007743F-92 1,9567:7E-01 1,999742E=03 1.309371E=03 7,560841E=04 4,654741E=04 2,714940E=31 2,9631,6E=01 2 1 2 7 02 . v 3 2

PEMICAL CUMPOSITOR

HINETIC STEERNTUSE CONDITIONS AXIAL POSITION 3,340996401

					MOLEC/CG	7,7633E+14	7,10376-19	2,31906-13	1.65166.09	9,5422E-14
	1,024794196-05	1.62400E+01	10 1 4 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1		HOLE FRACTION	3,2164674-02	2.9763898-01	***********	4.842926E-12	3,9534321-02
TERPS	1.00		•		NO, SPECIES MASS FRACTION HOLE FRACTION MOLEC/CG	7,216597E-02 3,2164g7k-02 7,7633E+14	4,250928E-01 2,976309E-01 7,1837E-19	8,310096E-64 9,5944eb-64 8,3199E-13	4,190070E-12 6,842926E-12 1,6516E-05	2,031549E+03 3,953432E+02 9,5422E+14
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d I NE				THENICAL COMPOSITION	MOLEG/CC	133426-15	1,4422E-15	1117506-13	1.0472E-13	130506-19
	6.05704573E-00 2.69181152E-02	1.002998286.00 1.002998286.00 1.002998286.00 1.0029988886.00	4) 0.8724972668 1.044928668 1.04498668 1.044799668 44.646988		- 48, SPESIES MASS FARETISM MOLE FUAGTION MOLEC/CC	1,972645F-01 1,3814;6E-01 1,3342E-15	1,801552t-02 1,848445E-01 4,4422E-15	P. 0120576w03 1:3156PEF-03 4117506-13	1,000024E=03 6,741759E=04 1,6472E+13	2,802941E=81 3,0517'5E-01 7,3059E-19 10
147168	1981A)	TEPERATURE (DEGAR) DENSITY (LB/F13) DMTMALPT-MG (810/LB)	MEAT CAPELITY(011/18-DEGR) AND CONSTITY(011/18-DEGR) BUT C(11-4(1) (FT2/SEC2)		MASS FARETIBE	1,9726656-01	1,8015524-02	2,0120576-03	1,0000246-03	2,8029416=81
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2,8667715-84 3,5398946-84 8,84216-13

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RIVETIC STREAMTURE CONDITIONS ANIAL POSITION 3,401106-01

Flow Properties		KINETIC COUPLING TERMS	
ACH NUMBER	2,418764146-82	COUPLING TERM A	9,19687383E-86 -9,1727362E-86
PAPERATURE (DEG-R)	9.738695646.02	INTEGRATION PARAMETERS	
PERSONAL CONTRACTOR	1.047401206-02	COLDENT STEP BIZE	2.040000000
AS MOLSCULAR MESONT	1.961991736-81	PRECENT ENTRAINT CHANGE	1.61799786601
BALL CAPACITY COLONIAL STREET		Tree-wer of the City	
AB 60467(F72/8662/DE0-F)	2,934779906-03	BOVERS EDUATION	

	1458 FRACTION	NO. SPECIES MASS PARCTION HOLE FARCTION MOLEC/CC NO. SPECIES MASS FRACTION MOLE FRACTION MOLEC/CC	MOLEC/CC	÷	SPECIES	HASS FRACTION	HOLE FRACTION	10/8C/CC	
	1,9726546-01	1 G0 1,9726546m81 1,3814;9E-01 3,8797E-15 2 C02 7,21643E-02 3,216481E-02 7,16146-14	3,07976-15	~	200	7,2167936-02	3,216401E-02	7,16166-16	
	3003006300"3	67-960000 11-060006014 11-90006014	4100306049	+	+	**************************************	18-968696943	***************************	1
	E. 012057E-03	2,012957E-03 1,319022E-03 2,9296E-13 6 0w	2,92966-13	•,	ī	0,200420E-04	8,2564206-04 9,5499016-04 2,12636-13	2,12636-13	
	1,1000306-03	1,1888286-03 6,7429596-84 1,50136-13	1,56136-13	•	v	4,1A9778E-12	4,189779E-12 6,842437E-12 1,9834E-83	1,92346-63	
T	100388886919	67	-97-9010c-1-	*	*	2,034546645	2-938000-0- 20-9007490-2 20-909880-2	*******	
	1,6783696-67	1,678369E-07 2,539054E-07 5,2878E-09 12 0	5,2070E-09	7		2,9051436-04	2,9051436-64 3,5616176-64 7,92986-12	7,92966-12	

KIMETIC STREAMTURE CONDITIONS AKIAL POSITION 3,625826-81

		MOLEC/CE	6,4951E-14	9,0730E-19	1,00476-13	1,37366-09	7,03306.14	7,21206-12
1, 02400000000000000000000000000000000000	2 - 30074004505	HOLE FRACTION	7,210072E-02 3,210574E-02 6,4951E-14	4,250929E-01 2,976309E-01 9,9730E-19	8,2292916-64 0,409000-84 1,00476-13	4,1894816-12 6,8419346-12 1,37366-89	2,031928E-83 3,093384E-82 7,0336E-14	2,9316496-84 3,5041146-64 7,21206-12
		NO. SPECIES HASS FHACTION HOLE FRACTION HOLEC/CO	7,2160726-02	4,250929E-01	8,2292816-84	4,189401E-12	2,031520E-03	2,9316496-84
MINETIC COUPLING TERMS COUPLING TERM B INTEGRATION PARAMETERS CUMMENT STER SIZE PHOTON TOWATON CLINCE PHOTON TOWATON CLINCE PHOTON TOWATON CLINCE PHOTON TOWATON CLINCE PHOTON TOWATON CLINCE PHOTON TOWATON CLINCE PHOTON TOWATON CLINCE PHOTON TOWATON CLINCE PHOTON TOWATON CLINCE PHOTON TOWATON CLINCE PHOTON TOWATON CLINCE PHOTON TOWATON CLINCE PHOTON TOWATON CLINCE PHOTON TOWATON CLINCE PHOTON TOWATON CLINCE PHOTON TOWATON CLINCE PHOTON TOWATON TOWATON CLINCE PHOTON TOWATON T	COMPOSITION	. SPECIES	2 Cr2	F.	å	U	1	0
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		0m mule F4,	1,30139	1,8464	1,85900	8 6.74495	3,09182	* 1,380m
FLOW PROPERTIES MACH WUNDER PRESSURE (TELA) TEMPERATURE (DEG-R) DEWRITY (LDFT3) BENTALLY (LDFT3) BAS MOLECULAR HEIGHT MALY CAPACITY (STATA)	Freger Garma Bas Comst(f72/SEG2/DEG-H) But C(1)+H(1) (f72/SEG2)	-401 -00E0166-4466 F486710m mule F446710m MOLEC/CC	1,972641E=01 1,381339E=01 7,7722E-15	1,801545E=C2 1,8464V7E=01 3,6936E-15	E-012007E-01 1,01502E-08 p.6406-13	1,100240E=03 6,744952E=04 1,3959E+13	2,402949E=01 3,091022E=01 4,1249E-15	25 -0
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FINETIC STREAMTURE CONDITIONS AXIAL POSITION 3,09367E-01

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.0u PROPERTIES		KINETIC COUPLING TERMS	•
IGH NUMBER ISPANS (PSIA)	1,941767846-00	COUPLING TERM A	9,70082196-00
MODEL TURE (DES-R)	0.544727406-02	INTEGRATION PARAMETERS	91
Property of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of the contract of th	1.100040715-02	SUMPRIT STEP SIZE	4,00600006-11
LAT CAPACITY (STU/LS-DESA)	3.0,0201306-01	1.0 - SUMMATION CELL	1, 430130906-11 -1, 430130906-11
18 COMET (FT2/8EG2/DEG-R)	2,934779366-83	GOVERNING SQUATION	21

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3		HASE FRACTION	NO. SPECIES MASS FIACTION HOLE FIACTION HOLEC/CC	MOLEC/CC	£	SPECIES	NO. SPECIES MASS FRACTION NOLE FRACTION MOLEC/CC	HOLE FRACTION	HOLEC/CC
*	8	1,9726106-01	1 CO 119726196-81 1,3813428-81 2,28448-15 2 CO2	2,26446-19	~	200	7,2173396-62 3,2167366-82 5,27396-14	3.2167306-02	5.27306-14
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•	7	2,0129976-03	2,012997E-03 1,315022E-03 2,1969E-13 6 0H	2,19696-13	•.	ē	0,13000E-04	8,139000E-84 0,377562E-84 1,93726-13	1,93726-13
~	8	1,1007006-03	1,1807006-03 6,7470146-04 1,1001E-13	1,10616-13	•	v	4,1007036-12	4,1867836-12 6,8488266-12 1,12146-89	1,12106-09
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11		1.6672346-07	11 N 1.667234Em87 2,334599Em87 3,8278Em89 12 0	3,02706-19	77	•	2,9492996-14	2,9892996-64 3,6647366-64 6,68746-12	4,0874E-12

MINGTIC STREAMTURE CONDITIONS ANIAL POSITION 4,136715-61

PLOW PROPERTIES		KINETIC COUPLING TERMS	
PRESSURE (PSIA)	1,304130846-02	COUPLING TERM A	9,00000006-00 -5,1001033E-00
	1.216291996-02	INTEGRATION PARAMETERS	
•	.374470401-02	CURRENT STEP SIZE	
-	1,941551894-01	PRICENT RETEALPY CHANGE	
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4,136714946+01	1809080769871	2.091000006-01	2,7390000E-01	-4-1296724946-91	2,74700005-01	4,13671494E+01	4,136714546-01	4,134714946-61	1,709598666+90	4,136714546+01	4,136714546+31	1,7950800E-81	1,635000006-01	4,136714946-01	2,891000006-01	7,3735000E+60	4,21500000E-01	3,1230000E-01	2,9230000E-01	4,136714546-81	4,136714546-81	4,13671494E+81

5. SURFACE EFFECTS DETERMINATIONS

The calculation of the effects of the impingement of the plume gas, large fuel and oxidizer droplets, and small condensation droplets upon the space venicle appendages and surface is carried out in the segment <u>SURFACE</u> of <u>CONTAM</u>.

SURFACE is discussed in detail in Appendix D and the reader is referred to that section of the report for a complete description of all program overlays and inputs. The material that will be covered below will be directed at the problem of interfacing the outputs of TCC, MULTRAN, and KINCON with SURFACE.

It is assumed that the user has operated <u>SURFACE</u> under "STPFLG = 1" and has obtained a table listing the surface segments that the plume will impinge upon. In the sample case, a "p" value (one-half semi-latus rectum) of 0.14 has been used as a nominal value to obtain the data for thrustor No. 2, as illustrated.

Thrustor No. 2 is located 6 inches off the skin of the vehicle and 3.6 feet back from the leading tip. The thrustor is part of the RCS.

Inspection of the table "Thrustor-Based Coordinates of Plume-Impinged Segments" reveals that segments 52, 62, 72, 82, 91, 92, 93, 101, 102, and 103 will be impinged by the thrustor plume. The outputs of the preceding programs will be interfaced with the input data required for <u>SURFACE</u> at location 52. This will amply demonstrate the techniques required to solve for the surface effects on any segment on the vehicle.

Referring back to the first kinetic streamtube condition printout of KINCON, the "Input Normalizing Axial Scale Factor (FT) is approximately 0.008. This figure is simply the throat radius of the nozzle given in the proper system of measure. By dividing the values obtained from SURFACE by this factor the whole problem is quickly converted into the normalized units employed in MULTRAN and KINCON, i.e., R/RC = 62.5 and Z/RC = 215.6.

At this point it is suggested as good practice to make a plot of R/RC vs Z/RC for the particle limiting streamline, obtained from the printout of <u>TD2</u> and <u>TD2P</u> and the gas limiting mass streamlines, obtained from the printout of <u>SLINES</u>. The location of the nozzle wall can also be ascertained from <u>TD2</u>.

Referring to the first page of the characteristic calculation printout of the Gas-Particle Flow (Nozzle), e.g., <u>TD2</u>, any streamline identified by a "4" is part of the limiting particle streamline, and any printout identified by "5" is part of the boundary or in this case the nozzle wall. When looking at <u>TD2P</u> output the latter identification is not of concern since the calculations of <u>TD2</u> end at the exit of the nozzle. The limiting particle streamline, however, continues and may be traced out to the end of the <u>TD2P</u> solution.

THRUSTOR-SASED COORDINATES OF PLUME-IMPINGED SEGMENTS

FROM THRUSTOR NO. 2 MMEN Ps .14

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* AMELE PROM LINE JOINING AXES OF SATELLITE AND THRUSTOR

USE COORDINATES I AND W IN DETERMINATION OF PLUME CHARACTERISTICS FOR IMPUT TO REPLACE DEFAULT VALUES ON EDATA & TAPFII

The printout of <u>SLINES</u> or the Streamline Generation Program gives a table of various gas phase limiting mass streamlines in terms of Z/RC, R/RC, and S/RC. A direct plot of the first two items will produce the needed curve and correlation of this data with the last item will allow identification of proper <u>KINCON</u> information that will later be required as input to <u>SURFACE</u>.

The drawing shown displays the particle limiting streamline and the mass limiting gas phase streamlines for the values of 0.0, 25.0, 50.0, and 80.0 percent of the total mass. Inspection of this graph quickly reveals that the required location, as given by <u>SURFACE</u>, is far beyond the calculated data of CONTAM to this point of the solution.

When dealing with very small RCS engines, the normalizing factor of the throat radius will produce a characteristic grid mesh so fine in comparison to the characteristic size of the space vehicle that a computer solution taken out to this point is very costly. However, inspection of the streamline plot and the outputs of KINCON to the point they have been carried out demonstrate that the complete MOC calculation is not necessary. The streamlines have all become straight lines and the chemistry has frozen, thus enabling accurate and relatively simple extrapolation.

The methods suggested to obtain the required information at the called for <u>SURFACE</u> segments will now be discussed item by item, including the assumptions used whenever an extrapolation of <u>MULTRAN</u> or <u>KINCON</u> results is carried out.

Attention is directed to Appendix D. The array PLMCHR of the data set EDATA contains the needed input information for each of the segment locations called for. Note should be taken of the proper formats and the fact that three blank cards must precede input of the item NPMCR. These three cards may contain comments as they are not read by the program.

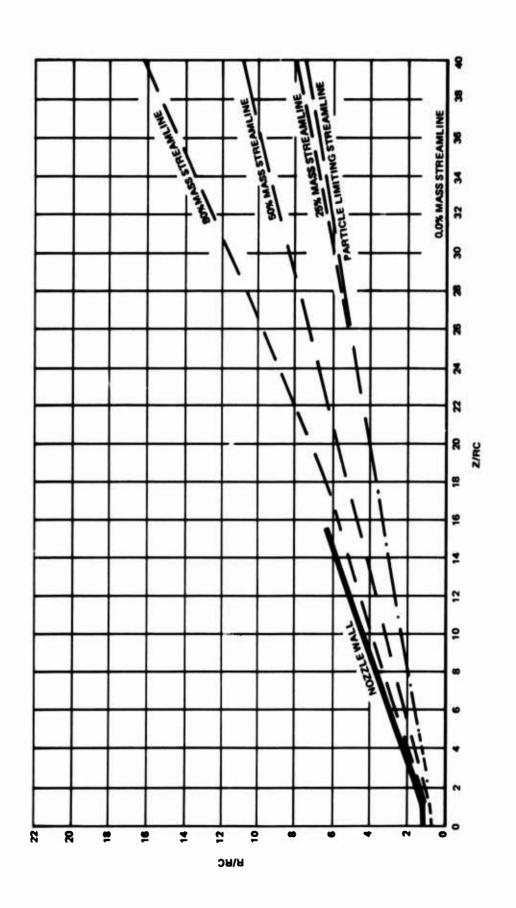
Extrapolation of the 25 percent mass limiting streamline and the particle limiting streamline data to the approximate region of interest on the surface have been used to obtain the data that will now be discussed.

PLMCHR(1) and PLMCHR(2)

SURFACE always assumes that two drop sizes will be given. The small droplet diameter will represent any products of condensation occurring in the plume, and the large droplet will refer to the fuel or oxidizer drops as predicted by TCC.

If the region of the plume of interest is hot or the density is very low (i. e., very few nucleation centers), no condensation will take place. In cooler regions of the plume of sufficient density, the condensation droplets formed will usually have a diameter of the order of one micron or less (3.94E-05 ft). This value is suggested for PLMCHR(1).

The large particle size was taken from <u>TCC</u> and <u>MULTRAN</u>; in this case 50 microns (1.97E-03 ft), and this value was used for PLMCHR(2).



PLMCHR(3)

The small droplet is assumed to move at the gas velocity. At the end of the <u>KINCON</u> solution the chemistry is frozen as indicated by the constant values of molefractions and the velocity of the gas is nearly constant.

Consequently, assuming the velocity to remain constant in value from this point will produce little or no error in the kinetic energy imparted to the surface of the vehicle. Accordingly, the last velocity figure from KINCON is taken as the input for PLMCHR(3), e.g., 11,417 ft/sec.

PLMCHR(4)

The item VPK in the output of <u>TD2P</u> is the particle velocity in units of ft/sec. The above argument certainly holds also in the case of the particles, and the last value given in <u>TD2P</u> is taken as input fot PLMCHR(4). Referring back to the <u>TD2P</u> printout identified by "4", the value of 1,697 ft/sec is obtained.

PLMCHR(5) and PLMCHR(6)

The parameters of THETA-G and THETA-K of the <u>TD2P</u> output give the gas and particle streamline angles, respectively. Since the streamlines has become straight, it introduces no error to take the last values given for the particle limiting streamline as the inputs for PLMCHR(5) and PLMCHR(6). Care must be taken, however, to make sure the values given in <u>MULTRAN</u> in degrees are converted to radians, e.g., 0.170 and 0.218, respectively.

PLMCHR(7) and PLMCHR(8)

Since there are no small droplets present in this flow a very small number is input into PLMCHR(7), in this case say 10⁻⁶. It should be noted that <u>SURFACE</u> will not accept "0" as an input here, but a very small value will accomplish the same task.

The mass flow rate for the larger droplets is not explicitly given in any of the outputs of the previous programs. This value may be determined in the following manner:

From the appropriate page of KINCON kinetics output a value for the gas density at the last S/RC position can be found, e.g., 2.44E-05 lb/ft³. The value of density can be extrapolated from this last printout position to the surface point by using a "inverse square of the distance" procedure, i.e., using the radius vector distance at the end of the MOC calculation in comparison to the radius vector in the vicinity of the surface segment, a value for the gas density at the satellite of 8.39E-08 lb/ft³ is calculated.

Referring back to the limiting particle <u>TD2P</u> output at the closest value of R/RC and Z/RC for the S/RC value used above gives the value for SDK/DG, the ratio of total particle density to gas density, 3.076. The product this value, the extrapolated gas density, and VPK, the particle velocity, give the estimation of the flow rate needed for input at PLMCHR(8), i.e., 4.38E-04 lb/ft²-sec.

PLMCHR(9) through PLMCHR(31)

At stated earlier, the chemistry has frozen and, therefore, the mass fractions given in the later pages of output from <u>KINCON</u> remain unchanged and are assumed to continue to remain so until the surface point is reached. If any of the species called for in the PLMCHR list are not here, they are input in the proper array location as 0.0 (if the card is simply left blank that will also suffice in the case of EDATA). When a species on the PLMCHR list does exist, the mass fraction is read from the <u>KINCON</u> output and put into the proper position of the data card. In this case all that is matched is water and carbon, i.e., PLMCHR(15) = 0.2803 and PLMCHR(25) = 4.19E-12.

PLMCHR(34) through PLMCHR(53)

If there were any small drops present, they would probably be condensed water. Looking at the table of "Physico-Chemical Properties of Plume Deposits" inthe output of <u>SURFACE</u> it is seen that the Code No. for water is "7". This value is input in PLMCHR(34). The mass percentage of water is entered into PLMCHR(35), in this case 100.0. Since no other items make up the composition of the small drops, PLMCHR(36) through PLMCHR(53) are set to 0.0 or let blank.

PLMCHR(54) through PLMCHR(73)

This portion of the data array is the same as the previous, only for the large drops. In this example only the fuel drops are considered, and reference to the <u>SURFACE</u> output gives a code No. of "3" for MMH. This is used as the input for PLMCHR(54) and PLMCHR(55) is set to 100.0. All other portions of this segment of the array are set to zero or left blank.

PLMCHR(84) through PLMCHR(86)

These three inputs describe the plume gas phase at the surface of the vehicle. SURFACE uses this information to determine any additional heat transfer to the surface of the satellite due to latent heat release upon condensation. In the case, such as this example, when the rocket motor is so relatively far from the surface, this additional heat release is very small. No significant error is introduced, accordingly, if the temperature, PLMCHR(84) for the plume is taken to be the last value given by KINCON, and the pressure and density, PLMCHR(85) and (86) are extrapolated from the last output of KINCON to the point on the surface via an "inverse square of the distance" method, i.e., PLMCHR(84) = 890°R, PLMCHR(85) = 4.69 E-05 psia, and PLMCHR(86) = 8.39 E-08 lb/ft³.

PLMCHR(87)

This input is not actually needed as <u>SURFACE</u> will calculate the viscosity. If this item is left blank, <u>SURFACE</u> makes the calculation. If a non-zero value is input, the program uses this figure rather than calculate one.

At this point the file EDATA is rewritten, as discussed in Appendix D. At the present time the portions of the program SURFACE are still being checked out, and consequently, this subprogram is not in a fully production status.

SECTION VI

EXPERIMENTAL DETERMINATION OF GAS DEPOSITION RATE ON CRYOGENICALLY COOLED SURFACES

NOMENC LATURE

Symbol	Definition	Units
At	Nozzle throat area	ft^2
c_{D}	Nozzle discharge coefficient	
Сp	Specific heat at constant pressure	ft ² /sec ² -°R
$\Delta \mathbf{F}$	QCM beat frequency	Hz
h	Axial distance between nozzle and sensor	in
K	QCM constant	gm/cm ² -Hz
k	Hypersonic similarity parameter (Eq. 2)	
M	Mach number	
m	Mass flow rate	lb _m /sec
Δm	Mass deposited	gm/cm ²
P _c .	Nozzle stagnation pressure	mmHg or lb _f /ft ²
R	Gas constant	ft ² /sec ² -°R
$Re_{ extbf{D}}$	Throat Reynolds number	
r _e	Nozzle exit radius	in
T	Temperature	° R
Δt	Time increment	sec
77	Velocity	ft/sec
v lim	Limiting gas velocity (Eq. 3)	ft/sec
α	Sticking coefficient (deposited mass flux/incident mass flux)	• • •
Y	Ratio of specific heats	
θ	Flow angle	deg
ρ	Gas density	lb _m /ft ³
ρV	Incident mass flux	lb _m /ft ² -sec or gm/cm ² -sec
	Subscripts	
С	Nozzle stagnation condition	
e	Nozzle exit condition	

1. OBJECTIVE AND APPROACH

The objective of this experimental program was to measure the steady-state deposition rates of various gases on cryogenically cooled surfaces. These data will eventually be used to predict contamination effects on spacecraft borne optical devices and thermal control surfaces.

It was originally planned to perform these tests as functions of gas type, surface temperature, and prior surface deposit. The items actually covered consisted of variations in gas type (NH₃, CO₂, and H₂O) and surface material (gold and silver).

All testing was performed in a laboratory scale vacuum facility the key elements of which were: a gas supply system, sonic nozzle, LN₂ cooled shroud, measuring equipment, and a quartz crystal microbalance (QCM).

Each combination of gas type and surface materials was run as a function of incident mass flux (which was accomplished by varying the gas stagnation pressure). After data reduction the results were presented as the sticking coefficient (deposited mass flux/incident mass flux) versus incident mass flux. Other methods of data normalization are possible; e.g., sticking coefficient versus Knudsen number (based on gas properties at the surface); but these were not explored during the course of the present investigation.

2. EXPERIMENTAL FACILITY

The experimental facility consisted of four subassemblies:

- 1. Vacuum system.
- 2. Gas supply and feed system.
- 3. QCM electronics and sensor.
- 4. Measuring equipment.

Figure 13 is a schematic of the facility and indicates the functional and geometric relationship of he above subassemblies.

a. Vacuum System

The vacuum system consisted of a 14-in, -diameter glass belljar; a mechanical pump; and a 6-in,-diameter oil diffusion pump with an $\rm LN_2$ cold trap. A Veeco Type RG-31X unit was employed to monitor the belljar pressure via a Veeco Type RG-75 ionization gage and the diffusion pump foreline pressure using a Veeco Type DV-1M thermocouple gage.

Feedthroughs were provided in the belljar baseplate to accommodate the shroud LN₂ lines, electrical power to the sensor, signals from the sensor, nozzle heater, and thermocouple leads.

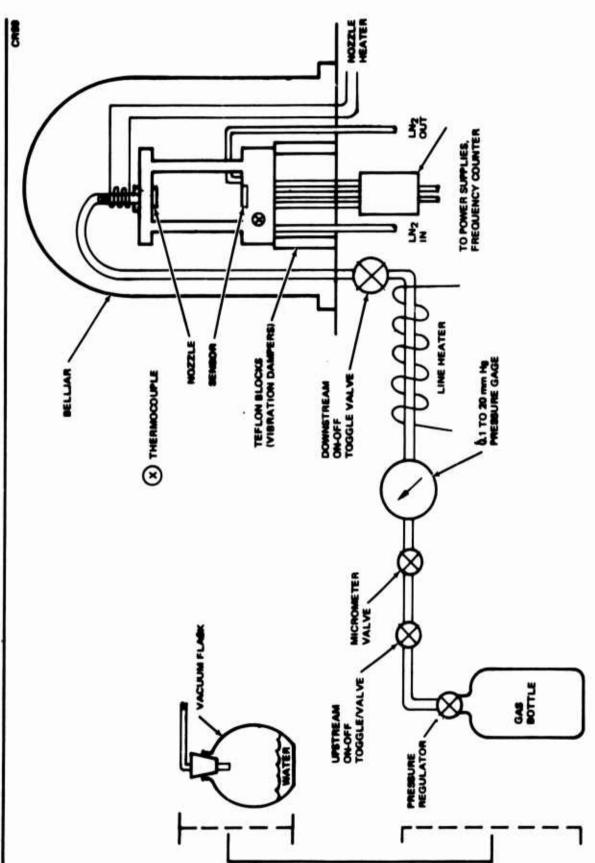


Figure 13. Facility Schematic

O

b. Gas Supply and Feed System

This system schematic is also shown in Figure 13 and includes the LN₂ cooled shroud (onto which is mounted the sonic orifice and QCM sensor). The CO₂ and NH₃ gases were supplied from commercially available cylinders while the water vapor was drawn from a vacuum flask.

An electrical line heater, powered by a Variac, was wrapped over the 1/4-in. stainless steel lines from the 0.1 to 20 mm Hg absolute pressure gage to the belljar baseplate in order to avoid NH3 and H2O condensation in the line. In addition, a bead heater was wrapped around the line just upstream of the nozzle orifice fitting (again, powered by a Variac). The use of these heaters was necessitated by evidence of nozzle plugging by condensed NH3 during some of the earlier test runs.

Figure 14 presents a detailed cross section of the LN₂ cooled shroud. The nozzle orifice (Figure 15) was mounted in a Swagelock fitting and secured to the top of the shroud with two Teflon washers in order to provide some degree of thermal isolation. A copper-constantan thermocouple was inserted through the side of the feedline just upstream of the nozzle fitting such that the junction was within the gas stream. This reading was taken as the gas stagnation temperature because of the high contraction area ratio between the feedline and nozzle throat.

SENSING CRYSTAL

REFERENCE CRYSTAL

SENSOR CAP
(STAINLESS STEEL)

SENSOR SOCKET
(PLASTIC)

TO QCM

SELECTRONICS
(LEADS EPOXIED TO TEE)

Figure 14. Sketch of QCM Sensor/Shroud Installation

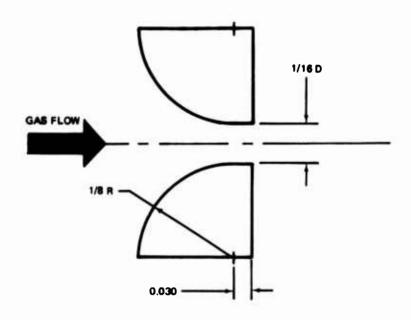


Figure 15. Sonic Nozzle Configuration

A second copper-constantan thermocouple was spot-welded to the outside of the shroud LN2 reservoir and was used to verify the shroud cooldown. Typically, this reading very quickly approached the LN2 boiling point at one atmosphere pressure and the use of this thermocouple became of secondary importance.

The QCM sensor was installed through the bottom of the shroud and was located at its forward edge by a smaller diameter lip within the brass shroud. The sensor was secured to this lip by an aluminum tee member which beared on the sensor plastic socket. This member was also a means of cooling the sensor by providing a conduction path from the sensor to the LN2 reservoir. Additionally, the six leads from the sensor to its remote electronic package were bonded to the aluminum tee by a high thermal conductance, high electrical resistance epoxy (Wakefield Delta Bond 152). This design feature "short circuited" the energy being conducted from the warmer electronics package to the sensor. The shroud assembly was then loosely covered with aluminum foil with the foil being secured by wrapping it around the LN2 lines. A narrow slit was made in the foil so that the sensor surface could be visually examined.

c. QCM Electronics and Sensor

The QCM system employed for this experimental program was manufactured by Faraday Laboratories, Inc., La Jolla, California, and consisted of an electronics package (Model 71-DE, Serial No. E8) and a series of plug-in crystal sensors.

The electronics was located on the test stand external to the $\rm LN_2$ vacuum system since its specification indicated a minimum allowable temperature of -50°C. Electrical power was furnished by a standard 28-vdc supply while power for the sensor heater was furnished by a variable dc supply.

The QCM sensor consisted of two 10-MHz crystals matched to the temperature coefficient (so as to minimize beat frequency shifts with crystal temperature). The outer or sensing crystal was exposed to the incident gas stream and could be supplied with a variety of coatings. The program described herein used gold or silver plated sensing crystals. In addition, both crystals could be heated via an aluminum undercoat in order to drive off prior deposits. A solid-state sensistor was supplied to measure the crystal temperature; however, the manufacturer did not furnish calibration information for these units in the temperature range between -40°C and -200°C. To remedy this situation a copper-constantan thermocouple was spot-welded to the ground pin of the sensor plug and, the output of this thermocouple was one of the primary readings recorded during a test.

The mass deposited on the sensing crystal causes a change in its natural frequency of oscillation which when it is mixed with that of the inner, or reference, crystal causes a "beat" frequency (ΔF , the difference between the reference and sensing crystal frequencies) to be generated. This beat frequency was recorded on a digital frequency counter and was then related to the mass deposited by:

$$\Delta m(gm/cm^2) = K \Delta F (Hz)$$

where K was either 4.1 x 10^{-9} or 3.5 x 10^{-9} gm/cm² -Hz depending on the sensor employed.

The manufacturer's specifications for the sensors indicated that they were capable of measuring deposits as low as 1.0 \times 10⁻⁹ gm/cm² to as high as 1.0 \times 10⁻³ gm/cm².

d. Measuring Equipment

The voltage output of the copper-constantan thermocouples (in millivolts) located on the gas supply line, shroud, and sensor were measured on two Hewlett Packard Model 3440A digital voltmeters one of which employed

a H-P model 3433A high gain/auto range plug-in and the other with a H-P model 3444A dc multifunction plug-in. The output of the sensistor (in volts) was displayed on the DVM with the auto range plug-in. These units have a lowest scale of 100 millivolts which yielded three-digit representation of the thermocouple voltages.

A H-P Model 5245L digital electronic counter was used to display the sensor beat frequency and was set on its most sensitive scale of 0.1 vrms. The input to the counter was split to also drive a H-P Type 561B oscilloscope with a Type 3A72 dual-trace amplifier. The oscilloscope was was used primarily during the initial facility setup to check for ground loops; subsequent usage was restricted to measuring the sensor output level.

3. TEST PROCEDURE

At the beginning of each day certain preparatory steps were taken prior to the initiation of the deposition runs. These included: evacuating the belljar, filling the thermocouple junction bath with an ice-water mixture, energizing the line and nozzle heaters (if NH₃ or H₂O was to be run), trimming the digital voltmeters against their internal checks, and filling the shroud with LN₂ to cool the sensor. The gas supply system was purged with dry GN₂ if a different gas was to be run.

The sensor cooldown period typically required 1 hour as evidenced by stabilizing of the sensistor and sensor thermocouple readings. At this point the micrometer valve was opened and adjusted to give the desired feed pressure and the clock was started. The beat frequency, gas stagnation thermocouple, and sensor thermocouple readings were manually recorded at time intervals which varied with the feed pressure; e.g., every 1 minute at 0.5 mm Hg and every 10 seconds at 4.0 mm Hg. The sensistor output was occasionally recorded and adjustments were made to the micrometer valve, if necessary, to maintain a constant feed pressure.

The run was terminated when the beat frequency reached 250 to 300 kHz (which corresponded to the maximum sensor loading of 1.0 x 10^{-3} gm/cm²) at which time the micrometer valve was adjusted to the next feed pressure. The upstream toggle valve was closed cutting off the gas flow and the sensor heater was energized to drive off the deposited material and return the sensor beat frequency to its nominal value. The next run was initiated by opening the upstream toggle valve and starting the clock.

The use of an automatic data recording system would be required in order to extend the gas feed pressure (and gas flowrate) to higher levels since it was not possible to manually record the data for time intervals less than 10 seconds.

It is to be noted that the sensor was not removed from the belljar once it was installed even though some change in the sensor nominal beat frequency was noted over a period of time. Earlier attempts to clean the gold plated sensors with a solvent usually resulted in removal of some of the gold. Two of these damaged units were returned to the manufacturer who successfully replated one of them with gold.

4. DATA REDUCTION

The reduction of all test data involved the following steps:

- 1. Determining the steady-state mass deposition flux.
- 2. Estimating the incident mass flux at the sensor location.
- 3. Calculating the sticking coefficient from the ratio of steps 1 and 2.

The final reduced data consisted of graphs showing the sticking coefficient versus incident mass flux for each of the gases and surfaces employed.

a. Steady-State Mass Deposition Flux

As previously mentioned the sensor beat frequency, ΔF , was recorded as a function of time, t. The first step in the data reduction process was to plot ΔF versus t and establish when this function achieved a constant slope. A typical result is presented in Figure 15 and indicates that approximately 5.5 minutes was required to establish a steady-state deposition condition for this particular run.

The corresponding steady-state mass deposition flux, $\Delta m/\Delta t$, was then found from:

$$\frac{\Delta m}{\Delta t}$$
 (gm/cm²-sec) = K $\frac{\Delta F (Hz)}{\Delta t (sec)}$

where K is a theoretical value supplied by the manufacturer and was either 4.1 x 10^{-9} pt 3.5 x 10^{-9} depending on the particular sensor.

b. Inviscid, Continuum, Incident Mass Flux

The inviscid, continuum, incident mass flux, ρV , at the sensor location was calculated using the approximation by Roberts (Reference 19).

In this formulation the gas density is given by:

$$\rho = \rho_{e} \frac{k}{2} \left(\frac{r_{e}}{h} \right)^{2} (\cos \theta)^{k}$$
 (1)

$$k = Y(Y-1) M_e^2$$
 (2)

The opening in the cap over the sensing crystal was 3/8 in. which indicated that this crystal subtended an angle of ± 3.7 deg; therefore, $(\cos \theta)^k$ was taken as unity (since $k \approx 0.45$).

The gas density at the nozzle exit was found from the measured feed pressure, stagnation temperature (T_c) , gas constant (R), and isentropic expansion to an M_e zone. The value of $Y = C_p/(C_p-R)$ was found from Reference 20 as a function of stagnation temperature. The gas velocity, ρV , at the sensor location was taken as the limiting gas velocity, i. e.:

$$V = V_{\lim} = \sqrt{2 \, YRT_{c}/(\gamma - 1)}$$
 (3)

The inviscid, continuum, incident mass flux, ρV , was found by multiplying Equations (1) and (3).

c. Pitot Pressure Measurements

A pitot pressure probe was fabricated from 0.064 in. Od by 0.024 in. Id by 10 in. long stainless steel with a square lip and the pressure was measured on a Type 77 MKS Baratron Pressure Meter utilizing a secondary vacuum reference source.

The measured pitot pressure was divided by the feed (stagnation) pressure and with the normal shock total pressure relationship was used to calculate the free-stream Mach number. Results of these calculations for ambient temperature air, as shown in Figure 16, indicated that the free-stream Mach number was approximately 10.6 (over a feed pressure range of 0.5 to 3.0 mm Hg) while the Mach number found from the plume approximation (an extension of Equation 1) was 19.0. This difference is due to the rarefaction of the gas during the expansion process which "freezes" the velocity, static temperature, and Mach number. The error thus incurred amounts to overpredicting the air velocity by 2.2 percent when Equation (3) is employed, i.e., V(M=19)/V(M=10.6)=1.022.

The inverse square dependency of density with distance (Equation 1) is still valid at the sensor location since it is a consequence of satisfying the conservation of mass. To more clearly illustrate this point Equation 1 may be rewritten (with $\cos \infty = 1$) as:

$$\rho V = \rho_{e} \frac{k}{2} \left(\frac{r_{e}}{h}\right)^{2} \sqrt{2YR(T_{c}-T)/(Y-1)}$$
 (4)

At some axial location, h_f , the static temperature (T) and velocity (V) "freeze" and the free-stream mass flux, ρ V, still varies as (h)-2. If it is assumed that the expansion suddenly freezes; i.e., the expansion follows continuum theory up to H_f and then T (and V) becomes constant; then the value of h_f can be calculated. For the air results cited above it was found that $h_f = 0.68$ in. compared to the nozzle/probe separation distance of 2.85 in.

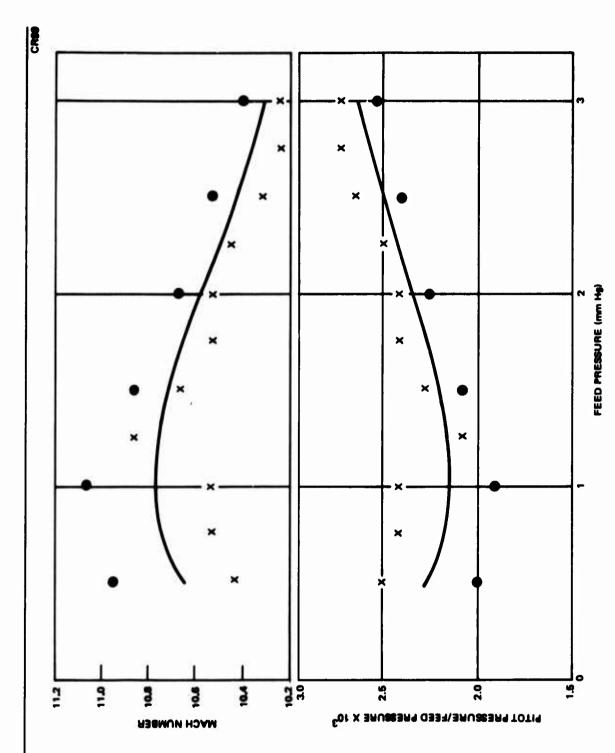


Figure 16. Pitot Pressure Results for Ambient Temperature Air

(3) Multiplying the result from step 2 by the nozzle discharge coefficient corresponding to a square-edged orifice to arrive at the incident mass flux.**

The deposited mass flux was then divided by the incident mass flux to yield the sticking coefficient. The last step in the data reduction process involved performing least squares polynomial (cubic) curve fits to all test data. No attempt was made to preselect those data points to be used in this process; i. e., all data points were employed.

d. Sensor Surface Temperature

At the conclusion of the test program a small copper-constantan thermocouple was bonded directly to the sensing crystal of an inactive sensor. The shroud was then cooled with LN₂ and the reading of this thermocouple was compared to that attached to pin C of the sensor plug after a steady-state condition had been attained. It was found that the crystal was 5°C warmer than the plug thermocouple. (The latter thermocouple was thermally closer to the LN₂ reservoir than the former.) Therefore, all quoted surface temperatures were found by adding 5°C to the plug thermocouple reading.

5. REDUCED DATA

The finalized test data are presented for steady-state deposition of NH₃, CO₂, and H₂O on both gold and silver surfaces. Attempts were made to deposit O₂ and CH₄ in a flowing system but these were unsuccessful. Also presented are the results of a test performed to measure the NH₃ evaporation rate in a vacuum environment.

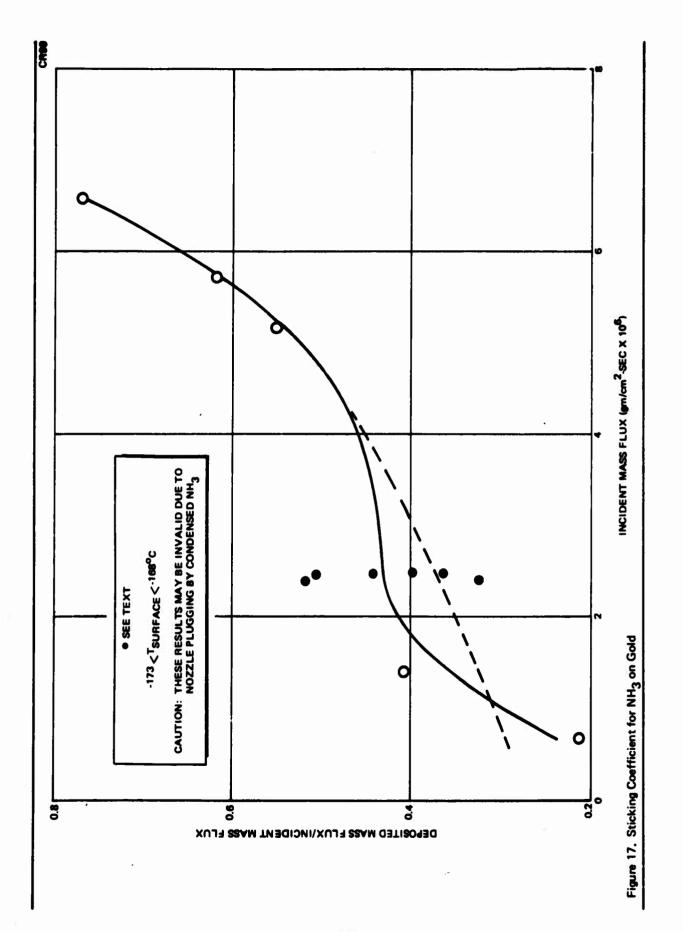
a. Ammonia (NH₃) Results

The HN₃ deposition rate on a gold plated QCM sensor was the first of the test series conducted and was subject to more data scatter than the subsequent tests. These data are presented in Figure 17** and depict the sticking coefficient (defined as the ratio of the deposited mass flux to the incident mass flux) as a function of the incident mass flux.

The tests conducted at an incident mass flux of approximately 2.5 x 10^{-6} gm/cm²-sec were the only ones which exhibited a <u>reduction</u> in the sticking coefficient with time, i.e., the deposition rate was larger at the beginning of the run than later on during the run. (A similar occurrence was noted in Figure 6 of Reference 23 for N_2 but no explanation was offered therein.)

^{*}The use of the measured C_D 's, in preference to those from Reference 22 would result in sticking coefficients on the order of 2-4.

^{**}Surface temperature range noted on Figure 17 (and subsequent figures) refers to the minimum and maximum pensor temperatures recorded for all the data points. The variation in sensor temperature during the course of any one run was usually on the order of 2-3° C.



In point of time the NH_3 on gold results were the first ones obtained and these tests were performed prior to the installation of heaters on the gas supply line and nozzle. (These heaters were installed commencing with the H_2O on gold series to prevent condensation in the lines since saturated vapor was being withdrawn from the liquid flask.) Thus, there is some reason to suspect the NH_3 on gold results due to partial line plugging (especially at the higher $\rho v's$).

Corresponding results for NH₃ deposition rates on silver are shown in Figure 18. It is to be noted that the sticking coefficient was monotonically increasing with time to its steady-state value at each data point. Figure 19 compares the NH₃ sticking coefficient for both the gold and silver plated sensors. Both results agree up to a ρv of 2 x 10^{-6} gm/cm²-sec with the gold results then exhibiting a lower sticking coefficient and different trend. (This difference is attributed to nozzle plugging by condensed NH₃ at the higher incident mass fluxes.)

b. Carbon Dioxide (CO₂) Results

The gold and silver results of sticking coefficient versus incident mass flux are presented in Figures 20 and 21, respectively. Both sets of data show very similar trends: a nearly linear increase of α with ρV followed by an asymptotic approach to an α of unity. This is more clearly depicted in Figure 22 which shows that $\alpha \approx 1$ for ρV between 10 to

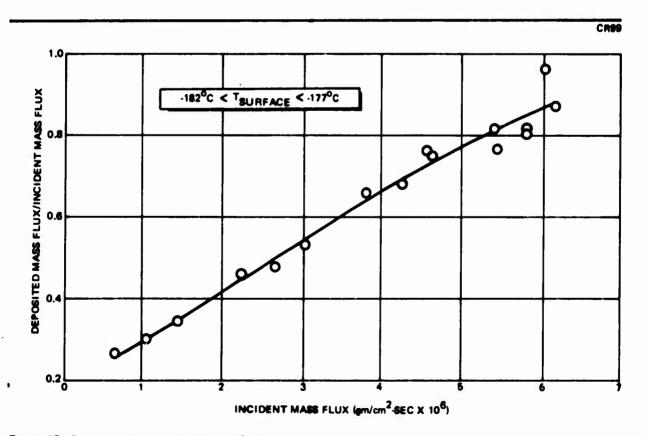


Figure 18. Sticking Coefficient for NH₃ on Silver

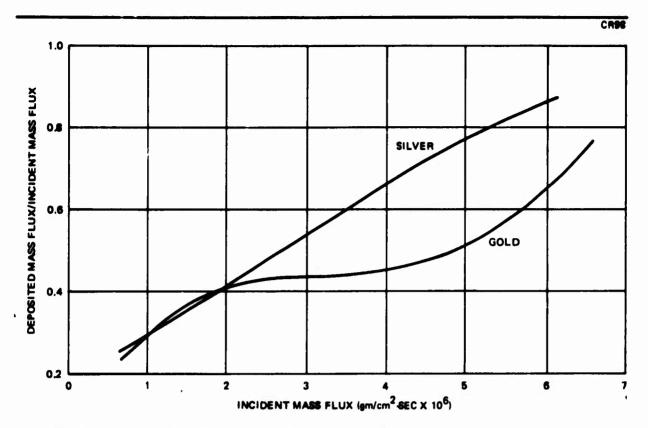
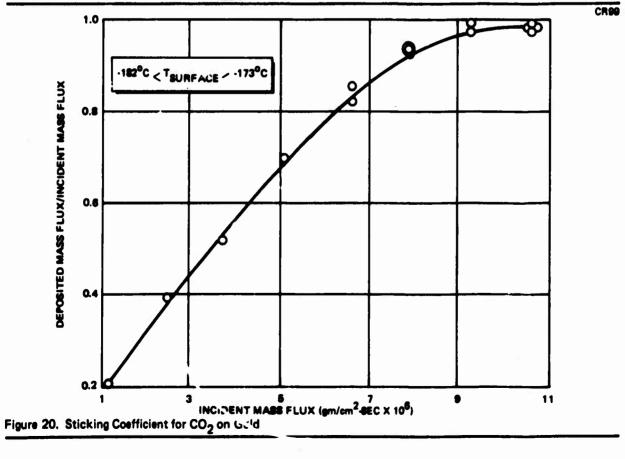
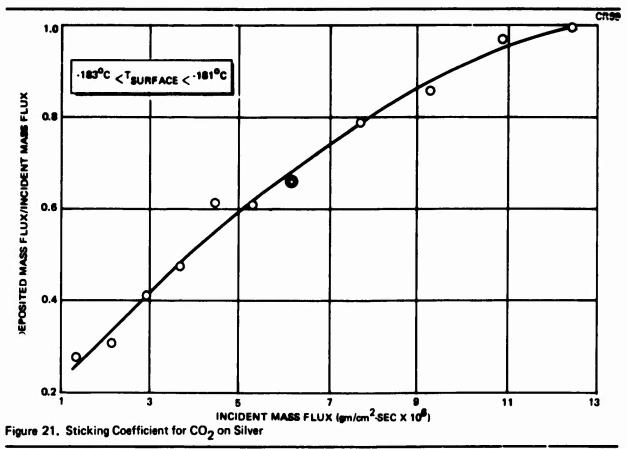
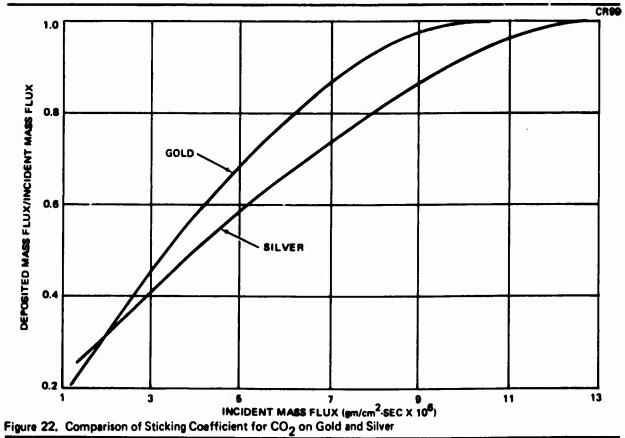


Figure 19. Comparison of Sticking Coefficient for NH₃ on Gold and Silver







12.5 x 10^{-6} gm/cm²-sec. In addition, the differences in α between the gold and silver plated sensors were judged to be relatively minor, i.e., approximately a 19 percent difference at most. This observation, in conjunction with the close similarity in surface temperatures, indicates that there is essentially no difference in the steady-state α produced by the surface materials. This is supported by calculations which showed that at the sensor's upper measurement limit (1 x 10^{-3} gm/cm²) there are approximately 2.5 x 10^4 monolayers of CO₂ deposited on the surface; i.e., the incident gas does not "see" the sensor surface.

The use of a theoretical clean surface would result in deposition rates which were dependent on the surface material due to differences in the potential energy function between the gas and the surfaces. However, it was not feasible to achieve this ideal condition nor accurately measure the deposition rate with the experimental setup, e.g., the buildup in feed pressure (incident flux) was too slow to attribute a measured deposition rate to a corresponding incident flux.

c. Water Vapor (H2O) Results

The H₂O results for gold and silver surfaces are contained in Figures 23 and 24, respectively, while a cross-plot is shown in Figure 25.

The use of all the silver data points in the curve fit resulted in a dropoff in α at the higher values of V. A modified fit, as shown by the

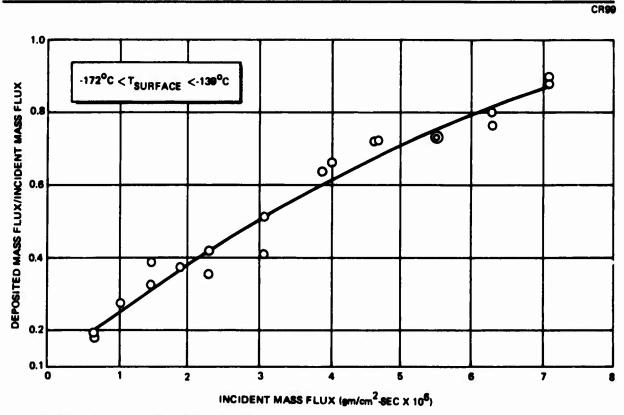


Figure 23. Sticking Coefficient for H₂O on Gold

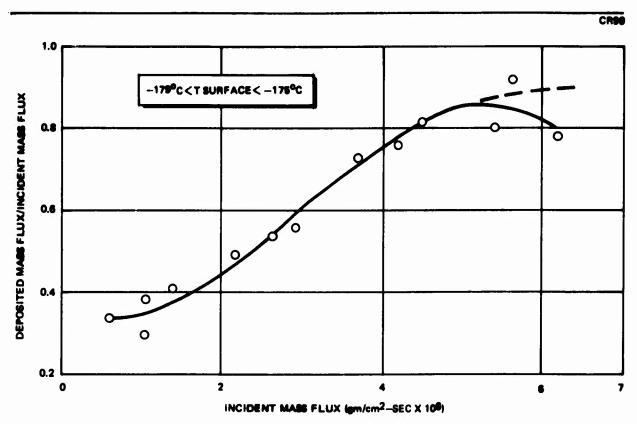


Figure 24. Sticking Coefficient for H₂O on Silver

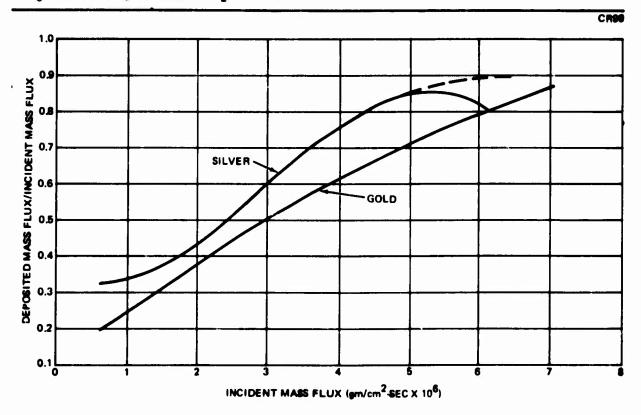


Figure 25. Comparison of Sticking Coefficient for H₂O on Gold and Silver

dashed lines in Figures 24 and 25, was employed in this area and the resulting comparison (Figure 25) indicates the following:

- 1. The gold results were obtained over a much larger range in surface temperature.
- 2. The silver α 's for the H₂O are higher than the gold values (as opposed to the opposite trend for CO₂ shown in Figure 22) although they both appear to be approaching a common limit. This is probably due to the higher evaporation rate for the gold results which would produce lower net deposition rates, i.e., lower values of α .
- 3. The spread between the results is greater than those obtained for CO_2 with the maximum difference being 24 percent for $\rho V > 2 \times 10^{-6}$ gm/cm² -sec.

For engineering level contamination studies the H₂O results, especially item 3 above, suggest that there is essentially no difference between steady-state deposition on gold and silver surfaces.

d. Composite Summary of Sticking Coefficients on Gold and Silver

The reduced data for NH₃, CO₂, and H₂O on the gold plated sensor are shown in Figure 26 with the corresponding silver information presented

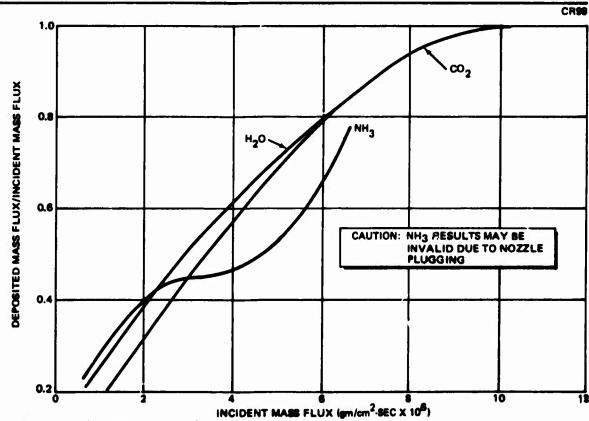


Figure 26. Sticking Coefficient for NH2, CO2, and H2O on Gold

in Figure 27. The α for H_2O on gold is seen to approach, and then coincide with, the CO_2 results. It is to be noted that the surface temperature was significantly higher, 10 to 34°C, for the H_2O on gold than for the CO_2 test series. This higher temperature will cause a greater evaporation rate and a corresponding lower net deposition rate.

By comparison the NH₃, CO₂, and H₂O deposition results with a silver plated sensor (Figure 27) appear to be the most consistent set of data gathered: there were no anomalies concerning a decrease in α with time (as was the case with NH₃ on gold) since line and nozzle heaters were installed for all runs to prevent plugging; and, the maximum difference in surface temperature under any condition was 8° C. The α for CO₂ is seen to reach unity at an incident mass flux of \approx 12.5 x 10^{-6} gm/cm²-sec and it is apparent that both H₂O and NH₃ will achieve an α of unity at even lower values of V. (The more pronounced curvature of H₂O on silver at low ρ V is due to the nozzle discharge coefficient effect which was more significant for these runs.)

e. Vacuum Evaporation

A preliminary vacuum evaporation experiment was performed wherein various amounts of heater current were applied to a gold sensor containing a prior NH₃ deposit. (The sensor manufacturer's specification indicated a 400 ma nominal heater current.) This test was planned to determine the evaporation rate as a function of surface temperature.

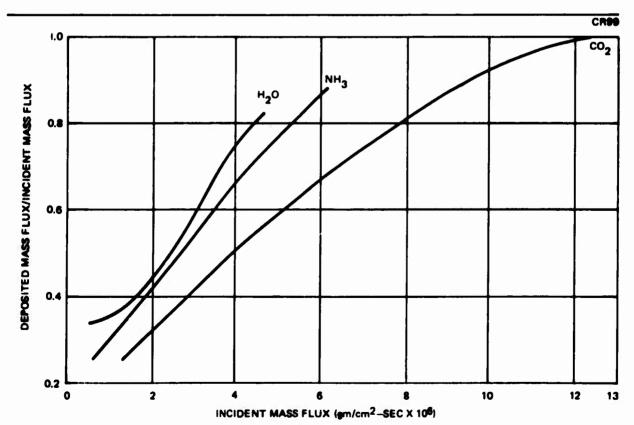


Figure 27. Sticking Coefficient for NH₃, CO₂, and H₂O on Silver

The results of this experiment were as follows:

- (1) With 100 ma heater current the sensor crystal stabilized at -171°C with a belljar pressure of 4.4 x 10⁻⁶ mm Hg. Additional evaporation was occurring at a rate of 7, 823 x 10⁻⁸ gm/cm²-sec over an 18 minute time period.
- (2) With 200 ma heater current and the belljar at 4.0×10^{-6} mm Hg the sensor crystal stabilized at -169.5°C. An evaporation rate of 4.106×10^{-8} gm/cm²-sec was noted over a 92 minute time period.
- (3) With 300 ma heater current and the belljar at 3.6 x 10^{-6} mm Hg the sensor crystal stabilized at -167.5°C and the corresponding evaporation rate was 1.002 x 10^{-6} gm/cm²-sec averaged over a 13 minute time period.

The additional evaporation (item 1) was probably due to residual NH3 gas in the belljar which condensed on the sensor even though the heater was partially activated. The other results (items 2 and 3) indicate that the evaporation rate is strongly dependent upon the surface temperature; i. e., a 2.0°C increase in surface temperature produced a factor of 24.4 increase in the evaporation rate.

6. INTERPRETATION OF RESULTS

All the data generated during the present investigation exhibited the common trend of increasing sticking coefficient (α) with increasing incident mass flux (ρ V). In addition, each gas was run at essentially the same kinetic energy level as evidenced by the relatively small variation in stagnation temperature.

The deposition process involves the collisional transfer of energy from the incident molecule to the surface; and if this transfer is of sufficient magnitude or if the incident energy is low enough, then the molecule is "trapped" by the surface's potential energy field. This is clearly shown in Figure 6 of Reference 24 which presents the fraction trapped and thermal accommodation coefficient for argon on nickel as a function of incident energy. This particular study was theoretical in nature and considered only a single molecular impacting on a "clean" surface. Above a certain incident energy level there was no trapping predicted, i.e., $\alpha = 0$, while these results also shown that the thermal (energy) accommodation coefficient is $\approx 1/2$ when the sticking coefficient is 0, i.e., the molecule is partially accommodated to the surface energy without being trapped.

The present study involved vast numbers of incident molecules which possessed distributions of velocity (energy) about the mean flow velocity (energy). The results presented herein are qualitatively interpreted as follows:

- a. Low values of incident mass flux, ρV , were obtained by low values of density, ρ , which implies fewer gas molecules in the path of the incident molecule.
- b. Therefore, the incident molecule suffers few collisions and impacts the surface with small attenuation in its velocity (energy).
- c. The reflected molecule also experiences few collisions after rebounding from the surface and has a low probability of being redirected back toward the surface for a second encounter.
- d. Some of the incident molecules should be trapped by the surface at low PV due to the distribution of velocity (energy) about the mean; i. e., statistically, a fraction of the incident molecules will have energies low enough to be trapped.
- e. As the incident mass flux increases, through increases in ρ , a given molecule will undergo more collisions with those molecules which have already impacted the surface and will arrive at the surface with a reduced energy content, and the probability of trapping will increase.
- f. Those molecules which have not been trapped during their first impact with the surface may experience collisions with molecules near the surface such that they are directed back toward the surface for an additional encounter(s), thereby increasing the trapping probability.
- g. In the continuum limit the molecule will reach the surface with zero kinetic energy along the stagnation streamline. The deposition should then depend on whether the heat transfer rate to the surface is sufficiently high to permit extracting the sensible and latent energy from the gas.

The matter of particle density on surface effects has received attention in the area of surface damage due to high speed particles (Reference 25). In this instance increasing the incident particle flux did not produce a proportional increase in surface damage. Indeed, the relative (per particle) target damage was observed to rapidly diminish with increasing particle flux. This effect was attributed to shielding of the surface by particles which were already in the shock layer thereby causing significant energy attenuation of incoming particles and reduced damage per particle.

The above results have a direct parallel with the deposition experiment in that at high incident mass flux the incoming particle energy is attenuated to the point where they are more effectively trapped by the surface.

7. RECOMMENDATIONS FOR THE FUTURE EFFORT

Recommendations are contained, herein, for future effort in two areas:

- a. Improvements to experimental system hardware.
- b. Additional tests.

The recommended system hardware changes are based upon experience gained during the course of the present investigation and are suggested to improve the ease of data gathering, data reduction, and general flexibility of the system.

The additional tests proposed are those which would extend the range of the key parameters and thereby increase their applicability to full-scale systems.

a. System Hardware Changes

The sensor surface temperature is one of the key parameters to any deposition study and, for the present investigation, was difficult to control with any degree of precision. There was a tendency for the sensor temperature to increase with time during the course of a run due to heat transfer from the gas to the sensor. This drift can be corrected by redesigning the sensor/reservoir interface so that the sensor will be thermally closer to the reservoir.

A thermostatically controlled sensor electrical heater and/or controlled shroud LN₂ flow rate must be included in the design so that preselected sensor temperatures can be achieved without the necessity of opening the belljar. Attempts were made during the present investigation to increase the sensor temperature by, for example, inserting Teflon washers between the sensor cap and the reservoir, and by wrapping a small heater around the sensor cap. These attempts were very time consuming (requiring 3 hours just for a cooldown and warmup cycle) and produced either an insignificant temperature change or too large a change.

The incident mass flux, ρV , was varied during the present investigation by changing the stagnation pressure (and stagnation density) at a fixed nozzle/sensor separation distance. Variations in ρV could also be accomplished by changing the separation distance via a remotely controlled electric motor. The data thus generated would serve as a cross-check on the results presently available and would possibly allow the sensor to span the range from continuum to rarefied fluid mechanics.

A fast opening valve, externally controlled, must be placed as close to the nozzle as possible if it is desired to follow the transient portion of the deposition process. The present configuration had manually operated toggle valves outside the belljar and may have been responsible for the extended time period required to achieve a steady-state deposition rate (see Figure 16).

The need for an automatic data recording system becomes critical if results are to be obtained at higher values of incident mass flux. The test equipment employed for this investigation have BCD coded terminals on the chassis which can be connected directly to a recording system. If possible, the output of the frequency counter should be displayed on a stripchart so that

a quick check can be made of the run, e.g., the run could be terminated when the mass deposition flux achieved a constant valve as evidenced by a linear variation of ΔF versus time. These plots could also be used as the first step in the data reduction process (as described in Subsection 4. a).

Higher rate vacuum pumps would also have to be used if mixtures of condensible/noncondensible gases are to be tested (Subsection 7. b).

b. Additional Tests

The primary additional test that is suggested involves measuring the deposition characteristics of a condensible/noncondensible gas mixture. This data will supply information as to whether the deposition rate of the condensible gas is a function of its partial density or of the total density. Commercially available mixtures of, for example, carbon dioxide in argon could be used at various CO₂ concentrations.

As mentioned in Subsection 7. a the attempts to systematically vary the sensor surface temperature were unsuccessful. However, the proposed system modifications should make it possible to more closely regulate this temperature so that gas deposition characteristics can be measured as a function of surface temperature. It is recommended that CO_2 be used for the initial series of tests and, based on these results, followed by NH_3 and H_2O .

The molecular weight ratios employed in the present investigation were NH₃: CO_2 : H_2O : 1.0:2.584:1.058. It is suggested that a higher molecular weight gas such as SF₆ be tested to extend this ratio to 8.577 in order to establish if the deposition rates can be correlated in terms of molecular weight and/or inter-molecular force constants.

The effect of incident particle energy on deposition rate should be investigated. As discussed in Section VI, this is another key parameter in that it can indicate if a particle can be trapped by the surface's potential energy field. This will be somewhat difficult to simulate in the laboratory since high energies require high temperatures and the experimental facility may not be capable of achieving these conditions without a major redesign.

Section VII CONCLUSIONS AND RECOMMENDATIONS

A computer program has been developed to predict the sensitivity to engine operating parameters and impingement geometry for the production and transport of bipropellant engine contaminants. An associated program to analyze the effect of plume impingement on sensitive thermal and optical surfaces in terms of changes in surface properties has also been completed. The contaminant production and contaminant transport models and subprograms have been checked out and seem to be working correctly, although verification with experimental data has not been attempted due to a lack of detailed experimental data relating engine operation to contaminant effects on surfaces.

Based on the analysis of a single engine, the Marquardt R-6C MMH/NTO 5-lb thrustor, it appears that significant amounts of unburned fuel and oxidizer will be ejected from the engine during the transient portion of the pulse and that this contaminant may cause damage to sensitive thermal and optical surfaces if they are impinged upon by the central core of the plume, or if they are spattered with wall-film material blown off at the nozzle lip.

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Appendix A

TCC

TRANSIENT COMBUSTION CHAMBER DYNAMICS COMPUTER PROGRAM

A Bipropellant Contaminant Production Model

Program Number H607

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LIST OF ABBREVIATIONS AND SYMBOLS

Α	Area, Acceleration, Molecular collision frequency
a	Acoustic velocity
В	Coefficient in vapor pressure equation
С	Specific heat, Coefficient
D	Diameter
E	Activation Energy, Accommodation Coefficient
F	Stoichiometry (mass fraction which is fuel)
f	Fanning friction factor
G	Entrainment parameter
Н	Enthalpy
(I)	Subscript indicating a particular member of an array
K	Drop size distribution coefficient, Thermal conductivity
K¹	Droplet burning rate coefficient
L	Length
M	Mass
N	Dimensionless modulus
P	Pressure
[P]	Parachor
Q	Heat of reaction
q	Volumetric flow rate, Heat flow rate, Quenching distance
R	Universal gas constant, Coefficient for resistance to flow
T	Temperature
t	Time
U	Chamber volume, Gas velocity
v	Velocity, Any time-dependent variable, Chamber volume
Wm	Molecular weight
x	Mass fraction which flashes, Axial location
Y	Radial location
β	Resultant stream angle
Y	Ratio of specific heats
Δ	Difference
•	Expansion area ratio

δ Thickness of a liquid film

μ Viscosity

P Density

σ Surface tension

Ratio of Circumference to diameter of a circle

SUBSCRIPTS

c Chamber, Critical point

cond. Condensed phase d Drag, Difference

e External, Effusion correction

F Fuel, Thrust coefficient

film Liquid film

flash Flash threshold

fuel Fuel g Gas gas Gas

i Droplet group identifier, Chamber axial segment identifier

inj Injection conditions

j Jet conditions

L At axial location L

1 Liquid

m Wall film material

mn Droplet mean $D_{mn} = \left[\frac{\sum Ni \ Di^{m}}{\sum Ni \ Di^{n}}\right]^{\frac{1}{m-n}}$

n Injection group identifier

Nu Nusselt number

O Oxidizer, Orifice, Stagnant

Oxid Oxidizer

P Propellant p, At constant pressure

R Resultant after impingement

r Reduced variable (ratio to critical value), ripple correction

rel. Relative

Re Reynolds number

S Surface

s Spin-derived

Sat Saturation conditions

Stream Stream conditions

T Throat, Total of fuel plus oxidizer, Tangential direction,

Temperature correction

t Tank

V Vapor

Vap. Vapor

w Wall

We Weber number

x Component in the x direction

y Component in the y direction

Appendix A

TCC

TRANSIENT COMBUSTION CHAMBER DYNAMICS COMPUTER PROGRAM

A Bipropellant Contaminant Production Model

A. 1 INTRODUCTION

a. The TCC Program

The computer program described in this appendix is a subprogram to the Plume Contamination Effects Prediction Computer Program, CONTAM, and performs the time-varying analysis of the chemical and physical processes occurring in the feed system, injector, combustion chamber, and nozzle-throat inlet of a bipropellant rocket-engine system operating under unsteady conditions. As a subprogram to CONTAM, the TCC program is the first link in the analysis of contamination effects, providing information about production of contaminants in the combustion chamber and the dynamic and thermodynamic state of combustion gases entering the nozzle throat. Unburned propellant droplet distributions and liquid wall film flow are computed for the entire transient pulse as well as gas-phase properties. TCC may also be used as an independent computer program on any third-generation computer with a core exceeding 104,000(8) words and a Fortran IV processor.

Development of the <u>TCC</u> program has been supported by MDAC Independent Research and Development over a period of approximately 5 years with additional modification supported by the current Air Force Rocket Propulsion Laboratory study. References A-1 through A-6 discuss the detailed modeling of the combustion processes included in the basic <u>TCC</u> program. Reference A-7 describes the development of the <u>TCC</u> program at the end of 1971, relates it to the earlier work in combustion chamber modeling, and gives an illustration of its use in doing a parametric variation study on a small rocket engine (the Marquardt R6-C). Reference A-8 illustrates the application of the <u>TCC</u> program to perform a detailed rocket engine analysis related to contaminant production and its potential effect upon a satellite. As far as is practical, this appendix will cover all aspects of the modeling and computer program.

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b. Development of the MDAC Transient Combustion Chamber Program

The combustion chamber processes important to large rocket engines have been modeled successfully for several years using the method introduced by R. J. Priem and subsequently refined by others (References A-9 through A-12). In these methods, numerical integration is used to calculate the trajectories and evaporation rates of the propellant droplets, assuming steady conditions in the chamber and one-, two-, or three-dimensional flow of the gas and droplets.

This same basic analysis has been extended to include the more complex events of importance in small pulse-modulated rocket engines. Small rocket engines differ from large engines in two important ways: (1) the proximity of the injection points to the chamber wall, the larger surface-to-volume ratio of the chamber, off-nominal injection and atomization during the transients, and the wide-spread use of film-cooling makes deposition of unburned propellant on the walls much more important in small engines; (2) small rocket engines are typically used to control vehicle attitude or for minor trajectory corrections, where the firing duration is typically as short as 10 to 100 msec, thus the transient behavior of a small engine system is at least as important as the steady-state behavior.

The <u>TCC</u> program was started at MDAC in 1967 under IRAD funding as a general method for solving transient problems in liquid rocket engines, and in particular as a basis for aiding in the development of very fast response control engines. It very quickly became evident that the original one-dimensional program had the capability to model low-frequency combustion instability with great precision, that it had a considerable potential for assisting in the analysis of hard starts, and that it offered a feasible approach to the study of contaminant production in pulse-mode rocket engines.

Since 1970 the AFRPL has supported the extension of the program to model contaminant production in liquid bipropellant rocket engines. Under this support, the droplet trajectories were made two-dimensional (three-dimensional in some experimental versions) to model deposition of propellant on the walls, hypergolic ignition and extinguishment were modeled, the wall processes have been modeled with progressively greater precision, and the postfiring dribble and vacuum evaporation have been modeled.

The intent has always been to use the simplest acceptable mathematical approximation for each combustion chamber process but to model all of the processes which are known to be important. Thus, the program is quite general in its capabilities.

The operational problems that can currently be analyzed include vacuum hypergolic ignition and the start transient, pulse-mode specific impulse, low-frequency system instability, production of contaminant material in the plume and as wall film, explosion pressures that can be obtained from chamber detonations, and engine response to off-nominal operating conditions. Good agreement with experiment has generally been obtained.

The present program, however, is still incomplete, in that there are some important processes which are not yet represented, i.e., axial and radial variation in gas stoichiometry, secondary atomization of droplets, pressure waves in the propellant feed systems, heat transfer to and through the walls, etc. It is also obvious that the present experimental correlations for stream breakup distance and primary atomization droplet size, which were all obtained with large orifice sizes, form a very questionable basis for design or prediction when extrapolated to the very small injection orifice sizes found in pulse-mode engines. The complexities of very reactive impingement or stream blow-apart have not been modeled.

c. Scope

The transient combustion chamber program uses numerical integration of the detailed system processes to calculate the events in a rocket engine system operating under unsteady conditions. The system which is modeled consists of a pressure-fed bipropellant feed system, an injector, a combustion chamber, and a nozzle (Figure A-1).

The initial flow of propellant is calculated as the valves open, the fluids accelerate through the resistive-inertial feedlines, and the injector fills. If the vapor pressure of the first injected propellant is sufficiently above the initial chamber pressure, and the stream Weber number is sufficiently high, the atomization is calculated to take place by vacuum-flashing of the stream. If one stream is injected before the other and does not flash, it is calculated to undergo single-stream breakup if there is sufficient distance for it to do so.

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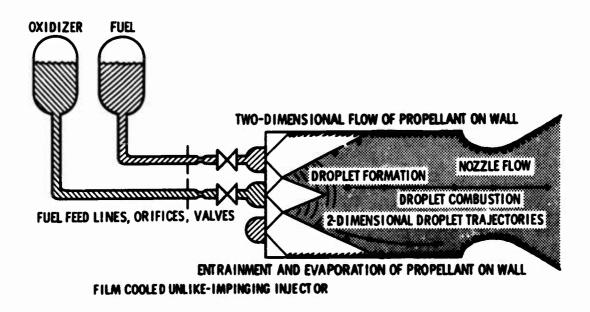


Figure A-1. Rocket System Diagram

Alternatively, it impinges upon the chamber wall, where part of it sticks as wall film and the rest is calculated to atomize by the wall impact. Later in the firing, atomization occurs by the impingement of the two unlike streams. The time-varying initial mean droplet size, the droplet size distribution, the initial droplet velocity vector, initial velocity distribution, and the distance traveled before the breakup process is completed are all calculated as time-varying functions of the injection and chamber parameters.

Two-dimensional trajectories of all of the droplets injected into the chamber are calculated with Reynolds-number-dependent drag coefficients recalculated for each droplet group at each time interval. The trajectories are followed until the droplets either burn completely, pass through the nozzle incompletely burned, or impact upon the wall. The normal bipropellant combustion rates of the droplets are based on droplet diameter and physical properties, droplet Reynolds number, combustion gas temperature and composition, and the presence or lack of ignition in the chamber. When a propellant constituent such as hydrazine has appreciable monopropellant character, this may also be modeled. The combustion rates are recalculated for each droplet group at each time interval to reflect the changing conditions in the chamber. The chamber gas properties are recalculated at each time interval, as is the chamber's velocity profile. The velocity profile is based on the axial distribution of contributions of vapor from burning droplets, igniter flows, vacuum flashing propellant streams, propellant burned off the wall-film and propellant vacuum-evaporated from the wall-film.

Hypergolic vacuum ignition is calculated in the vapor phase, based upon global chemical kinetics (Reference A-13). Various locations are examined to determine where ignition will occur first—the well-mixed vapors in the chamber, the most recently formed vapor mixture from the flashing streams, the boundary layer around the fuel droplets, or the boundary layer around the oxidizer droplets. Once ignition occurs anywhere, the entire chamber contents are presumed to ignite and will continue to burn until the criterion for extinguishment is met. Before ignition, the values for the temperature, molecular weight, and specific heat of the chamber gas are calculated, assuming well mixed but urreacted fuel and oxidizer vapor. After ignition, properties appropriate for equilibrium combustion products are used. After extinguishment, a distinction is drawn between quenced combustion gas and newly formed propellant vapors, so as to properly calculate reignition, if it occurs.

The film of liquid that builds up on the wall may flow in the axial and tangential directions, from its initial momentum, and under the influence of shear forces from the combustion gas and chamber wall. When the film has a tangential velocity component, the effects of spin-generated forces are considered. The film is partially burned off by the heat transfer from the hot combustion gases flowing past it. When threshold levels for an entrainment parameter are exceeded, entrainment from the liquid film is calculated. The axial variation in the wall-deposited material is approximated by dividing the chamber wall into 100 discrete axial segments. The deposition, flow, burnoff, entrainment, and vacuum evaporation from each segment are treated separately, so as to give the film thickness and composition profile and to correctly reflect the influence of unevenly distributed propellant. If the amount and rate of flow of the material on the wall are sufficiently high, some of the wall film material will be carried through the throat and ejected (one form of contaminant production).

The rate of burnoff from each axial segment of the wall is calculated from a heat transfer coefficient based upon the time-varying local Reynolds number, corrected for counter-current mass transfer and rippling of the surface. Entrainment from the surface is modeled based upon the local film thickness and velocity, the local velocity and density of the combustion gas, and the local surface tension of the film (Reference A-14).

When the local chamber wall temperature is sufficiently above the boiling temperature of the propellant droplets (at the momentary chamber pressure), an impacting droplet will bounce off instead of sticking (Reference A-15). When wall-film flows onto an axial segment that is sufficiently hot to cause the transition to film boiling, the film is detached from the surface, converted to a group of droplets, and entered into the ensemble of droplet groups in the chamber.

In calculating the axial and tangential velocities of the liquid material deposited on the wall, each axial element of fluid on the wall is treated at each time interval as an inertial free body being acted upon by shear from the wall, shear from the combustion gas, spin-induced forces, mass and momentum addition from the impinging droplets, mass and momentum loss from burnoff and entrainment, and with convection of fuel and oxidizer mass, axial and angular momentum at the upstream and downstream boundaries. The axial and tangential shear forces at the gas-liquid and liquid-wall interfaces are calculated based on correlations with appropriate local Reynolds numbers. It is presumed that the material in each axial segment is well-mixed at each time interval. When a hydrazine-derivative fuel and nitrogen tetroxide oxidizer are mixed on a wall segment, the neutralization reaction is presumed to take place to form the hydrazinium nitrate salt plus a remainder of whichever reactant is in excess of the stoichiometric ratio. This time-varying twocomponent system is used to estimate the physical properties of the wall film at each segment.

The assumption is implicitly made that injected fuel or oxidizer will react to give thermochemical equilibrium combustion products only when it is vaporized from the surface of a stream, droplet, or liquid film and the vapors subsequently are mixed with a mass of hot, ignited, combustion product gases.

After the valves have closed, the pressure in the chamber decays, most of the droplets leave the chamber, and the combustion in the chamber is calculated to be extinguished as soon as the time-varying calculated quenching distance exceeds the chamber diameter.

If the chamber pressure falls below the vapor pressure of the propellant in the injector dribble volumes, this material will flow out of the injector. The same technique is used to compute flow rate, atomization, and droplet trajectory as was used during the rest of the firing; with the exception that injector vapor pressure is now used instead of tank pressure to produce flow, and the flow impedances of the injector orifices are used instead of the whole system flow impedances. This dribbled material will be calculated to burn, be expelled through the throat unburned, or be deposited upon the chamber wall, depending upon the injector and chamber conditions.

When the chamber pressure falls below the vapor pressure of the material on the walls, it starts to evaporate, absorbing heat from the chamber

walls. The rate of evaporation from each segment of chamber length is calculated separately, based upon simultaneous solution of the Knudsen-Langmuir evaporation law and the heat transfer through the liquid film.

At any time, the propellant valves may be reopened to initiate a second pulse of firing. The propellant dribbled from the injector following the first pulse will constitute the injector void volume that must be filled to prime the injector for the second pulse; the propellant buildup on the walls will continue from the values attained during the vacuum evaporation following the first pulse, etc.

d. Comparison With Experiment

(1) Low Frequency Combustion Instability

The <u>TCC</u> program calculations were first tested by comparison with high-amplitude, low-frequency combustion instability produced in a small experimental engine (Reference A-16). The engine was operated with a family of four injectors, which differed only in impingement distance. The input to the program consisted of blueprint values for the engine and literature values for the propellant physical parameters, with no fitted coefficients of any kind. The calculated and experimental frequencies differed by no more than 7 percent while the amplitudes agreed within 5 percent for the entire family of injectors (References A-1 and A-2).

Figure A-2 shows the calculated chamber pressure for one of the injectors. It agrees in waveform with the experiment. The rate of rise is two to three times faster than the rate of fall, and it is cusped at the minimum points. The calculated fuel feedline flow rate was interesting in that it periodically assumed negative values; i.e., reverse flow. The integral of the reverse flow with time predicted a maximum backflow bubble volume of 0.4 cc. The back side of the injector was observed with high-speed motion picture photography through a transparent window, and the observed bubble size was in agreement with the calculations.

(2) Vacuum Hypergolic Start Transients

Figure A-3 illustrates a McDonnell Aircraft Company research engine that has been fired to investigate vacuum hypergolic start transients and pulse-mode behavior (Reference A-17). Figure A-4 shows a comparison of an experimental 25 msec duration pulse and a corresponding computed value. The two curves show surprisingly good agreement, especially in the prediction of events. The injector priming delay, the ignition delay, the initial surge from flash-atomized fuel, the succeeding pressure trough, the second pressure surge, the steady-state chamber pressure, and the cutoff delay are all in good agreement. The residual differences between the curves are associated with pressure waves in the experimental engine feed system, which was equipped with very long propellant lines. Pressure waves are not modelled by the lumped-parameter flow algorithm of the <u>TCC</u> program.

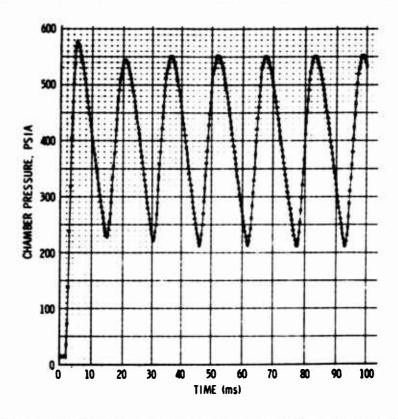


Figure A-2. Calculated Chamber Pressure Versus Time During Low-Frequency Combustion Instability

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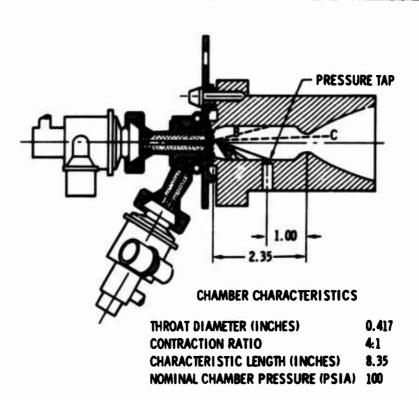


Figure A-3. Research Rocket Engine



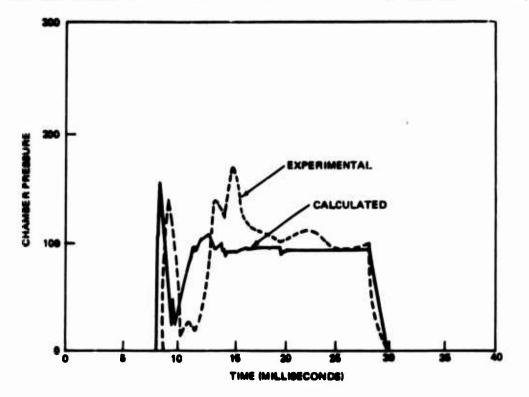


Figure A-4. Comparison of Calculated and Experimental High Altitude Start

(3) Contaminant Production

Contamination from a conventional bipropellant RCS engine takes one or more of three forms: reacted or unreacted propellant vapor (Reference A-18), incompletely burned droplets expelled through the throat (Reference A-19), and unburned propellant that impinges upon the chamber wall and is eventually ejected from the nozzle lip (Reference A-20). All of these sources are modeled by the TCC program. The only mode of contaminant production which has been experimentally measured is wall-film production. The wall-film production rate from the Marquardt R1-E engine was measured at three operating temperatures by P. J. Martinkovic. Calculations of wall film production from the similar R6-C engine have been calculated by the TCC program, and give acceptable agreement with experiment both in trend with operating temperature and in absolute values (See Section III of this report).

(4) Chamber Explosion Amplitude

Since the spatial distribution of unburned propellant in the form of streams, droplets, unreacted wall-film or hydrazinium-nitrate deposits is known in a time-dependent manner; it is possible to calculate the mean chamber pressure that could be produced at any instant by the explosive combustion of this material. The term "popping" is often used to describe the moderate amplitude explosions of the droplets and streams that may be periodically initiated by explosive reactions of the fuel and oxidizer streams near the impingement point. The stream hydraulic conditions and chamber conditions

that can produce impingement point explosions, and the conditions necessary for such an initiating explosion to propagate a detonation through the droplet spray contained in the chamber have been discussed in a number of publications (References A-21 through A-24).

Figure A-5 illustrates the calculated maximum volume-mean "popping" amplitude as a function of time for a particular 60-msec engine firing. Experimental bombings of this engine during the "steady-state" portions of the firing produced experimental chamber pressure amplitudes that had a mean value within 14 percent (or 1.2 experimental standard deviations) of the calculated value.

The time-dependent volume-mean chamber pressure amplitude that could result from a detonation of hydrazinium-nitrate deposits on the chamber walls is calculated in a similar way. Such detonations, which are often called "spikes", have been observed in a number of engines and can be quite destructive. Large transient accumulations of hydrazinium-nitrate have been calculated for two engines, which experimentally are known to produce "spikes" or "spikelike" behavior under certain conditions.

One of these engines is the McDonnell Aircraft Company research engine illustrated in Figure A-3 (Reference A-25). When this engine is operated in pulse mode under vacuum ambient conditions (using NTO and MMH propellants), the oxidizer flows from the injector dribble volume very much

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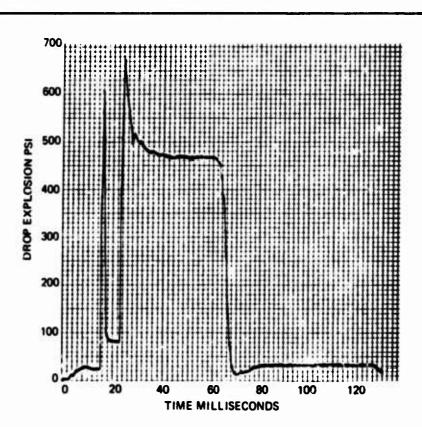


Figure A-5. Calculated Popping Amplitude Versus Time

faster than the fuel does at the end of a pulse because of its higher vapor pressure. Hence, in a train of very closely spaced pulses, successive engine pulses are likely to be started with imbalanced initial conditions, i.e., with an oxidizer dribble volume nearly empty and a fuel dribble volume nearly full. The firing of this engine (Figure A-4) has such an initial condition, corresponding to a previous pulse completed 95 msec prior to its start. The result of this initial condition is shown in Figures A-6 and A-7, which give the fuel and oxidizer flow rates as functions of time after first valve motion. The fuel side fills after 2.3 msec. This increases the resistance to flow and rather quickly limits the flow rate to a value about 40 percent greater than the steadystate value. The oxidizer side, however, takes 8.0 msec to fill, and in this time a flow rate almost 100 percent higher than the steady-state value is attained. The consequences are seen in Figure A-8, which shows the resultant stream momentum angle versus time. Between 2.3 and 8.0 msec, only fuel is entering the chamber. Consequently its momentum is centered along the axes of the fuel orifices, i.e., 35.5 degrees inward relative to the engine centerline. The fuel streams are flash-atomized to form a spray-cone of approximately 30-degree apex angle, as illustrated by A in Figure A-3. Some of the small droplets are turned sufficiently by the initial rush of vacuum vaporized fuel vapor to pass through the throat; however, much of this initial fuel is deposited on the chamber wall. When the oxidizer first enters the chamber 8.0 msec after first valve motion, its flow velocity is so disproportionately high that the resultant momentum angle is 13 degrees outward, as shown in Figure A-8 and as B in Figure A-3.

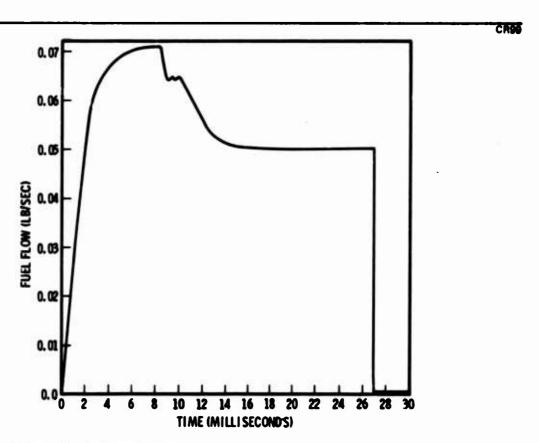


Figure A-6. Calculated Fuel Flow Rate Versus Time

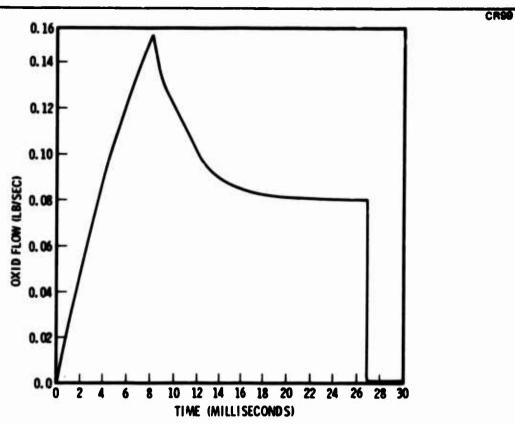


Figure A-7. Calculated Oxidizer Flow Rate Versus Time

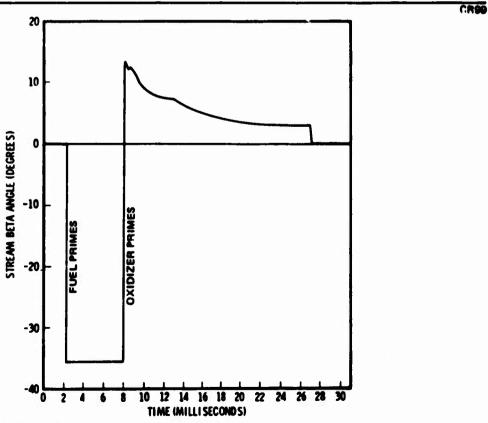


Figure A-8. Calculated Resultant Streem Angle Versus Time

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Eventually the propellant flow rates settle to the design value of 3 degrees outward, C in Figure A-3; but in the meantime, 272 milligrams of mixed fuel and oxidizer have been deposited onto the chamber wall, as shown in Figure A-9. This material can be burned off completely in an additional 60 msec of steady operation; however, if it detonates at the worst possible time, a volume-mean chamber pressure of 2,000 to 3,000 psi is produced, depending upon whether the unburned droplets and streams contained in the chamber are detonated also.

This is in general agreement with experiment which gave local measured peak pressures as high as 7,000 psi, with volume-mean chamber pressures on the order of 1,500 to 3,000 psi.

Another engine, for which calculations have been made, only exhibits spikelike behavior when it is run cold and in short pulses. Both of these conditions contribute to the accumulation of detonable material. Again reasonable agreement between calculated and measured overpressures are obtained (References A-5 and A-26).

Figures A-10 and A-11 show the results of very closely spaced repetitive firings of another small RCS engine. This engine will burn its walls clean very quickly after a start transient, however with firings which are too short and too closely spaced, the wall-deposited propellant can build up over a series of pulses.

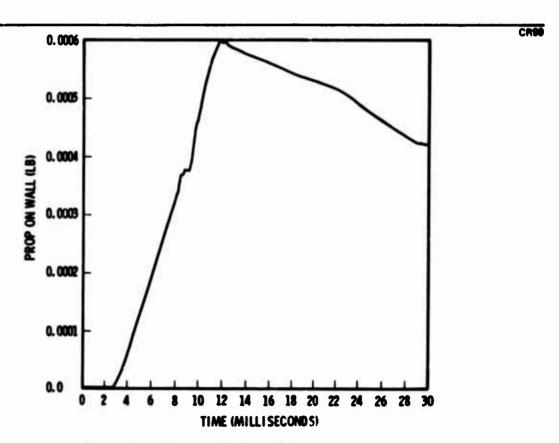


Figure A-9. Accumulation of Propellants on Chamber Wall Versus Time

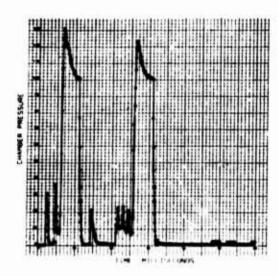


Figure A-10. Calculated Chamber Pressure for Two Short Closely Spaced Pulses

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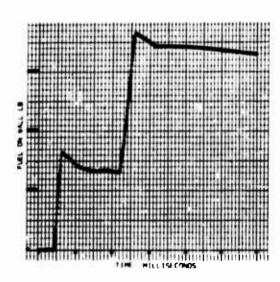


Figure A-11. Progressive Buildup of Fuel on Wall Over Two Short Closely Spaced Pulses

(5) Pulse-Mode Specific Impulse

The calculations of time-varying thrust and time integrated specific impulse are at present a rather uneven mixture of sophistication and oversimplification. The fully transient, two-dimensional treatment of the droplets, and the very detailed treatment of the wall-film behavior implies that the predicted variation of delivered specific impulse with pulse-width or with fraction of total fuel used as film-coolant should be accurate, and this has been verified by comparison with experiment.

The absolute values for calculated specific impulse have also given good agreement with experimental values; however, this depends in part upon the fortuitous cancellation of errors arising from oversimplications inherent in the present <u>TCC</u> program.

The most important si aplifying assumptions are (1) treating the combustion gases in the chamber as being well-mixed, (2) ignoring wall-film evaporation from axial heat conduction in the chamber wall, (3) ignoring secondary breakup of the droplets, and (4) using equilibrium thrust coefficients. The result of these simplifying assumptions is that the characteristic velocity is calculated a few percent too low, while the thrust coefficient is calculated a few percent too high. It is pleasant, but fortuitous that the absolute values of calculated specific impulse have been almost identical with the corresponding experimental values.

A. 2 ANALYSIS

a. Program Structure

The following logical sequence of events, with subroutine names and functions, illustrates the order in which calculations are performed during combustion chamber calculations, and for the two other modes of operation.

READER Read input data Write out input data IF "Work From Tape" G	O TO GRAPHS
START	
Do initializing steps Write out propellant and IF "Data Review Only" S	feed system derived values
INJECT	
Calculate propellant inje Calculate occurrence and	ependent propellant properties ection rates d extent of flashing es and median droplet diameters eces

DROPLT

Calculate whether ignition has occurred

Calculate aerodynamic drag on droplets

Calculate droplet burning rates

Calculate new droplet velocities, locations and masses

Delete droplets which have passed through the throat

Transfer wall-impinged droplets to wall-film arrays

WETWAL

Flow wall-film propellant

Burn off wall-film propellant

Entrain wall-film propellant

Transfer film-bound wall-film to droplet array

Vacuum-evaporate wall-film propellant

CHAMBR

Sum up axially distributed vapor sources and unburned propellant

Calculate gas outflow from chamber

Calculate new chamber state

Calculate whether extinguishment has occurred

Calculate new velocity profile and Reynolds number profile

Calculate thrust and chamber explosion capability

WRITER

Write output values on printer

Write parameters on data tapes

Increment model time

IF "Model time less than run duration" GO TO INJECT

GRAPHS

Plot droplet trajectories

Plot time-varying parameters

Plot wall-thickness of propellant

Calculate interval-average values

Plot movie frames

It may be seen that there are three principal modes of running the program.

If a nonzero value is given to "Data Review only" input Data Value 208, then the input data are read and they are printed with identifying headings. The initializing steps are performed including calculations of propellant physical property correlations and any feed system values obtained from the flow-rate overrides. Because of the extensive input data required for <u>TCC</u>, it is strongly recommended that any new data deck be checked out with this optional mode of operation before proceeding to make calculations. The person trying to assemble a data deck for the first time will find the input data print of value in assigning DDD1 array subscripts correctly to his data.

The second mode of operation is used to do combustion chamber calculations. After the initializing steps, the successive time increments are calculated by passing repeatedly through the subroutines INJECT through WRITER. The time-varying parameters are calculated and printed, and large amounts of time-varying output is stored on tapes 9 and 13. After the entire run duration time has been calculated, subroutine GRAPHS is entered and any designated plots and interval-averaged performance values are produced. Producing all the possible graphics or interval values would be quite time-consuming and expensive, so it is best to ask only for the most descriptive parameters such as chamber pressure and fuel and oxidizer injection rates at this time.

After the first set of graphics and printout has been examined, it is reasonable to remount tapes 9 and 13 and rerun the program, specifying "work from tape," input Data Value 10. In this mode any new parameter plots can be produced from the values recorded on tape 9. Chamber "snapshots" or "movies" can be produced from tape 13 and the cumulated and mean rate values may be calculated for the start transient, the steady-state portion of the firing and for the tailoff. The times at which these portions begin and end must come from the user's judgment, after examining the available graphics and printout, with the criterion for the time assignments based upon his particular needs. The "work from tape" option may be repeated as many times as desired, so long as the data tapes remain intact.

b. Subroutine READER

The <u>TCC</u> program has taken several forms through the course of its development. The READER subroutine has always read the input data block and reprinted it with descriptive headings. The CONTAM version uses four NAMELIST reads. Other versions exist which use formatted FORTRAN reads or input editors.

c. Subroutine START

The START subroutine converts english unit input to metric CGS units which are used for calculation inside the program. Curve-fit parameters are derived for later use in the propellant property calculations. When flow override values are specified, the corresponding hardware values are calculated and used in place of any conflicting data which might have been read into the input data array. Valve port areas are calculated first if flow rates and valve pressure drops have been supplied. If flow rates and injector pressure drops have been specified, the injection orifice discharge coefficients are calculated next. This calculation assumes that all of the injector pressure drop is localized in the injection orifices. When these calculations are done for multiple-ring injectors, the relative flow rates through the various rings must be specified. This data input is specified as a set of percentage values, but the total flow through all the rings is normalized to unity, so fractional flow or flow ratios or actual flows can be used equally well. The final flow override

value which can be calculated is an equivalent orifice to represent all the losses except valve and injector in the entire feed system. This is particularly useful for modeling complex systems with long lines, many fittings, filters, bellows, etc. The required input data are the difference between tank pressure and chamber pressure at a specified propellant flow rate. In making this calculation, it is presumed that the venturi is not cavitating. The START subroutine does all of the initialization required to commence the combustion chamber calculations and prints out derived values for propellants, feed systems, and chamber shape.

d. Subroutine INJECT

(1) Chamber Wall and Injection Temperature

The injector end and the throat end of the chamber are each assigned asymtotic temperatures for the firing and nonfiring conditions of operation, and half-times for the approach to these values. The time-varying values for injector end temperature and throat temperature are computed from these values. The axially varying chamber wall temperature is linearly interpolated with chamber length between these two values at each computational time interval.

The thermal transient of the propellant delivered after valve opening is approximated by using the injector end temperature until an amount has flowed from the injector equal to the hot slug of fuel contained in the injector at valve opening time. Following this the injection temperature varies linearly with flowed volume, from the injector temperature to the tank temperature, where it remains for the remainder of the firing (Figure A-12). The propellant in the injector is presumed to warm up to injector temperature whenever the valves are closed, so repeated firings start with propellant initially at injector temperature.

(2) Propellant Physical Properties

The density, viscosity, surface tension, vapor pressure, and liquid enthalpy are evaluated at the injection temperature, and are important in determining the dribble flowrates, the onset and extent of stream flashing, the flashing or impinging stream atomization droplet size, and the ignition delay time.

The densities of the liquid propellants are obtained from a mathematical approximation to a general correlation of reduced density versus reduced temperature along the saturation line (Reference A-27). The critical density is obtained from the propellant density given at a reference temperature.

The viscosity is approximated from a mathematical approximation to a generalized correlation of reduced viscosity versus reduced temperature. The critical viscosity is obtained from the propellant viscosity given at a reference temperature.

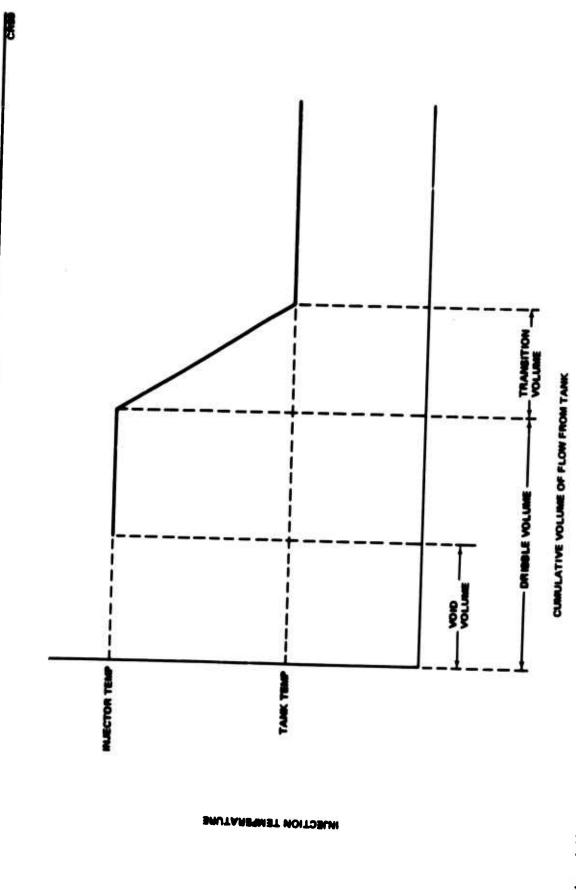


Figure A-12. Injection Temperature Profile Versus Flow from Tank

The surface tension is approximated using the parachor and a mathematical approximation to a generalized correlation of reduced density difference (liquid density-vapor density) versus reduced temperature along the saturation line. The parachor is a parameter which is closely related to the critical properties of a fluid.

$$[P] = \frac{Wm \sigma^{0.25}}{\rho_{\ell} - \rho_{V}} \tag{A-1}$$

For most materials it varies by no more than a few percent over the entire liquid range. The parachor for each propellant constituent is calculated from the propellant surface tension given at a reference temperature, and is used to calculate surface tension at other temperatures.

The vapor pressures of the propellants are obtained from a modification of the equation of Calingaert and Davis (Reference A-28):

$$ln P = A - \frac{B}{T - C}$$
 (A-2)

The value used for C is from an empirical correlation with the critical temperature of the material:

$$C = -7.39 + 0.03785 T_c + 0.000149 T_c^2$$
 (A-3)

A and B are evaluated for the fuel and oxidizer using the pressures and temperatures corresponding to the normal boiling point and the critical point.

Because of the scarcity of thermodynamic data for many propellants, the specific heats of vapor, liquid, and the solid form of each propellant are assumed to be constants which are neither temperature- nor pressure-dependent. When these values are supplemented by the melting point, heat of fusion, normal boiling point, latent heat of vaporization at the boiling point, and critical temperature, the straight-line relationship of Figure A-13 is obtained. This relationship is used to approximate the enthalpy of the vapor or condensed phases as a function of temperature.

(3) Feed System

The feed systems are approximated with single lumped parameters representing the inertial and resistive aspects of the feed systems:

$$\frac{dq}{dt} = \frac{P_t - P_c - \frac{1}{2} \rho R q |q|}{\rho \Sigma L/A}$$
 (A-4)

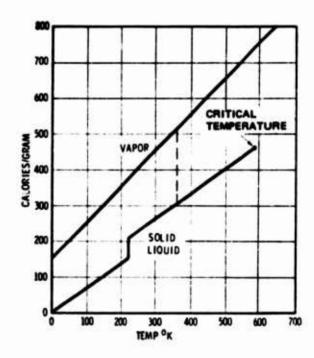


Figure A-13. Approximate Enthalpy of Monomethyl Hydrazine

where

q = volumetric flow rate through the line

P_t = tank pressure

P_c = chamber pressure

R = a coefficient representing the steady-flow pressure drop through the feed system

ΣL/A = the summation of segment length divided by crosssectional area for the series of segments which make up the feed system

Opening and closing of the valves are modeled by varying the value of R as a function of time. Flow reversals or initial start conditions which result in partially or fully gas-filled feedlines are simulated by varying both R and $\Sigma L/A$ as functions of time, so as to delete the resistive and inertial contribution of components which are not liquid filled at any particular instant. Equation A-4 does not model the compressibility of the propellant or the pressure waves in the lines. For cases in which wave action is important, a method-of-characteristics solution for a distributed parameter resistive-inertial-compressible feed system would be more accurate; however, the

present approximation is quite adequate to model flow events which take longer than five or six acoustic periods of the feed system (see Reference A-29).

After the valves close at the end of a firing, the chamber pressure decays, approaching the external pressure. When the chamber pressure falls below the vapor pressure of either propellant, it will start to flow from the dribble volume (the liquid-filled volume between the valve and the injector face). The flow rate out of the dribble volume is calculated using Equation A-4, but with R and $\Sigma L/A$ now restricted to the injector orifice values and with P_t replaced with the propellant vapor pressure evaluated at the injector temperature. As soon as the dribble volume is emptied, the dribble flow rate is set to zero.

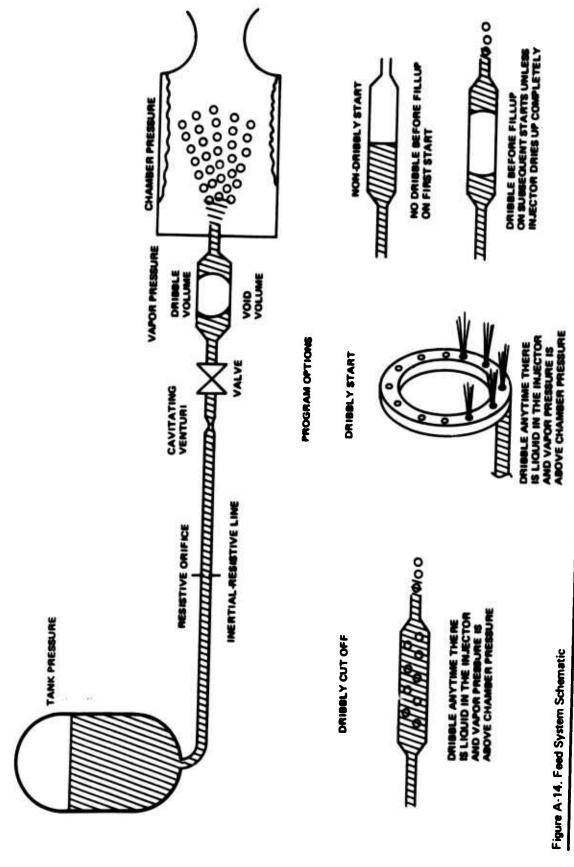
There are two optional assumptions which may be used to describe the injector priming process (Figure A-14). If a ring-type injector is being filled in a vacuum environment, some of the orifices will be wetted even before the ring volume is filled, and flow will be expected through these wetted orifices under the influence of the propellant vapor pressure. The initial dribble option is for this situation, and initial dribble flow rate is calculated the same as a postfiring dribble. The flow from the propellant tank into the dribble volume is calculated according to Equation A-4 but with R and $\Sigma L/A$ restricted to the values for the upstream hardware and with $P_{\rm C}$ replaced with the vapor pressure evaluated at the injector temperature. As soon as the injector is primed, the flow follows Equation A-4 as written.

The other injector priming option which may be used is appropriate for a smooth unbranched injector passage which would be expected to fill progressively from the valve out to the injector orifices without any dribbling of propellant before the passages are completely filled.

If high-amplitude combustion instability or high-amplitude pressure spikes temporarily reverse the flow in the propellant feedlines, the reverse flow is time-integrated to give the time-varying gas volume behind the injector. So long as there is gas behind the injector, the injection rate is set equal to zero, and the inertial and resistive contributions of the injector orifices are deleted from Equation A-4.

The static pressure in the throat of the cavitating venturi is monitored, and if it falls below the vapor pressure of the propellant, Equation A-4 is evaluated for the impedance of the upstream hardware only, with the venturi throat pressure set to the vapor pressure.

There are several complex modes that flow from the injector could take which have not been programmed in the current model. The propellant in the dribble volume could boil in a vacuum environment, with the well-mixed foam flowing through the orifice. Chugging or popping during the dribble periods could force noncondensible combustion gases into the dribble volumes to form foam at a pressure higher than the vapor pressure. If these modes of flow were found to be important they could be modeled.



(4) Flash Atomization and Flash Vaporization

The atomization process is calculated for one of several modes, depending upon the chamber pressure, the propellant injection temperature and the propellant injection rates. If the chamber pressure is sufficiently below the vapor pressure of the propellant at its injection temperature, the stream is presumed to flash atomize. If streams of both propellants are being injected and are not flashing, then impinging stream atomization is calculated. If only one stream is being injected, and it does not flash, the stream breaks up in a single-stream mode unless it hits the wall first, in which case it is calculated to break up by wall impact. If both streams are being injected, and only one of them flashes, the other stream is presumed to be unaffected by passing through the spray of small flashed droplets and behaves as though the other stream were absent.

Bushnell and Gooderum (Reference A-30) investigated the flashing of streams of water over a range of subatmospheric pressures, and found that the ratio of stream temperature to saturation temperature was almost a constant at the onset of flashing (with temperature measured on an absolute scale). Brown (Reference A-31) investigated the flashing of streams of superheated water at atmospheric pressure, varying temperature, orifice diameter, and stream velocity, and found that the temperature at the onset of flashing was a function of the stream Weber number. If these findings may be combined, the supersaturation temperature ratio at the onset of flashing may be represented as a function of stream Weber number. The following correlation is taken from the experimental data of Brown, but does not disagree with the data of Bushnell and Gooderum.

$$0 < N_{\text{we}} < 12.5$$
 $\frac{T_{\text{flash}}}{T_{\text{sat}}} = 1.138 - 0.005 N_{\text{we}}$ (A-5)

12.5 <
$$N_{we}$$
 < 24 $\frac{T_{flash}}{T_{sat}} = 1.085 - 0.00353N_{we}$ (A-6)

$$\frac{T_{flash}}{T_{sat}} = 1.0 (A-7)$$

Where the Weber number is

$$N_{we} = \frac{\rho_g V_j D_o}{2\sigma}$$
 (A-8)

There is a discontinuity in the function at a Weber number of 12.5 corresponding to some unknown change in the mechanism. The supersaturation temperature ratio is known to depend upon the surface roughness of the orifice; however, this effect has not been modeled in the TCC computer program, and the above relationship is for ordinary drilled holes, flowing full.

Brown has investigated the mean droplet diameter and droplet size distribution obtained when superheated water is flash-atomized at atmospheric pressure.

Gooderum and Bushell (Reference A-32) flashed water through drilled holes into a vacuum (~2 mm Hg) under otherwise comparable conditions and also obtained measurements of mean droplet diameter.

The functional effects of orifice size and temperature were quite different for the two experiments, probably due to the large differences in Weber number. Until the relationships are better understood, we will take 40 microns as a typical mean droplet diameter for the flash atomization of water, with reasonable values for orifice diameter, stream velocity, and degree of superheat. Brown described the flash atomization process as resembling the gas atomization process, with the gas being supplied by the explosive growth of bubbles in the superheated stream. For this reason we have applied a physical properties correction which would be suitable for the gas atomization process. The equation used in the program to describe flash atomization droplet size is:

$$\overline{D}_{\mathbf{p}} = 0.0065 \left(\frac{\sigma_{\mathbf{p}} \mu_{\mathbf{p}}}{\rho_{\mathbf{p}}} \right)$$
 (A-9)

where

 \overline{D}_{D} = median droplet diameter of propellant P, in centimeters

σ = surface tension of propellant, P

 μ = viscosity of propellant, P

 ρ = density of propellant, P

The drop size distribution obtained by flash atomization is much narrower than that obtained by impinging stream atomization.

When a stream of propellant is injected into a combustion chamber having a total pressure lower than the vapor pressure of the propellant, and if the Weber number is sufficiently high, then flash atomization will take place from explosive growth of vapor bubbles at nucleation sites in the stream. A certain amount of evaporation of propellant will occur during the flash atomization process. Following this, evaporation from the droplets will proceed, initially following the Langmuir-Knudsen law. If the chamber pressure increases to a value above the vapor pressure of one or both of the constituents, the vaporation can continue by molecular diffusion. Molecular-diffusion-evaporation can continue until the droplets have either equilibrated with the chamber vapors, are expelled through the throat, or impact upon the chamber wall. There are large areas of uncertainty in this complex process. One unknown is the amount of evaporation which takes place during the atomization

process, before the droplets are completely formed. Another is the chemical processes which could occur on the surface of a cold droplet of nitrogen tetroxide which is cryopumping vapors of hydrazine onto its surface.

Seamans, Vanpee, and Agosta (Reference A-13) have calculated the preignition chamber pressure history for the injection of one propellant component, based upon Langmuir-Knudsen evaporation of droplets having a mean diameter of 100 microns in a chamber 1 in. long. They found that the droplets had nearly equilibrated by the end of a time period (1.5 to 5.5 msec) corresponding to a nominal droplet residence time. Since their droplets were quite large for flash-atomized propellant, and since their chamber was quite small, it is reasonable to assume that phase equilibrium is generally attained, and to use this asymptotic value to describe the preignition evaporation.

The <u>TCC</u> program tests each portion of injected propellant by Equations (A-5 through A-8) to determine whether flashing occurs. If it does, the program calculates the fraction of the propellant which will vaporize to obtain local phase equilibrium at the instantaneous chamber pressure. The vapor formed is added to the vapors already in the chamber, the addition being made at the propellant injection point. The liquid which is unvaporized is assigned droplet properties, and the droplets are entered into the chamber droplet array. No further evaporation from or condensation on the droplets is calculated until they either impact upon the wall or until the ignition of the chamber contents occurs.

When liquid propellant is injected at a prescribed injection temperature, the relationships shown in Figure A-13 fix the injection enthalpy. The value of the chamber pressure at the time of injection is used in Equation (A-2) to determine the equilibrium temperature of the flashing propellant. When the enthalpies of the vapor and condensed-phase propellant are evaluated at the equilibrium flashing temperature, the fraction of the propellant which evaporates may be obtained:

$$H_{inj} = X \cdot H_{vap} + (1-X) H_{cond}$$
 (A-10)

or

$$X = \frac{H_{inj} - H_{cond}}{H_{vap} - H_{cond}}$$
 (A-11)

Since internal boiling and convective heat transfer inside the droplet both decrease after freezing of the surface, the program has the option of terminating the evaporation process once the liquid droplets have reached the triple point.

(5) Impinging Stream Atomization

When the chamber pressure is sufficiently high that the injected propellant streams do not flash, atomization occurs by either the impinging stream or single stream mode of breakup. The expression for droplet diameter

used in our program is based upon the work of Ingebo (Reference A-33), who examined the atomization of two identical streams of heptane, impinging at 90-degree included angle in air streams having velocities typical of rocketengine thrust chambers. Ingebo correlated his data:

$$D_{30} = \frac{D_{j}}{0.30 \sqrt{D_{j} V_{j}} + 0.0125 D_{j} \Delta V}$$
 (A-12)

where

D₂₀ = volume-number-mean droplet diameter

D: = injector orifice diameter

V; = injection velocity

 ΔV = velocity difference between the injected streams and the gas flow at the atomization point.

Preim et al reported (Reference A-9) that for rocket-engine conditions, where ΔV varies with axial distance from the injector, that 4,560 cm/sec (150 ft/sec) is an appropriate average value to use for ΔV .

Preim also offers a correction for liquid physical properties:

$$\frac{D_{A}}{D_{B}} = \left(\frac{\rho_{B} \sigma_{A} \mu_{A}}{\rho_{A} \sigma_{B} \mu_{B}}\right)^{1/4} \tag{A-13}$$

where

 D_A = mean diameter obtained with fluid A

ρ = density

σ = surface tension

= viscosity

Equations (A-12 and (A-13) are combined, along with the above value for ΔV , the physical properties of heptane and the multiplier need to convert from volume-number-mean diameter to mass median diameter. This yields a more general expression for initial median droplet diameter expected from a symmetrical self-impinging doublet injecting a fluid of specified properties.

$$\bar{D} = \left[\frac{D_{j}}{0.20 \sqrt{D_{j} V_{j}} + 39 D_{j}} \right] \left[1.725 \left(\frac{\sigma \mu}{\rho} \right)^{1/4} \right]$$
(A-14)

This expression is still insufficiently general for the analysis of unlike impinging streams during transients, where the momenta of the two streams may be very different, or where one stream may be missing completely. In the absence of experimental data, a simplifying assumption is used to estimate the effects of relative stream momenta. The simplifying assumption is that a small element of liquid in the stream approaching the impingement point is not affected by the condition of the other stream except through the formation of a quasi-stationary planar stagnation surface which acts to redirect the stream into the initial planar flow pattern leading to the formation of the fan. This leads to the conclusion that stream velocity is not the variable controlling fan formation in Equation A-14, but rather the component of stream velocity relative to the resultant direction of the combined stream. This leads to the generalization:

$$\overline{D}_{p} = \left[\frac{D_{p}}{0.24 \sqrt{D_{p} V_{p} \sin \left[\frac{\beta_{p} - \beta_{R}}{P_{p}} + 39 D_{p}} \right]} \left[1.725 \left(\frac{\sigma_{p} \mu_{p}}{P_{p}} \right)^{0.25} \right]$$
 (A-15)

where

 \overline{D}_{p} = median droplet diameter for propellant p

D_P = orifice diameter of the injector for propellant p

V_n = velocity of the jet of propellant p along its own centerline

 β_{D} = angle of the stream of propellant p relative to the engine centerline

 β_{R} = angle of the resultant stream relative to the engine centerline

The resultant angle of the combined fuel plus oxidizer stream is calculated assuming conservation of momentum in the axial and radial directions. When a stream which has not flashed intersects a stream which has flash-atomized, it is assumed that there is no momentum interaction; i.e., each stream continues in its original direction instead of being redirected to the "resultant" direction.

If a single stream impacts against the wall before it has traveled a sufficient distance to atomize in the single-stream mode, then atomization is presumed to take place by wall impact. Atomization by wall impact is based

upon Equation (A-15); however, the value used for relative velocity is simply the absolute value of the radial velocity of the stream, $\begin{vmatrix} V_y \end{vmatrix}$ which is substituted for the term V_p Sin $\begin{vmatrix} \beta_p - \beta_R \end{vmatrix}$.

Experimental verification of Equation (A-15) is still rather sketchy. It is obviously as correct as Ingebo's expression for the special case (symmetrical 90-degree impingement of heptane) from which it was derived. When only one stream is being injected, the stream angle and resultant angle are identical, and the following special case is obtained.

$$\overline{D}_{\mathbf{p}} = \frac{1}{39} \left[1.725 \left(\frac{\sigma_{\mathbf{p}} \mu_{\mathbf{p}}}{\rho_{\mathbf{p}}} \right)^{0.25} \right]$$
 (A-16)

This is an expression for single-stream breakup (with an assumed value for mean gas velocity) and was compared with the findings of Weiss and Worsham (Reference A-34) for this special case. The functional relationships appear to be in approximate agreement and the values predicted for mean droplet diameter under typical rocket conditions were in adequate agreement.

Since Equation (A-15) should account for stream impingement angle, it was compared with the findings of Heidmann and Foster (Reference A-35), who measured mean drop size as a function of impingement angle for symmetrical impinging streams. Again the predictions were in reasonable accord with the data. The droplet sizes obtained from the unlike impingement of streams of water and molten wax have been reported by Rocketdyne (Reference A-36). Evaluated at several points, the Rocketdyne droplet diameter values differed from the values of Equation (A-15) by no more than 5 percent.

All of the early experimental investigations of impinging stream atomization were done with turbulent flow through round orifices flowing full. Unfortunately most real engines have short, sharp-edged orifices with the flow detached. There are some, very small rocket engines whose orifices are in the laminar flow regime, and noncircular injection orifices are being used in some injectors. The effects of these parameters have recently been investigated by L. J. Zajac and others (Reference A-37). Detached flow and laminar flow each result in droplet diameters approximately half the correlation value of Equation (A-15). Droplet diameter correction factors may be used in TCC to account for these effects. The Zajac correction values for detached flow or laminar flow appear to give TCC results which agree with experiment.

(6) Atomization Distance

The distance which the impinged streams travel before the atomization is complete is calculated based upon investigations performed at NASA (References A-35 and A-38). The calculations used in this study presume that the distance to breakup is a prescribed number of orifice diameters in the absence of impingement (i. e., when β_p - β_R = 0 degrees), that it is as specified in Reference A-38 for impingement when β_p - β_R = 45 degrees, and that it varies linearly with sin (β_p - β_R) for intermediate values.

$$L_{p} = L_{45p} + \left(L_{op} - L_{45p}\right) \left(1 - \frac{\sin \left|\beta_{p} - \beta_{R}\right|}{\sin 45^{\circ}}\right)$$
 (A-17)

 L_{p} = fan length for complete atomization of propellant p

L_{45p} = experimental fan length for symmetrical 90-deg impingement

L_{op} = distance for single stream breakup of propellant p

 β_p = injection angle of propellant p

 β_R = resultant angle of the fuel plus oxidizer fan

When β_p - β_R is greater than 45 degrees, L_p is taken equal to L_{45p} degrees. When the injected streams are flash-atomized, the droplets are presumed to be formed immediately at the injection point.

The atomization is completed at an axial location in the chamber equal to the impingement distance plus the breakup distance multiplied by the cosine of the resultant angle. This axial location is a droplet group property called XIMP (I) in the <u>TCC</u> program.

To approximate the relatively inert condition of the propellants prior to breakup, we assume that there are no combustion, aerodynamic drag, or momentum interaction between particles of injected propellant in the interval between the injection point and the point of droplet formation, i.e., when X (I) < XIMP (I). Since the injection velocity is time-varying, this implies propellant bunching in this region. This behavior may be visualized from a plot of particle location versus time (Figure A-15). When the first propellant is injected, it is moving quite slowly due to the restriction of the partly opened valve and the inertia of the fluid in the lines. The propellant, which is injected a few milliseconds later, is moving much faster and overtakes the previously injected material.

Where the lines are close together, the bunching is most extreme, and the effects of velocity modulation produce the maximum effect in varying the arrival rate of propellant. In many rocket engines, this point of least-stable propellant flow is close to the location where droplets are formed and become available for combustion, i.e., XIMP (I). The injected propellant moves from the injection point to the impingement point along the direction of injection. After impingement, the stream moves in the direction of the resultant angle. It moves in this new direction until its atomization is complete, after which its two-dimensional trajectory is determined by the aerodynamic drag forces exerted by the chamber combustion gases. At the present time, the spreading of the fan is not modeled. When droplets are formed by flash-atomization, they are presumed to be formed at the injection point, and they are presumed to be immediately subject to aerodynamic forces and available for combustion.

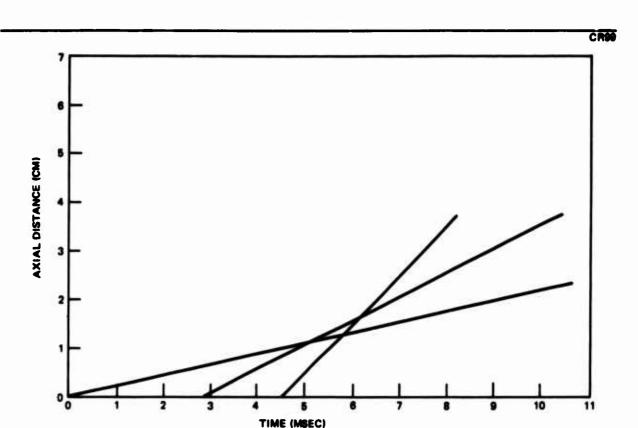


Figure A-15. Preimpingement Trajectory of Propellant

(7) Loading the Droplet Array

The droplets injected into a real combustion chamber are distributed in droplet size, initial direction, initial speed, in point of origin as newly formed drops, and possibly in composition, if the fuel and oxidizer are miscible and not excessively reactive. It is possible to model these distributed parameters by adding more than two droplet groups (one fuel, one oxidizer) to the droplet array during each computational time interval; however, the more droplet groups that are added, the more time-consuming the calculations become. If M values are used to approximate the distribution of each parameter and if N properties are treated as distributed, then the number of droplet groups per propellant is M^N . If M and N were each taken equal to five, then 6, 250 droplet groups would have to be added to the computational array during each time interval, instead of the 10 which are presently added. The storage location requirement for the droplet array would increase from 12,000 to 7,500,000 and the computing time on a high-speed computer would increase from about 2 minutes to about 20 hours. Obviously great restraint must be exercised in "improving" the calculations by this particular approach. It is this particular problem which keeps the TCC model from using a truly three-dimensional formulation at this time. It is still possible that a threedimensional version might be made economically practicable by restricting the initial directional variation to the larger droplet groups, as they are the only ones which can fly cross-wind any distance, and by introducing the directional variation in a time-dependent Monte Carlo approach where only a limited portion of the possible distribution is introduced at any given time iteration.

In the present computer program, the initial droplet diameter is the only parameter which is treated as distributed during "normal" operation; five values are used to approximate the experimental size distribution. For impinging stream atomization, the fifth mass quintile of the propellant has a droplet diameter approximately nine times that of the first quintile. This distribution is so wide that the distribution must be modeled if meaningful results are to be obtained.

During the periods that flash atomization is occurring, the droplet size distribution is much narrower, the fifth quintile diameter being only about twice the first quintile value (Reference A-31). However, at this time the initial directions of the droplets from the flashing stream are distributed over a cone having an apex angle of about 30 degrees. This distributed initial direction must be modeled in order to adequately describe the thickness profile of the propellant which is deposited on the chamber wall during the pre-ignition period.

For these reasons, only the droplet size distribution is modeled for impinging stream atomization, while only the initial direction distribution is modeled for flash atomization.

The droplet size distribution about the median value is approximated by dividing the injected mass of each propellant into five equal portions. Each of these portions is converted to a group of droplets having a diameter which is a prescribed multiple of the median droplet diameter. The droplet group size ratios come from a table of coefficients chosen to approximate the experimental droplet size distribution. The droplet diameter of the nth group is:

$$D_{n} = \overline{D} K_{n}$$
 (A-18)

The coefficients used to represent Ingebo's experimental distribution, a Rocketdyne experimental distribution and a widely used analytical approximation are given in Table A-I.

Table A-I. DROPLET DIAMETER DISTRIBUTIONS MID-QUINTILE DIAMETER/MEDIAN DIAMETER

	Ingebo Experimental (like doublets)	Log-Probability σ = 2.4	Rocketdyne Experimental (unlike doublets)
ĸ ₁	0.198	0.333	0.60
K ₂	0.759	0.645	0.81
к ₃	1.00	1.00	1.00
K ₄	1.23	1.55	1. 22
K ₅	2.30	3.00	1.55

After the droplet diameters of the fuel or oxidizer size groups are calculated, the mass per droplet and number of droplets in the group can be computed. The mass of each droplet in the nth group is:

$$M_{n} = \rho \frac{\pi}{6} D_{n}^{3} \tag{A-19}$$

The number of droplets in the nth group is:

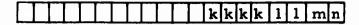
$$N_{n} = \frac{0.2 \rho q \Delta t}{M_{n}} \tag{A-20}$$

The feed system and atomization calculations provide values for the mass, median droplet diameter, initial velocity, etc., for the propellant which is injected into the combustion chamber at each time interval. Each group of droplets injected at each time interval is described by twelve variables, shown in Table A-II.

Table A-II. DROPLET PARAMETERS

AM (I)	Mass per droplet
AN (I)	Number of droplets in this group
X (I)	Axial location
Y (I)	Radial location
V (I)	Axial velocity
VY (I)	Radial velocity
SPAMOM (I)	Specific angular momentum
XIMP (I)	Axial atomization point
NUMS (I)	A "packed" integer number containing flags indicating whether or not the droplet group has passed the impingement point, whether it is fuel or oxidizer from which ring it originated, and the iteration at which it was injected.
VM (I)	Axial velocity immediately after impingement
VYM (I)	Radial velocity immediately after impingement
DM (I)	Mass evaporated from this group this iteration

The four values packed into the integer NUMS may be visualized by writing NUMS as an 8-digit decimal integer



n has a value of 1 before impingment, 0 after impingement m has a value of 1 for a fuel droplet, 0 for an oxidizer droplet ll has a value corresponding to the ring from which the droplet originated

kkkk has a value equal to the iteration at which the droplet was injected

As an example, a postimpingement fuel droplet which originated at the third ring during the 157th iteration would have a value for NUMS of 1570310.

NUMS may be assembled in the following manner: NUMS = K * 10000+IMRING*100+IFRAC*10+IXIM

The final complication of NUMS is that the iteration number is used only to designate whether a trajectory plot is to be made. For this reason, the iteration number is given a nonzero value only if the droplet is from the size group designated to be plotted.

The droplet array has 10 new droplet groups added to it from each injector ring at each integrating time interval, and has droplet groups deleted when the mass of the droplet falls to zero, the axial location exceeds the chamber length, or the radial location exceeds the chamber radius. This data array grows to represent the entire droplet population of the combustion chamber.

Each droplet group has the mathematical character of a vector in that 12 independent values are required for its description. The total droplet population of the chamber is then an array of droplet vectors or a matrix. The droplet population of the chamber can usually be adequately described using no more than 1,000 droplet groups or 12,000 computer storage locations. (The number is determined by the size of the integrating time interval.) The combustion chamber calculations consist of working on the array of droplet groups at each computational time interval to obtain the succeeding new value for droplet mass, droplet axial velocity, droplet radial velocity, droplet axial position, droplet radial position, and droplet evaporation rate. The storage locations reserved for the droplet group parameters are reused when the previous droplet group is burned completely, escapes through the throat, or sticks to the wall. If the droplet arrays fill up, leaving no place for newly injected propellant to go, an error message is printed and the calculations are stopped.

e. Subroutine DROPLT

(1) Ignition

Several types of ignition are modeled by the computer program. Calculations may be performed for sea-level or high-altitude ambient conditions, and for hypergolic or nonhypergolic propellants.

When a motor is being started at atmospheric pressure, using a liquid propellant igniter, it is assumed that droplets of the main propellants ignite instantaneously. When the fuel and oxidizer flow rates to the igniter and the axial location of the igniter port are specified, the initial composition, temperature, and axial velocity profile of the chamber gas are established. Although droplet ignition is presumed to be instantaneous, it would be erroneous to presume that the calculated start transient will necessarily be either smooth or simple, as experience has frequently shown it to be otherwise

When nonvolatile hypergols are injected at atmospheric pressure, the ignition results from complex liquid and vapor phase processes and it is necessary to obtain ignition delay values from the published literature based on experimental tests. When such an ignition delay time is prescribed, the droplet burning rates are set to zero during the preignition period. The delay time is counted as starting only after both fuel and oxidizer streams have reached the impingement point in the chamber.

When volatile hypergolic pairs are injected under near-vacuum conditions, the ignition depends upon the physical processes which contribute to the production of reactive vapors in the chamber. The calculation of the ignition delay time in the vapor phase is based on an equation given by Seamans, Vanpee, and Agosta (Reference A-13):

$$\tau_{\text{ignition}} = \frac{RT^2 \rho_{\text{gas}} C_{\text{p}}}{EAQ C_{\text{fuel}} C_{\text{oxidizer}}} Exp \left(\frac{E}{RT}\right)$$
(A-21)

where E, A and Q are the activation energy, frequency factor, and heat of reaction for the vapor phase reaction responsible for ignition. C_p is the specific heat of the vapor mixture in the combustion chamber and C_{fuel} or $C_{oxidizer}$ are the concentrations of fuel and oxidizer vapor in the chamber gases.

This function may be evaluated in either of three ways. One option is to assume that there is no axial mixing of vapors. In this mode, the relative concentrations of fuel vapor and oxidizer vapor come from the relative amounts flashed at each instant. A second option is to assume that all the vapors in the chamber are well-mixed. In this mode, the values for fuel and oxidizer vapor concentration used in Equation (A-21) are derived from a mass balance on the chamber considering the entire previous history of vapor generation and outflow. The third option presumes that at any instant there will be some location in the chamber which has the optimum stoichiometry for the quickest possible ignition. This optimum stoichiometry is found by examining the well-mixed vapor composition in the chamber, as well as the most advantageous composition in the boundary layers surrounding

fuel droplets and oxidizer droplets (when they are present), the composition having the greatest value for $(C_{\text{fuel}} \times C_{\text{oxidizer}})$ is the one used in the ignition calculations. Any ignition delay time longer than the gas residence time is discarded, as external ignition is ineffective. Because of the necessity of calculating repeated ignitions and extinguishments, unreacted fuel and oxidizer vapors are distinguished from residual quenched combustion product gases in calculating the concentrations used in Equation (A-21).

The vapor-phase ignition delay time is recalculated at each time interval, and the chemical delay time is added to the model time at which the calculation is made to obtain a projected time of ignition. The smallest value for the projected time of ignition is retained. Ignition is presumed to occur when the model time exceeds the smallest value for projected ignition time.

(2) Droplet Drag

The aerodynamic drag, heat transfer, and diffusion rates associated with the droplets depend upon a number of dimensionless groups which characterize the regime of flow; thus, the Reynolds number, Mach number, and Knudsen number must be considered. The Reynolds number is of great importance to all of the transport processes. Drag coefficients and heat transfer coefficients are correlated with droplet Reynolds number. The Mach number of the flow about the droplets is generally less than 0. 1 and its effects are ignored. The Knudsen numbers associated with the chamber processes have been calculated for a large number of conditions. The Knudsen numbers may be large for the first computational time interval, when the chamber pressure is taken equal to space ambient pressure: however, the Knudsen numbers have always been well into the continuum flow regime as soon as the first portion of either propellant is injected. Thus, free molecular flow effects should be negligible and are not considered in the correlations for vaporization rate, aerodynamic drag, and nozzle flow.

The velocity of the chamber gas relative to each droplet group must be calculated at each time interval in order to obtain the droplet aero-dynamic drag and droplet combustion rate.

The absolute value of the chamber gas velocity relative to the droplet is:

$$V_{rel.} = \sqrt{(V_{xgas} - V(I))^2 + (V_{ygas} - VY(I))^2}$$
 (A-22)

The Reynolds number is calculated for each droplet group at each time interval based upon the droplet diameter (obtained from the droplet mass, presuming spherical geometry) and the velocity difference between the droplet and the gas.

$$D = \left(\frac{6M_i}{\pi \rho_i}\right)^{1/3} \tag{A-23}$$

$$N_{Re} = \frac{\rho_{gas} D_{drop} V_{rel}}{\mu_{gas}}$$
 (A-24)

If the Reynolds number is less than 0.5, Stokes law acceleration is calculated:

$$A = \frac{18\mu_{gas} V_{rel}}{\rho_{drop} D^2}$$
 (A-25)

If the Reynolds number is greater than 0.5, the acceleration is calculated from the Newtonian drag law:

$$A = \frac{0.75 \text{ C}_{d} \rho_{gas} \text{ V}_{rel}^2}{\rho_{drop}}$$
 (A-26)

The drag coefficient data of Rabin (Reference A-39) are approximated by the functions:

$$0.5 < N_{Re} < 70$$
 $C_{d} = 27 N_{Re}^{-0.84}$ (A-27)

$$70 < N_{Re} < 59,200$$
 $C_{d} = 0.414 N_{Re}^{0.1433}$ (A-28)

$$59,200 < N_{Re}$$
 $C_{d} = 2.0$ (A-29)

(3) Droplet Evaporation Rate Calculations

The evaporation rate (combustion rate) of a stable fuel or oxidizer droplet is determined by the heat transport to the droplet surface and diffusional mass transport away from it. In the regime encountered in a rocket combustion chamber, the vapor effusion from the droplet has a very large effect on the heat transport process, and the effect of the forced convective flow about the droplet is also of great importance. An additional, important factor in determining the droplet life history is the transport of heat from the surface of the droplet to its interior.

If the droplet has appreciable internal circulation, the convective transport of heat to the interior of the droplet will be large, and there will be a considerable unsteady period when the droplet is warming from its initial injection temperature to the final steady temperature at which it evaporates. The unsteady droplet evaporation problem may be solved by simultaneous calculation of the heat and mass transport about the droplet together with the droplet state (Reference A-40).

If the droplet lacks internal circulation, then heat transport into the interior of the droplet is negligible (Reference A-41), and all the heat which reaches the droplet surface is used in vaporizing the outermost layer of fluid. When this is the case, the droplet state and vapor diffusion equations need not be solved.

According to Bond and Newton (Reference A-42), as cited by Hughes and Gilliland (Reference A-43), droplets will only circulate when the surface shear forces are sufficient to overcome the surface tension forces. An approximate criterion for the onset of circulation is

$$\Delta V = \sqrt{\frac{\sigma^2}{\mu_{gas} \rho_{gas} D}}$$
 (A-30)

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where ΔV is the difference in velocity between droplet and combustion gas at which circulation will start. This threshold is illustrated in Figure A-16.

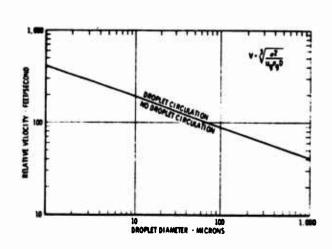


Figure A-16. Droplet Circulation Criterion

The gas properties in Figure A-16 are appropriate for a combustion chamber operating at 500 psia on liquid oxygen and RP-1. The surface tension is for room-temperature RP-1. According to this figure, circulation is expected in large droplets even at low relative velocity, but is not expected during most of the lifetime of the smallest droplets found in a rocket combustion chamber. In order to model accurately the evaporation of both large and small droplets in a time-varying environment, the presence or absence of circulation should be determined for each droplet group at each integration time interval, with heat transport into the droplet permitted only when circulation is present.

For the present calculations, the simplifying assumption is made that there is no heat transfer into the interior of any of the droplets. This permits accurate calculations for the small droplets responsible for much of rocket transient behavior, at the expense of accuracy in modeling the large droplets which largely determine combustion efficiency.

Heat transfer is correlated in terms of a Nusselt number. For a sphere

$$q = N_{NU}^{\pi K} D \Delta T$$
 (A-31)

where q is heat transfer rate and K is thermal conductivity of the vapor film.

From the assumption that all heat reaching the surface of the droplet goes to evaporate liquid

$$\mathring{\mathbf{M}} = \frac{\mathbf{q}}{\Delta \mathbf{H}} = \frac{\mathbf{N}_{\mathbf{N}\mathbf{U}}^{\mathbf{\pi}\mathbf{K}} \mathbf{D} \Delta \mathbf{T}}{\Delta \mathbf{H}}$$
 (A-32)

where \mathring{M} is the rate of evaporation and ΔH is the sum of the latent heat of the liquid and the sensible heat required to raise it from the injection temperature.

A noneffusing sphere in a nonconvective environment has a Nusselt number of 2.0, but the value for a droplet evaporating into a high-temperature environment is much lower. Godsave (Reference A-44) gives a simple but adequate estimate for the heat transfer to a burning droplet, assuming that thermal conductivity and specific heat of the vapor film are constant. His equation may be rearranged

$$N_{NU} = 2.0 \left[\frac{\Delta H}{C_p \Delta T} \ln \left(1 + \frac{C_p \Delta T}{\Delta H} \right) \right]$$
 (A-33)

The rate of fuel consumption from burning fuel-wetted spheres has been determined experimentally under forced convective conditions by several investigators (Reference A-45, A-46, and A-47). These experimental values are correlated (Figure A-17) by the equation

$$\frac{\mathring{M} - \mathring{M}_{0}}{\mathring{M}_{0}} = 0.25 \text{ N}_{Re}^{0.50}$$
 (A-34)

where M and M are nonconvective and convective consumption rates. This is rearranged and combined with Equations (A-32) and (A-33) to give

$$N_{NU} = \left[2.0 + 0.5 N_{Re}^{0.5}\right] \times \left[\frac{\Delta H}{C_p \Delta T} \ln \left(1 + \frac{C_p \Delta T}{\Delta H}\right)\right]$$
 (A-35)

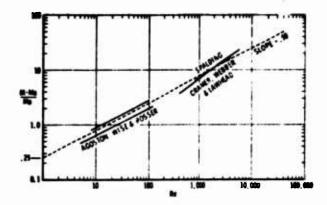


Figure A-17. Burning Rate Versus Reynolds Number

The term containing Reynolds number represents the effect of forced convection on a burning sphere. It is very similar in value to the correlation of Ranz and Marshall (Reference A-48) for the rate of evaporation from wetted spheres, but is preferred for this application because it was obtained in the presence of combustion and with temperature differences up to 3,000°K and Reynolds numbers up to 6,000. The Ranz and Marshall experiments were limited to temperature differences of 100°K and Reynolds numbers of 200.

Equations (A-32) and (A-35) may be combined to give

$$\dot{M} = \left(\frac{\pi K}{C_p}\right) D \left[2.0 + 0.5 N_{Re}^{0.5}\right] \times ln \left[1 + \left(\frac{C_p}{\Delta H}\right) \Delta T\right]$$
 (A-36)

which is the expression used to calculate the burning rate of a single droplet. In the computer calculations, $(\pi K/C_p)$ and $(C_p/\Delta H)$ are the parameters used to characterize the combustion properties of the fuel and oxidizer materials. These values are not treated as time-varying in the analysis in order to simplify the calculations. To obtain agreement with experimental burning rate values, C_p and K are evaluated at the arithmetic mean of the expected droplet surface and hot gas temperatures.

The temperature difference used in the burning-rate equation is the difference between the instantaneous droplet boiling temperature obtained from Equation (A-2), and the flame temperature or gas temperature surrounding the droplet.

The flame or gas temperature used in the burning-rate equation is obtained from the curve of chamber temperature versus composition (Figure A-18), the calculated composition of the chamber gas, and the known composition of the droplet which is carried as a subscripted variable. It is presumed that when a droplet of a given stoichiometry is evaporating into chamber gas having a different stoichiometry, every intermediate stoichiometry will be found in the diffusion zone surrounding the droplet and that the local temperature in the diffusion zone will correspond to the local stoichiometry. Thus the flame temperature used in the burning-rate equation

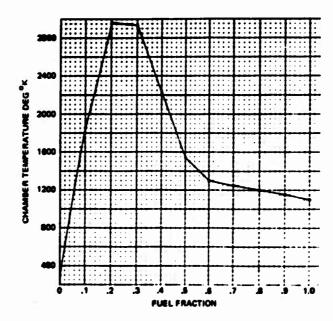


Figure A-18. Temperature Versus Stoichiometry

is the highest temperature found on the temperature versus composition curve in the interval between the chamber gas stoichiometry and the droplet stoichiometry.

There are many common propellant materials for which the vapor thermal conductivity is unknown in the 1,500 to 2,000°K temperature range. Nitric acid and UDMH are good examples. Experimental droplet burning rates have been obtained for these materials, and these burning rates may be used to infer this necessary information. Normally experimental burning rate data are given in terms of a burning rate constant, K'

$$\frac{dD^2}{dt} = -K' \tag{A-37}$$

in terms of our droplet burning rate expression

$$K' = \frac{8K \Delta T}{\rho \Delta H} \left[\frac{\Delta H}{Cp \Delta T} \ln \left(1 + \frac{Cp \Delta T}{\Delta H} \right) \right]$$
 (A-38)

This expression is solved for K using the experimental value for K'. When the mean value for thermal conductivity is obtained this way, the droplet combustion equations are no longer being used to derive the burning rates from basic principles, but merely become functions for extending a burning rate measurement to a different flame temperature.

Some propellant materials have the capability of burning as a monopropellant. Examples are hydrazine, propyl nitrate, and hydrogen

peroxide. When the monopropellant burning rate is known as a function of chamber pressure from liquid strand tests, the burning rate may be entered, correlated in the form

$$r = A + BP^{n} \tag{A-39}$$

where r is the burning rate in cm/sec and P is pressure in psia. The corresponding mass burning rate of a droplet is

$$\mathring{\mathbf{M}} = \mathbf{r} \pi \rho \mathbf{D}^2 \tag{A-40}$$

When propellant droplets can burn as either a bipropellant or as a monopropellant, the computer program calculates the burning rate for each droplet group in both ways at each time interval, and the larger of the two values is used.

(4) Droplet Velocity and Location

The simplifying assumption is made that the combustion chamber contour consists of seven butted cylindrical segments or conical frustum shapes. The corresponding assumption for gas flow direction is that it consists of a series of butted parallel flows or radial source or sink flows. This permits easy evaluation of the axial and radial components of chamber gas velocity at any point in the chamber. The direction cosines of the relative wind producing aerodynamic forces on the droplet are

$$C_{x} = \frac{V_{xgas} - V(I)}{V_{rel.}}$$
 (A-41)

$$C_{y} = \frac{V_{ygas} - V_{Y(I)}}{V_{rel.}}$$
 (A-42)

where $V_{rel.}$ is the absolute value of relative velocity as defined in Equation (A-22).

After the droplet acceleration and direction cosines have been calculated, the new velocity and location of the droplet group are calculated

$$V_{X(t+\Delta t)} = V_{X(t)} + A C_X \Delta t$$
 (A-43)

$$V_{Y(t+\Delta t)} = V_{Y(t)} + A C_{Y} \Delta t$$
 (A-44)

$$X_{(t+\Delta t)} = X_{(t)} + V_{x(t)} \Delta t + 0.5 A C_X \Delta t^2$$
 (A-45)

$$Y_{(t+\Delta t)} = Y_{(t)} + V_{Y(t)} \Delta t + 0.5 A C_Y \Delta t^2$$
 (A-46)

When the axial location of a droplet group exceeds the length of the chamber, the droplet group is removed from the array and its mass is added to the summation of unburned droplets that escape from the chamber.

When the radial position of a droplet group exceeds the radial dimensions of the chamber—and certain other conditions are satisfied—the droplet group is removed from the array and its mass is added to the summation of propellant mass which is coating the chamber wall at that particular axial location.

The droplets which remain in the chamber have their new mass calculated

$$M_{(t + \Delta t)} = M_{(t)} - \dot{M}\Delta t \tag{A-47}$$

where the value used for M comes from Equation (A-36) or (A-40).

Droplets are not permitted to decrease to negative masses. When the mass of droplet I decreases to zero all the other parameters for droplet group I are set to zero and the storage locations can then be used for a newly injected droplet group.

Each time that the mass decrement of a droplet is calculated from Equation (A-47), the value for the evaporation from that droplet group at that time interval is also calculated and is stored as DM(I) until it can be used in the summations which give the velocity profile in the chamber and the total burning rate for the chamber.

f. Subroutine WETWAL

(1) Wall-Film Flow Calculations

When a droplet group moves radially to the location of the combustion chamber wall, its fuel or oxidizer mass is deleted from the droplet array and is added to the appropriate location in a 100-member array which represents the axial distribution of fuel or oxidizer deposited on the chamber wall. At the same time its axial momentum and its tangential angular momentum are added to the corresponding locations in 100-member arrays holding these values. From these values the axial and tangential velocities, the gross stoichiometry, the masses of adduct and of free fuel or oxidizer can be calculated for each axial segment of chamber. The flows at each axial location may be regarded as two-dimensional in that axial and tangential momenta, velocities and drag forces are calculated, the wall treatment is one-dimensional, however, in that there is only one lumped value for the wall variables at each axial location. Tangential variation could easily be programmed, but computer time would increase considerably. The first process calculated for the wall film is change of velocity due to chamber gas shear force, wall shear force, and spin-generated axial force.

The gas shear force acting upon each axial segment of the liquid film is taken from pipe-flow theory:

$$F_{g} = f(N_{Re}) \pi D \Delta X 1/2 \rho_{gas} (V_{gas} - V_{film})^{2}$$
(A-48)

f is the Fanning friction factor, evaluated as a function of Reynolds number based upon chamber gas density, local axial gas velocity, chamber diameter, and gas viscosity. D is local chamber diameter, ΔX is the length of the wall segment.

The shear force from the chamber wall is:

$$F_{w} = f(N_{Re}) \pi D \Delta X 1/2 \rho V_{film}^{2}$$
(A-49)

here the Reynolds number is based upon the hydraulic radius of the film

$$N_{Re} = \frac{4 \delta V_{film Pfilm}}{\mu_{film}}$$
 (A-50)

where δ is film thickness.

The spin-derived net axial force on element i is:

$$F_{s} = \left\{ \frac{\rho_{i} \ V_{Ti}^{2} \delta_{i}^{2} \left(D_{i} + \delta_{i}\right)}{D_{i}} - \frac{\rho_{i+1} \ V_{Ti+1}^{2} \delta_{i+1}^{2} \left(D_{i+1} + \delta_{i+1}\right)}{D_{i+1}} \right.$$

$$\left. + \left[\frac{\rho_{i} \ V_{Ti}^{2} \delta_{i}}{D_{i}} + \frac{\rho_{i+1} \ V_{Ti+1}^{2} \delta_{i+1}}{D_{i+1}} \right] \times \left[\frac{D_{i+1}^{2} - D_{i}^{2}}{4} \right] \right\}. \quad (A-51)$$

where ρ is film material density, V_T is tangential velocity of the film, δ is film thickness, and D is chamber diameter. The net force acting upon the element of fluid is the sum of the above contributions, and the resulting change in axial velocity of the element is

$$\Delta V = \frac{F_g - F_w + F_s}{M} \Delta t \tag{A-52}$$

The wall shear force acting in the tangential direction is:

$$F_{T} = f(N_{Re}) \pi D \Delta X 1/2 \rho_{film} V_{T film}^{2}$$
(A-53)

where Reynolds number is:

$$N_{Re} = \frac{4 \delta V_{T film}^{\rho} film}{\mu_{film}}$$
 (A-54)

The tangential velocity decrement for one time interval is:

$$\Delta V_{T} = \frac{F_{T}}{M} \Delta t \tag{A-55}$$

The axial flow calculation is very simple. The axial displacement of the material in segments i during one time interval is:

$$\Delta L_i = V_i \Delta t \tag{A-56}$$

The corresponding uniform mass flow across the downstream boundary is:

$$\Delta M_i = \frac{V_i M_i \Delta t}{\Delta X}$$
 (A-57)

where ΔX is the length of the chamber segment.

For a small axial displacement of the material on segment i (less than half the segment length) the corresponding fraction of the mass of segment i is removed from segment i and added to segment i+1. Corresponding fractions of the axial and tangential momentum of segment i and of the gross fuel and oxidizer composition of segment i are transferred as well. When larger displacements are involved (which is very rare), a different algorithm is used for the sake of stability. The large-displacement algorithm distributes the mass of segment i over several downstream segments in amounts proportional to what would result by shearing segment i from the initial shape of a rectangle to the final shape of a parallelogram, with the base still in place, but with the top removed far enough downstream to move the center of mass of the element one displacement distance. The mass which flows out of the downstream end of the chamber at each time interval is considered to be the wall-film contaminant production.

(2) Wall-Film Burnoff

The burnoff from each axial segment of the wall is calculated from a heat transfer coefficient taken from the work of Gator (Reference A-14). The heat transfer coefficient is basically a function of chamber Reynolds number based on length, but is corrected for "blowing" from the evaporating surface, rippling of the surface, and temperature ratio. Evaporation rate is:

$$\dot{\mathbf{M}}_{i} = \frac{\mathbf{H}_{i} \mathbf{A}_{i} \Delta \mathbf{T}_{i}}{\Delta \mathbf{H}_{i}} \mathbf{C}_{\mathbf{e}_{i}} \mathbf{C}_{\mathbf{r}_{i}} \mathbf{C}_{\mathbf{T}_{i}}$$
(A-58)

 \mathring{M}_i is the rate of burnoff of propellant from the ith wall segment. H_i and A_i are the heat transfer coefficient evaluated for the conditions at i and the chamber wall surface area of the ith segment. The value for H_i is calculated:

$$H_i = .0295 C_{p_{gas}} \rho_{gas} V_{i_{gas}} N_{Re_i}^{-0.2} (D_i/L_i)^{0.2}$$
 (A-59)

 N_{Re_i} is the Reynolds number based on chamber diameter evaluated at the ith segment, and L_i is the downstream distance of the element.

The correction for simultaneous heat and mass transfer is calculated:

$$C_{e} = \left(\frac{\Delta H_{i}}{C_{p_{i}} \Delta T}\right) \ln \left(1.0 + \frac{C_{p_{i}} \Delta T}{\Delta H_{i}}\right)$$
(A-60)

Where ΔH_i is the heat required to evaporate or pyrolyze unit mass of the propellant mixture at the ith wall segment, C_{pi} is the specific heat of the vapor produced and ΔT is the difference between the chamber gas temperature and the surface temperature of the evaporating propellant, the ripple correction is calculated:

$$C_r = 1.0 + 35.4466 \rho_{gas}^{-0.4} V_{gas}^{-0.8}$$
 (A-61)

with the maximum value for C_r limited to 2.5. The temperature ratio correction is:

$$C_{T} = \left(\frac{T \text{ gas}}{T \text{ surface}}\right)^{0.25} \tag{A-62}$$

(3) Wall-Film Entrainment

Whenever the chamber gas shear stress is high enough and the surface tension small enough, the wall film will be expected to atomize from the surface and the droplets entrain into the combustion chamber gas. An entrainment parameter is calculated:

$$G_{i} = \frac{\rho_{gas}^{0.5} V_{gas_{i}}}{\sigma_{i}} \left(\frac{T gas}{T surface}\right)^{0.25}$$
(A-63)

Whenever the entrainment parameter is greater than 1.5 for an element of wall film, the entrainment rate for the element is calculated

$$\dot{M}_{i} = M_{i} V_{i} \left(1.0 - e^{-.001313 (G_{i} - 1.5) \Delta X}\right)$$
 (A-64)

The entrained mass is removed from the mass on the element at wall, and is presumed to burn immediately; i. e., it is added to the chamber gas array at the ith segment of the chamber.

(4) Wall-Film Detachment by Film Binding

It is well known that a liquid, boiling through contact with a hot surface, will detach itself from the surface when the surface temperature exceeds the equilibrium boiling temperature by too great a margin. This appears to be the mechanism responsible for the phenomenon illustrated in Figure 6 of McFarland's paper on internal regenerative cooling (Reference A-49), which shows the heat transfer coefficient from the hot chamber wall to a film of MMH decreasing as the chamber wall temperature increases from the boiling temperature to 1.5 times the boiling temperature. Below the boiling temperature, the heat transfer coefficient is appropriate for heat transfer to a nonboiling liquid film, while at 1.5 times the boiling temperature, the heat transfer is roughly what would be expected for heat transfer to saturated MMH vapor. This temperature ratio of 1.5 for detachment is not far from the value observed by Tamura and Tanasawa (Reference A-15) for the complete detachment of water from a hot surface. Several hydrocarbons

and alcohol appear to detach at considerably lower temperature ratios (1.15 to 1.3 times the boiling temperature). Attempts to correlate this behavior with the reduced temperature corresponding to equilibrium boiling (boiling temperature/critical temperature) as suggested by Bernath (References A-50 and A-51) were not successful.

In the TCC program, the wall film is detached from the wall whenever the liquid film advances onto a segment of wall surface having a temperature greater than 1.5 times the boiling temperature (because of hysteresis in the wetting process, droplets impinging against a dry portion of chamber wall are presumed not to adhere above a temperature ratio of 1.2). The detached wall film is converted to a group of droplets which are introduced into the droplet array. The diameter of the newly formed droplets is taken to be 3 times the thickness of the wall film from which they were produced. This ratio leaves the surface to volume ratio unchanged in the transformation from detached film to droplets. If the droplet formation was driven solely by surface tension forces, the droplet diameter to film thickness ratio would have to be higher than this. If droplet formation was aided appreciably by aerodynamic forces, the droplet diameter to film thickness ratio could be appreciably lower.

(5) Vacuum Evaporation Rate

When the chamber pressure falls below the vapor pressure of the fuel and oxidizer deposited on the wall, they undergo Knudsen-Langmuir evaporation, with heat simultaneously being transferred from the chamber wall.

The evaporating wall film is treated as being locally thermally quasi-steady (Reference A-52) and the local vacuum evaporation rate from each of the 100 axial segments of wall is presumed to balance the local heat transfer rate from the chamber wall to the evaporating surface. The expression for Knudsen-Langmuir evaporation is:

$$M_{pi} = A_{pi} E_{p} \left(P_{vap p}(T_{si}) - P_{c}\right) \left(\frac{Mw_{p}}{2 \pi R T_{si}}\right)^{1/2}$$
(A-65)

Where \dot{M}_{pi} is the evaporation rate of constituent p from the ith axial segment of the chamber wall, A_{pi} is the exposed surface area of constituent p on the ith segment. E_p is the accommodation coefficient of constituent p. P_{vap} p (T_{si}) is the vapor pressure of constituent p at the local surface temperature T_s . P_c is the chamber pressure. Mw_p is the vapor molecular weight of constituent p, and R is the gas constant. The effects of the neutralization reaction removing fuel or oxidizer from the mixed layer with the consequent formation of hydrazinium nitrate, and the vapor pressure reduction from solution effects are considered roughly by decreasing the vapor pressure of the free fuel or oxidizer to be proportional to its mass fraction in the wall-film mixture. Raoult's law is not presumed to apply to the very nonideal components of the wall film, and the experimental phase behavior of these mixtures has never been determined. The hydrazinium nitrate itself is treated as being nonvolatile in accordance with the experimental

findings of H. H. Takimoto et al at the Aerospace Corporation (References A-53, A-54, and A-55). The expression for heat transfer through the wall film is:

$$\dot{M}_{pi} = \frac{A_{pi} K_{p} (T_{wi} - T_{si})}{\delta_{i} \Delta H_{p}}$$
(A-66)

Where K_p is thermal conductivity of constituent p, T_{wi} is the chamber wall temperature at segment i, δ_i is the thickness of coating on segment i, ΔH_D is the latent heat of evaporation of constituent p.

Equations (A-65) and (A-66) must be simultaneously solved to obtain the values of \dot{M}_{pi} and T_{si} . This is done by Newton iteration. The fuel and oxidizer deposited on each axial segment of chamber wall surface are presumed to be well-mixed, reacted to completion to give hydrazinium nitrate plus fuel or oxidizer excess, and are regarded as uniformly covering the entire area of the wall segment.

g. Subroutine CHAMBER

(1) Axial Summations of Propellant Vapor

The 100-member arrays DXMF and DXMO are used to bring together the axially distributed contributions of fuel and oxidizer vapor mass from the igniter (if one is used), from flash vaporization of super-heated fuel or oxidizer, from droplet vaporization, from burnoff of wall film, from fuel or oxidizer mist entrained from the wall film (and presumed to burn rapidly), and from wall-film material vacuum-evaporated by heat transfer from the chamber wall.

At the same time that the vapor source summations are being obtained, sums are obtained representing atomized fuel and oxidizer in the chamber and fuel and oxidizer which has been injected but which is not yet atomized (i.e., streams and fans). These sums are used to obtain the chamber explosion capabilities of the engine.

Each member of the DXMF or DXMO array holds the production of fuel or oxidizer vapor at its particular axial loaction at that particular time interval. The DXMF and DXMO arrays are axially integrated to give the 100-member XMF and XMO arrays. Each member of these arrays holds the sum of all fuel or oxidizer vapor sources upstream of its axial location. These arrays are used to obtain the total vapor production at each time interval and to obtain the chamber gas velocity profiles.

(2) Gas Outflow from the Chamber

Conventional steady flow, compressible gas, nozzle flow equations are used to calculate the instantaneous mass flow rate through the nozzle throat. The use of these equations together with the assumption of constant pressure throughout the chamber restricts the validity of TCC

calculations to cases where the rate of change of pressure is sufficiently slow that only a small change in pressure occurs during a single acoustic time period of the chamber or nozzle.

The nozzle is sonic when

$$\frac{P_{e}}{P_{c}} \leq \left(\frac{2.0}{\gamma+1}\right)^{\gamma/(\gamma-1)} \tag{A-67}$$

Subsonic mass flow rate is calculated

$$\dot{M}_{T} = A_{T} P_{C} \sqrt{\frac{2.0 \ \gamma W_{m}}{(\gamma-1) \ RT_{C}}} \sqrt{\left(\frac{P_{e}}{P_{c}}\right)^{2/\gamma} - \left(\frac{P_{e}}{P_{c}}\right)^{(\gamma+1)/\gamma}}$$
(A-68)

Sonic flow rate is calculated:

$$\dot{M}_{T} = A_{T} P_{C} \sqrt{\frac{\gamma Wm}{RTc} \left(\frac{2.0}{\gamma+1}\right)^{(\gamma+1)/(\gamma-1)}}$$
(A-69)

The rates at which fuel- and oxidizer-derived mass leave the chamber are

$$\dot{M}_{F} = M_{T} \times F \tag{A-70}$$

$$\dot{M}_{O} = M_{T} \times (1 - F)$$
 (A-71)

where F is the mass fraction of the chamber gas which fuel-derived.

(3) Chamber Gas State

At each integration time interval, the expressions for flashing of the propellant streams, evaporation of droplets, burnoff of propellant on the walls, vacuum evaporation of propellant on the walls, and the assigned igniter flows are used to calculate the amounts of fuel-derived mass and oxidizer-derived mass which have been added to the chamber. Nozzle flow equations are used to calculate the amount of gas leaving the chamber, with the assumption being made that the material flowing through the nozzle has the same composition as the well-mixed material in the chamber. The new amounts of fuel-derived gas mass and oxidizer-derived gas mass contained in the chamber are then calculated:

$$M_{F(t+\Delta t)} = M_{f(t)} + \text{fuel evaporated - fuel exhausted}$$
 (A-72)

$$M_{O(t+\Delta t)} = M_{O(t)} + \text{oxidizer evaporated - oxidizer exhausted}$$
 (A-73)

The total mass of gas in the chamber is the sum of the fuel-derived and oxidizer-derived mass

$$M_{T} = M_{F} + M_{O} \tag{A-74}$$

The stoichiometry of the chamber gas is expressed in terms of the fraction of the total gas mass which is fuel-derived

$$F = \frac{M_F}{M_F + M_O} \tag{A-75}$$

When combustion is taking place in the chamber, the temperature, molecular weight, and thermal properties of the chamber gas are functions of the elemental compositions and heats of formation of the fuel and oxidizer, the stoichiometry and the chamber pressure. Since the composition and heat of formation of the fuel and oxidizer are fixed for a given set of propellants, and since the product composition is only a weak function of pressure, the temperature, molecular weight and thermal properties of the chamber gas are approximated as a function of F alone with only small error. Temperature, molecular weight and gamma are stored as 11-point tables with linear interpolation being used to obtain values for intermediate values of F. Figures A-18, A-19, and A-20 illustrate the values used for the white furning nitric acid - UDMH propellant combination.

When the combustion chamber contents are not ignited, the gas properties are calculated for the unreacted fuel and oxidizer vapors, assuming ideal gas behavior.

After chamber temperature and molecular weight are estimated as functions of the gas stoichiometry, the chamber pressure is calculated

$$P_{c} = \frac{\rho_{RT}}{W_{m}} \tag{A-76}$$

The density is obtained from the known total gas mass and total chamber volume, where \mathbf{U}_{c} is chamber volume

$$\rho = \frac{M_T}{U_c} \tag{A-77}$$

(4) Extinguishment

Since the droplet population of the chamber declines very rapidly after the propellant valves are closed, the vacuum evaporation of fuel and oxidizer film on the wall becomes the major source of fuel and oxidizer vapors very shortly after valve closure. For this reason the criterion for extinguishment is based upon the combustion chamber gas state rather than upon the conditions in the droplet boundary layers. Extinguishment in the gas is predicted from experimental quenching distance values taken from data on the combustion of premixed gases (Reference A-56). Quenching distance of premixed gases is a function of the pressure, and of

CRSS

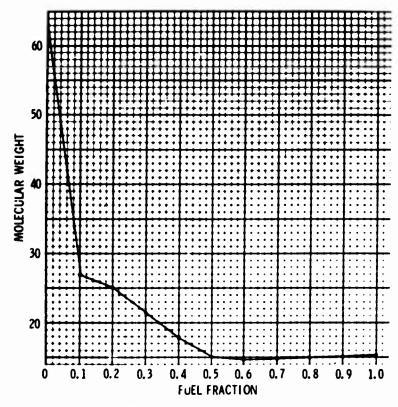


Figure A-19. Molecular Weight Versus Stoichiometry



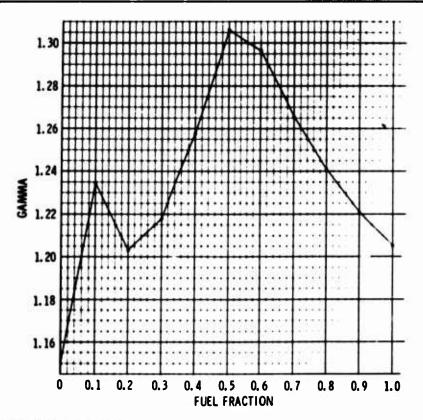


Figure A-20. Gamma Versus Stoichiometry

the initial temperature and stoichiometry of the combustible mixture. For the present purposes, the effects of stoichiometry and initial temperature can be lumped together in terms of the corresponding final gas temperature. The data on propane-air quenching distance are correlated adequately by:

$$q = 3.6 \times 10^{17} P^{-1.0} T^{-4.0}$$
 (A-78)

Where q is quenching distance in cm, P is pressure in dyne/cm², and T is equilibrium final gas temperature in °K. When the calculated value of q becomes larger than the chamber diameter, extinguishment is presumed to occur.

(5) Axial Gas Velocity

The axial gas velocity is calculated at 100 equally spaced axial locations at each integrating time interval to give the gas velocities used in the droplet drag and evaporation rate calculations, the wall burnoff calculations and the wall-film flow calculations during the next time interval. When velocities are required at intermediate locations, linear interpolation is used. The simplifying assumption is made that at any instant the pressure is constant throughout the chamber and the gas is well mixed. Since the computation time interval is typically of the order of 1 to 10 times the longitudinal acoustic period of the engine, the uniform chamber pressure assumption seems justified for engines with large contraction ratios.

The axial gas velocity at any axial location L_i may be obtained through the continuity equation, a mass balance on the chamber volume upstream of location L_i , and the assumption of uniform density throughout the chamber. The axial velocity at L_i is

$$V_{i} = \frac{M_{i}}{\rho_{gas} A_{i}}$$
 (A-79)

where \mathring{M}_i is the rate at which gas mass is flowing past the section at L_i and A_i is the cross-sectional area at L_i . The value for \mathring{M}_i is obtained from a mass balance on the chamber volume upstream of L_i .

$$\dot{M}_{i}$$
 = Evaporation rate upstream of L_{i} - accumulation rate upstream of L_{i}

$$\dot{M}_{i} = (XMF_{i} + XMO_{i})/\Delta t - \frac{U_{i}}{U_{100}} \left[(XMF_{100} + XMO_{100})/\Delta t - \dot{M}_{T} \right] (A-81)$$

 U_i is the chamber volume upstream of segment i, and U_{100} is the total chamber volume. M_T is the mass flow rate through the nozzle.

Reynolds number, based upon chamber diameter, is calculated at 100 axial locations:

$$N_{Re_i} = V_i \rho_{gas} D_i/\mu_{gas}$$
 (A-82)

(6) Thrust Calculation and Explosion Potential

The vacuum thrust is calculated as the sum of the momentum rate of the ejected droplets and the theoretical vacuum thrust of the gas:

$$F = P_c A_t C_F + \frac{1}{\Delta t} \sum_{\text{droplets ejected}} V_i M_i$$
(A-83)

When the chamber contents are ignited, the C_F is interpolated from a table of C_F versus stoichiometry of the chamber gas. When the chamber products are not ignited, C_F is interpolated from a table calculated for the mixture of unburned vapors. In calculating the C_F table for the vapors, first the ratio of exit pressure to chamber pressure must be obtained from the exit area ratio:

$$\frac{1}{\epsilon} = \left(\frac{\gamma+1}{2}\right)^{\frac{1}{\gamma-1}} \left(\frac{P_e}{P_c}\right)^{\frac{1}{\gamma}} \sqrt{\frac{\gamma+1}{\gamma-1}} \left[1 - \left(\frac{P_e}{P_c}\right)^{\frac{\gamma-1}{\gamma}} \right]$$
(A-84)

This is solved for each value of Y using a Newton iteration. After the pressure ratio is obtained, the thrust coefficient is obtained:

$$C_{F} = \sqrt{\frac{2\gamma^{2}}{\gamma - 1} \left(\frac{2}{\gamma + 1}\right)^{\frac{\gamma + 1}{\gamma - 1}} \left[1 - \left(\frac{P_{e}}{P_{c}}\right)^{\frac{\gamma - 1}{\gamma}}\right]} + \frac{P_{e}}{P_{c}}$$
(A-85)

Potential chamber explosion pressures are calculated, based on the assumption of well-mixed ideal-gas combustion products uniformly filling the chamber. Three assumptions are used: (1) that a "spike" or "total explosion" will involve all the fuel and oxidizer contained in the chamber, i. e., drops, streams, fans, gas and wall deposits, (2) that a "pop" will involve only gas, and drops (3) that an "adduct explosion" will involve only hydrazinium nitrate on the walls.

h. Subroutine WRITER

This subroutine prints 48 time-varying parameters at each time step and puts large volumes of time-varying data onto tapes 9 and 13.

At each Nth time step, where N is input data value 203, the cumulative amounts of injected fuel and oxidizer are printed, together with the cumulative amounts of the various forms in which oxidizer and fuel have been outflowed or accumulated. If a number has been specified for input data value 11, a large number of chamber parameters will be printed

characterizing the chamber state at that iteration. This print is most useful for clarifying chamber processes or diagnosing apparent program errors.

As soon as the prescribed model time has been calculated, or if the computer clock indicates that the time allocated for the calculations is expiring, end of file is written on tapes 9 and 13 and a summary print is written giving the average performance and contaminant production for the entire period of calculation. If data input value 9 has been given a nonzero value, masses of axially distributed wall deposits of fuel and oxidizer are punched out on cards. These can be used if it is desired to restart calculations at a later time with wall deposits corresponding to the values left at the end of this firing.

i. Subroutine GRAPHS

Subroutine GRAPHS can be entered either at the conclusion of a combustion chamber calculation or for use with data tapes 9 and 13 which hold output data recorded from a previous chamber calculation. If the program is being run from pre-recorded data tapes (indicated by a nonzero value on input data value 10), the information on tape 9 is transferred to disk storage to hasten the ensuing data handling processes and the number of records before end of file are counted for later use in subscripted read and write instructions. Any other necessary data is taken from the data array.

The first plots produced are droplet trajectory plots. Only the droplet specified during the initial combustion chamber calculation can be plotted and changing the droplet trajectory plot input data for re-runs from tape will not give trajectories for different droplets.

Next to be plotted are any time-varying parameters designated by non-zero values in namelist PRTDIR. These plots may be done either with or after the combustion chamber calculations, so long as the data tapes remain intact.

The third set of graphics to be produced is the set of wall-film thickness profiles. If a non-zero value N is given to input Data Number 204, either at the time of a combustion chamber calculation or later for a run from tapes, then the wall-film thickness profile is plotted at every Nth time iteration.

If time intervals are specified, the next output from subroutine GRAPHS is cumulative and time-averaged mean values for performance and contaminant production during the time intervals. The input values required for the MULTRAN subprogram are also produced for the specified time intervals.

The final output from subroutine GRAPHS is a set of snapshots or movie frames showing the combustion chamber contour, the instantaneous location of all droplet groups in the chamber and a wall-film thickness profile which may be given sufficient vertical exaggeration to make it easily visible.

These motion picture graphics give a very vivid description of the combustion chamber processes and make clear, in a very concrete way, many design problems which would go unnoticed if the data were presented in any other way.

j. Utility Subroutines and Functions

Most of the subroutines have available the statement functions PVAPF (TEMP) and PVAPØ (TEMP). These give the equilibrium vapor pressure of fuel or oxidizer in dynes per square centimeter when given the temperature in degrees Kelvin. The expressions used are:

$$P = e^{\left(A + \frac{B}{T - C}\right)} \tag{A-86}$$

The minimum value allowed is either zero or the triple-point pressure, depending on whether FORTRAN variables PFRZF and PFRZØ are set equal to 0.0 or the values held in HOLDF and HOLDO, respectively. The maximum value allowed is the critical pressure.

TEMPF (PR) and TEMPO (PR) give the equilibrium boiling temperature of fuel or oxidizer in degrees Kelvin when the pressure is given in dynes per square centimeter. The expression used is:

$$T = C + \frac{B}{\ln p - A} \tag{A-87}$$

The minimum value permitted can be either 0.0 or the freezing point of the material, depending upon the valve assigned to FORTRAN variables TFRZF and TFRZQ. The maximum values permitted are the critical temperatures.

HVAPF (TEMP) and HVAPO (TEMP) are values for the fuel vapor enthalpy and oxidizer vapor enthalpy in calories per gram as functions of temperature in degrees Kelvin. Contant C_p is assumed, and the zero datum is the solid at zero degrees Kelvin.

$$H = Hc + C_{p}T \tag{A-88}$$

TRIPOT (A, B) gives the length of the vector with an x component of 1.0, a Y component of Tan A, and a Z component of Tan B.

$$L = \sqrt{1 + Tan^2 A + Tan^2 B}$$
 (A-89)

Subroutine SHAPE (X, DCHAM, ACHAM, UPVOL) is given an axial chamber station x and returns the chamber diameter, chamber cross-sectional area, and the upstream volume. Units are centimeters, square centimeters, and cubic centimeters. The chamber contour is linearly interpolated from an input table of 8 stations and diameters.

Functions HCONDF (TEMP) and HCONDØ (TEMP) approximate the enthalpies of liquid or solid phase fuel and liquid or solid phase oxidizer as functions of temperature. Constant C_p is assumed, and the solid phase is assumed to have the same heat capacity as the liquid. The latent heat of fusion is added above the melting point. Values are in calories per gram and degrees Kelvin.

$$T_{freeze} > T$$
 $H = C_{p} T$ (A-90)

$$T \ge T_{\text{freeze}}$$
 $H = C_p T + \Delta H_{\text{fusion}}$ (A-91)

Function REDRHO (TRED) approximates the reduced density of a liquid as a function of reduced temperature along the saturation line, based upon the curves of Hougen and Watson:

$$0.8 > T_r$$
 $\rho_{r(\ell)} = 3.97 - 1.91T_r$ (A-92)

1.0 >
$$T_r$$
 > 0.8 $\rho_{r(\ell)} = 1 + \sqrt{34.7T_r - 25T_r^2 - 9.7}$ (A-93)

$$T_r > 1.0 \qquad \rho_{r(l)} = 1 \qquad (A-94)$$

Function REDROD (TRED) approximates the reduced density difference (reduced density of the liquid minus reduced density of the vapor) as a function of reduced temperature, along the saturation line. Based upon the curves of Hougen and Watson, the law of rectilinear diameters, and some supplementary physical property data:

$$0.8 > T_r$$
 $\rho_d = 3.97 - 1.91 T_r$ (A-95)

1.0 >
$$T_r$$
 > 0.8 ρ_d = -2.21 + 2.21 T_r + 2 $\sqrt{34.7 T_r}$ - 25. T_r^2 -9.7 (A-96)

Function REDVIS (TRED) approximates the reduced viscosity of the liquid as a function of reduced temperature along the saturation line up to the critical temperature, and the reduced viscosity of the gas along the P=0 line above the critical temperature.

$$0.6 > T_r$$
 $\mu_r = 0.55 T_r^{-5.68}$ (A-98)

$$1.0 > T_r \ge 0.6 \quad \mu_r = 2.0 T_r^{-3.15}$$
 (A-99)

$$T_r = 1.0 \mu_r = 1.0 (A-100)$$

$$2.0 > T_{p} \ge 1.0 \quad \mu_{p} = 0.45 T_{p}$$
 (A-101)

Function FANFAC (REN) approximates the Fanning friction factor as a function of Reynolds number.

$$16 > Re$$
 $f = 1.0$ (A-103)

$$2.000 > \text{Re} > 16 \text{ f} = 16 \text{ Re}^{-1.0}$$
 (A-104)

$$Re > 2,000$$
 $f = 0.06028 Re^{-0.2113}$ (A-105)

k. Limitations and Assumptions

The particular processes which have been emphasized in the current program have been dependent upon the immediate interests or problems of the users. Thus many aspects of the start transient and contaminant production are well represented, while pressure wave action in the propellant feed lines, heat transfer in the chamber walls, and sophistication in the dynamics of the combustion gas are conspicuously absent. There are several obvious places where the precision or generality of the present program could be extended if there was a sufficient requirement for the added capability to justify the expense of programming and the longer computing time which goes with greater sophistication.

(1) Feed System

Compressibility of the propellant and the resulting wave action in the lines could be modeled using the method of characteristics. The pressure waves in the lines are important in determining the response of attitude control engines which have very fast valves and extended lines. The two-phase flow in the injector may be important during injector priming and during postrun dribbling under vacuum ambient conditions. Neither wave action nor two-phase flow is modeled at present.

(2) Chamber Gas Dynamics

The axial and transverse variation in gas stoichiometry could be modeled to replace the current assumption of well-mixed gas. This could be done by following discrete "slugs" of chamber gas along stream-tubes and summing the additions of fuel and oxidant vapor from the passage of the "slug" of gas through the droplet array. Doing this, together with a numerical approximation of eddy-diffusive mass exchange between stream-tubes, would permit much more accurate estimation of delivered performance especially in heavily film-cooled engines.

At present the combustion chamber gas state is imperfectly modeled. It is assumed that the chamber gas pressure, temperature, and density at the nozzle throat and at the injector end of the chamber are equal. No consideration is made of the fall in gas temperature associated with the large gas expansions at cutoff or during chugging. To model the expansion or compression of hot thermochemically equilibrated gas with concurrent arbitrary additions of fuel or oxidizer will probably require that equilibrium thermochemistry calculations be performed at each time step. There are current thermochemistry programs which possess sufficient speed and reliability to make this feasible, but it is not warranted for the present uses of the program.

(3) Droplet Trajectories and Secondary Atomization

The present program treats droplet trajectories as being only two-dimensional (axisymmetric). This means that the lateral spreading of the propellant fans is not modeled. Instead, all the propellant droplets are regarded as being initially directed down the centerline of the fan. This leads to predictions of wall deposition more localized than would be the case with a correct description of the fan. Changing the droplet trajectories to be fully three-dimensional would require only minor program changes but would give more precise results, at the expense of longer computing time.

It is well known that when droplets are subjected to sufficiently high relative gas velocities, they undergo secondary atomization, and that this can have an important effect in increasing the combustion efficiency of the chamber. This phenomenon is not presently modeled, but it should be considered if it ever became desirable to better approximate the actual delivered impulse of the engine.

(4) Wall Effects

The heat transfer to the nonwetted portions of the chamber walls is not presently calculated. This is a very interesting possibility, since the axial gas velocity profile, temperature, density, heat capacity, Reynolds number, and even heat transfer coefficients are already calculated for other uses. A simple two-dimensional mesh for unsteady heat transfer into the chamber walls would permit rather sophisticated calculations to be made, such as internal-regenerative cooling, and the effects of thermal soakback on the evaporation of deposited propellant and on subsequent hypergolic starts. Thermal histories for multiple-pulse firings could be calculated, including effects of quenching from propellant leads, from injector dribbling, etc.

(5) Experimental Uncertainties

In addition to the computational limitations of the program, there are some important processes and physical properties which are inadequately known.

The droplet sizes resulting from primary atomization of reactive unlike streams have never been investigated. There have been no experimental determinations of droplet size for the small orifice sizes and extremes of momentum imbalance which can be found in pulse-mode engines. This uncertainty as to correct droplet size is one restriction on the present use of the program. Very little is known about the droplet sizes produced when either an unatomized stream or large droplets impact upon the wall.

When film-coolant fuel flows onto a sufficiently hot segment of chamber wall it will film-bind, be detached and returned to the free volume of the chamber as droplets. The ensuing history of this fuel will depend very strongly on the droplet sizes which are produced; however, this atomization mechanism has never been studied.

The breakup distances for impinging-stream and single-stream atomization have been determined for only a very limited range of liquid velocities, gas densities, and orifice diameters. Further investigations are badly needed for smaller orifice diameters.

Very little is known about the impingement of droplets against the chamber wall. The fraction of impinging droplets which will stick to either a cold chamber wall or to a hot chamber wall is not known.

The effects of droplet size or velocity of approach is not known. The velocity of rebound from the wall and the amount of heat transfer between a bouncing droplet and a hot wall are unknown.

The physical properties of the hydrazinium nitrates and their solutions are completely unknown. Accommodation coefficients for the hydrazines, for nitrogen tetroxide, and for the hydrazinium nitrates are unknown, but because of the highly polar, highly associated nature of the molecules, they are certainly very far from unity.

A. 3 NUMERICAL INTEGRATION METHOD

The numerical solution of sets of differential equations can be accomplished using a number of different techniques. The simplest technique is Euler's method. In this method, the next value for each dependent variable is obtained by linear extrapolation of the present value using the first derivative calculated at the present value. For a time-dependent function:

$$V_{(t + \Delta t)} = V_{(t)} + \frac{dv}{dt_{(t)}} \Delta t \qquad (A-106)$$

where V may be any variable value defined by a differential equation.

Other methods of numerical integration are available in which higherorder derivatives are calculated and used:

$$V_{(t + \Delta t)} = V_{(t)} + \frac{dV}{dt} \Delta t + \frac{1}{2} \frac{d^2V}{dt^2} \Delta t^2 + \dots \frac{1}{n!} \frac{d^nV}{dt^n} \Delta t^n$$
 (A-107)

The use of higher-order derivatives often allows larger values for Δt to be used without introducing excessive error or instability. A penalty must be paid for the use of the higher-order derivatives. In the present calculations, any higher-order derivatives would have to be estimated numerically by taking differences between previous values for the variables. Since 12,000 computer storage locations are required to contain the current values for the droplet array, the requirement that previous values be kept also would drastically increase the already large computer storage requirement. A further argument against the use of a higher-order integrating scheme is that many of the physical processes which are modeled do not have continuous derivatives. The feed system flow rates vary discontinuously when the dribble volumes fill up. The chamber gas properties and droplet burning rates vary discontinuously when the chamber contents ignite. The propellant primary atomization droplet size varies discontinuously when a decrease in chamber pressure or an increase in Weber number causes flashing to commence. When even the first derivative is not continuous, it is pointless to think of approximating higher-order derivatives.

One way that the Euler method can be improved in the solution of certain physical problems is by limiting variables to their physically defined asymptotic values. The time interval chosen for the calculations is based mostly upon the response time of the chamber pressure. If a fuel droplet is injected which is so small that it would burn up completely in less than one computing time interval, then Equation (A-106) would predict that it would burn to a negative diameter in one time interval and contribute more than 100 percent of its mass to the gas phase. Obviously a real burning droplet has an asymptotic value of zero for its diameter and mass. Hence it is appropriate to limit the derivative to a value which will just consume the droplet completely in one computational time interval. Doing this will obviously assign an incorrect value to the droplet burning rate, but will yield a value for chamber pressure at the end of the interval which is correct. Several other derivatives are limited to known asymptotic values. Droplets being accelerated by aerodynamic drag forces will not exceed the local velocity of the gas which is accelerating them, feed system flow rates will not accelerate past the steady-state flow rate corresponding to the current pressure drop. The instructions for limiting the value of the derivatives are all written so as to properly consider flow rates and velocities approaching from either side of the asymptotic value. It should be emphasized that this derivativelimiting procedure is to stabilize the calculations for exceptional conditions, and that most of the calculations are made according to Equation (A-106).

a. Computational Methods for Droplet Arrays

The methods used for the calculation of droplet combustion and trajectories differ according to the generality or restriction of the particular case. Three methods will be described to illustrate the similarities and differences. The first two methods illustrate the historical development of droplet analysis of rocket engine combustion chambers.

(1) Steady-State Chamber with Monodisperse Droplets

The simplest case is that of axially directed monodisperse droplets in a steady chamber. All of the droplets injected as a group at any particular time will behave alike, and the droplet group injected at any particular time will behave the same as any other group introduced at any other time. Thus, knowing the mass, velocity, and position history for any one droplet yields a general solution for the entire combustion chamber. The aerodynamic drag force and evaporation rate for a droplet may be computed, and the successive values for mass, velocity and location may be calculated for a sequence of time steps in a very straightforward manner:

$$M_{(t + \Delta t)} = M_{(t)} - \dot{M}_{(t)} \Delta t$$
 (A-108)

$$V_{(t + \Delta t)} = V_{(t)} + \frac{F_{(t)}}{M_{(t)}} \Delta t$$
 (A-109)

$$X_{(t + \Delta t)} = X_{(t)} + V_{(t)} \Delta t$$
 (A-110)

$$U_{(t + \Delta t)} = U_{(t)} + \frac{1}{\rho A} N\dot{M}_{(t)} \Delta t$$
 (A-111)

Where M is the mass of a droplet, M is the evaporation rate of the droplet, V is the velocity of the droplet, F is the aerodynamic drag force acting on the droplet, X is the axial location of the droplet, U is the combustion gas velocity, N is the number of droplets in the group, ρ is the density of the combustion gas, and A is the cross-sectional area of the chamber. Since U and X are both functions of t, U(X) is defined. This was the method used by Priem to evaluate chamber conditions with uniform droplets (Reference A-57).

(2) Steady-State Chamber with Distributed Droplet Sizes

When the droplets produced by the injector are distributed in initial diameter, the calculations are less straightforward. The small droplets are accelerated by aerodynamic forces much more rapidly than the large droplets, hence, if the droplets are initially together at some particular place and time, they will be separated a short interval of time later. For this reason, the same time interval cannot be used by all the different sized droplet groups to progress from one axial location to the next. In the steadystate chamber, no generality is lost by summing up combustion gas contributions from large, medium and small droplet groups which originated at different times, but which happen to be at the desired axial location in the chamber at the time the gas velocity summation is made. In marching down the chamber to develop the gas velocity profile, an axial distance interval is chosen, and then the time interval required for each different droplet size group to traverse the distance is calculated. The evaporation and acceleration effects for each droplet size group must be evaluated using its own correct time interval.

$$\Delta t_i = \frac{\Delta x}{V_{i(x)}}$$
 (A-112)

$$M_{i(x + \Delta x)} = M_{i(x)} - \dot{M}_{i(x)} \Delta t_{i}$$
 (A-113)

$$V_{i(x + \Delta x)} = V_{i(x)} + \frac{F_{i(x)}}{M_{i(x)}} \Delta t_{i}$$
 (A-114)

$$U_{(x + \Delta x)} = U_{(x)} + \frac{1}{\rho A} \sum_{i} N_{i} M_{i(x)} \Delta_{ti}$$
 (A-115)

The subscript i is used to distinguish between the various groups of different sized droplets. This is the method used by Priem (Reference A-9) and later by Lambiris (Reference A-58) to calculate chamber profiles with distributed droplet sizes. The Dynamic Science approach (Reference A-10) differed only in having groups of oxidizer droplets as well as fuel droplets.

It is apparent that the previous methods depend very strongly upon the assumption of steady conditions in the chamber and cannot be used to calculate the unsteady state. There are other more subtle restrictions, the methods cannot be used for droplets injected in the reverse direction, etc.

(3) Unsteady-State Chamber with Distributed Droplets

4 >

The method used in the present program differs from the earlier methods. Instead of following one or a few droplet groups progressively down the length of the chamber, the entire chamber population of droplets is represented simultaneously and reexamined at each time interval. When this method is used, a larger computer memory is required in order to store the description of the entire droplet population of the chamber, and more computing time is required since all of the droplet groups constituting the entire population are examined at each time interval, however it is now possible to calculate time-varying behavior and it is simple to calculate droplet motion in one, two or three dimensions with no arbitrary restrictions. The mass, velocity and location for each droplet group in the chamber is recalculated at each time interval:

$$M_{i(t + \Delta t)} = M_{i(t)} - \dot{M}_{i(t)} \Delta t \qquad (A-116)$$

$$V_{i(t + \Delta t)} = V_{i(t)} + \frac{F_{i(t)}}{M_{i(t)}} \Delta t \qquad (A-117)$$

$$X_{i(t + \Delta t)} = X_{i(t)} + V_{i(t)} \Delta t$$
 (A-118)

The forces, velocities, and locations may be treated as being one, two, or three dimensional with no difficulty.

The axial gas velocity may be calculated at any axial chamber location, based upon a mass balance on the upstream region. In its present form, this calculation depends upon the assumption that the gas in the chamber is well mixed and has a constant density (i. e., no gradients in composition, temperature, or pressure).

$$U_{x} = \frac{\dot{M}_{x}}{A} \tag{A-119}$$

$$\dot{M}_{x}$$
 = Evaporation rate upstream of X-Accumulation Rate (A-120) Upstream of x

$$\dot{M}_{x} = \sum_{All \text{ groups upstream of } x}^{N_{i}M_{i}} - \frac{V_{x}}{V_{c}} \left[\sum_{All \text{ groups in chamber}}^{N_{i}M_{i}} - \dot{M}_{Nozzle} \right]$$
(A-121)

Where V_X is volume upstream of x and V_C is total chamber volume. In the present program the axial gas velocity is calculated at each of 100 equally spaced intervals and interpolated between these points. In the present program, also, vapor from other sources than droplets is considered; however, the principle remains the same.

(4) Integrating Time Interval

When gas is flowing from a chamber through a sonic throat, the mass flow rate is:

$$\dot{M}_{t} = A_{t} P_{c} \sqrt{\frac{\gamma W m}{RT} \left(\frac{2}{\gamma+1}\right)^{(\gamma+1)/(\gamma-1)}}$$
(A-122)

The mass stored in the chamber is:

$$M = \frac{Pc \ Vc \ Wm}{RT}$$
 (A-123)

The ratio of M/M is often referred to as "gas residence time" and is a rough measure of how fast the chamber pressure can fall by flow through the nozzle. When Equations (A-122) and (A-123) are combined and simplified:

gas residence time =
$$\frac{L_c \frac{A_c}{A_t}}{a_c \sqrt{\left(\frac{2}{Y+1}\right)^{(Y+1)/(Y-1)}}}$$
 (A-124)

When this is evaluated using the values $\gamma = 1.25$, $(A_C/A_t) = 4$, $a_C = 3,600$ ft/sec and $L_C = 1.0$ inch, a value of 0.17 msec is obtained for gas residence time. It is obvious that if normal pressure changes are to be modeled with sufficient accuracy, the integrating time interval must be considerably less than the gas residence time of the particular chamber.

Fuel and oxidizer droplets are introduced at the injector end of the chamber and move downstream with a velocity which depends upon the axial injection velocity and the subsequent aerodynamic drag. To maintain sufficient accuracy in the calculation of the droplet trajectory and mass history, a sufficient number of calculations must be made during the stay time of a droplet in the chamber. Droplet stay time may be estimated as chamber length divided by average droplet axial velocity. A reasonable estimate for axial velocity is 100 ft/sec. Hence, for a chamber 1 inch long the droplet stay time is on the order of 0.8 msec.

In addition to the obvious restraints on maximum integrating time interval, there is a more subtle restraint on minimum time interval. At each time interval the axial gas velocity profile in the chamber is calculated based upon the mass of newly formed combustion gas which must be moved to restore the chamber to a uniform density and pressure. If an extremely short time interval is taken, it is possible that in some unusual cases, gas velocities will be calculated which are higher than are physically possible. This would introduce error in the velocity-sensitive functions such as droplet burning rate, drag, wall film flow and wall film burnoff. If the chamber is viewed as a closed-closed organ pipe a pressure relaxation physically requires one quarter of an acoustic period, i.e.,

$$\Delta t = \frac{Lc}{2 a_C} \tag{A-125}$$

This sets 0.012 msec as a minimum time interval for a combustion chamber 1-inch long.

Since the maximum and minimum permissible time intervals often differ considerably, there is leeway to trade off computer expense against precision of calculation. Most of the computations reported here were done near the maximum acceptable value for time interval to reduce the computation expense.

A. 4 PROGRAM USER'S MANUAL

The <u>TCC</u> program is the first subprogram of the <u>CONTAM</u> computer program. It may be run as a subprogram to CONTAM or as an independent program. The TCC program requires 104,000 words of core and is written in FORTRAN IV.

a. Input

A complete data set consists of 600 data values; however, not all of these are required for every calculation. These are normally punched as four NAMELISTS; PROCED, PRTDIR, RESRT1, and RESRT2.

The 318 values in NAMELIST PROCED describe the engine, propellants, operating conditions, and a number of general instructions or choices of program options. The 80 values in NAMELIST PRTDIR are set to 1.0 or 0.0 to indicate which time-varying parameters should be plotted or not plotted.

The 101 values in NAMELIST RESRT1 are the masses of fuel-derived wall film left in the chamber at the end of a previous firing. The 101 values in NAMELIST RESRT2 are the masses of oxidizer-derived wall film left in the chamber at the end of a previous firing.

The first step performed by the program is to read the input data and then print it out along with descriptive headings. The values in NAMELIST PRTDIR are not printed out as they do not affect the accuracy of the combustion chamber calculations.

This printout makes it easy to review the data to ensure that all the values have been written, punched, and entered correctly. The first program option is to terminate the run at this point to permit a new set of data to be examined thoroughly before time is spent on calculations. The input data to this program are sufficiently complicated that occasional mistakes in entering the data may be expected and careful periodic examination of the contents of the data deck is highly recommended.

Certain of the dein entries are not required for particular calculations and may be omitted (set to zero by the NAMELIST subroutine).

For a computation for which no motion picture is desired, NAMELIST PROCED items 205 through 207 may be omitted. When time-averaged values are not required, items 313 through 318 are omitted. When droplet trajectories are not desired, items 209 through 212 are omitted. When flow-rate override calculations are not desired, 61 to 64, 85 to 88, and 213 to 220 may be omitted.

For a computed vacuum hypergolic ignition, items 33, 34, 35, and 36 are omitted. When an igniter is used, items 33, 37, 38, 39, and 40 are omitted. When sea-level hypergolic ignition is being simulated by assigning a value for ignition delay, items 34, 35, 36, 37, 38, 39, and 40 are omitted. If injector heating of propellant and postfiring dribble are unimportant, the transition volumes (59 and 83) may be omitted. The monopropellant, burning-rate constants (126, 127, 128, 146, 147, 148) are omitted for materials which do not burn in a monopropellant mode.

When multiple pulses are not desired, 237 to 264 are omitted. When computations are being made for engines with less than three pairs of rings, no input values are required for rings which do not exist.

The option flags are all set to zero for "normal" calculations so that they may all be omitted unless an option is specifically desired.

If no time-varying parameter plots are required all of the values in NAMELIST PRTDIR may be omitted. To calculate a first start with a clean engine all values in NAMELISTS RESRT1 and RESRT2 should be omitted.

(1) Namelist Proced

An excellant way to assemble a data set for a new application is by having an old input data print to work from. If none is available, one can be obtained by running TCC with no input data except DDD1 (208) = 1.0, which will produce a "Data Review Only" print with no combustion chamber calculations attempted. The input data will be described in the order that they appear in the program input data print.

DDD1 Subscript	Description	Units
	GENERAL INSTRUCTIONS	
9	Restart punch option (1.0 or 0.0, a nonzero value will cause a deck of cards to be punched at the conclusion of the run. These cards are input values for RESRT1 and RESRT2)	
10	Work from tape option (1.0 or 0.0, a nonzero value will cause the program to omit any combustion chamber calculations and work from prerecorded output data tapes)	
11	Debugging print option (a nonzero value K will cause diagnostic prints to be printed on the Kth time iteration)	
12	No movie tape option (a nonzero value will prevent a movie tape from being generated)	
201.	Model time at which the calculations will be terminated	seconds
202.	Integrating time Interval	seconds

DDD1 Subscript	Description	Units
203.	Number of time intervals between printing of propellant mass disposition prints	
204.	Number of time intervals between plotting of contaminant thickness profiles	
205.	Film vertical exaggeration (wall-film thickness exaggeration for motion picture frames)	
206.	Start movie iteration (time iteration at which the movie starts)	
207.	End movie iteration (time iteration at which the movie ends)	
208.	Data review option	(0.0 or 1.0)
,	(When this flag is set to 1.0 the data are read, certain checks and initializing steps are made and the raw data and some derived information are printed out with headings; however, the chamber calculations are not performed. This makes it possible to check a new data deck for errors before proceeding with expensive calculations.)	
	TIME-AVERAGE PERFORMANCE VALUES	
313.	Time when the first interval starts	seconds
314.	Time when the first interval ends	seconds
315.	Time when the second interval starts	seconds
316.	Time when the second interval ends	seconds
317.	Time when the third interval starts	seconds
318.	Time when the third interval ends	seconds

DDD1 Subscript	Description	Units
	DROPLET TRAJECTORY PLOT	
	Fuel trajectory group	(1.0 through 5.0)
209.	(The chamber trajectory of one selected fuel or droplet group may be plotted each run. The chosen fuel size group is entered.)	
210.	Oxid trajectory group	(1.0 through 5.0)
	(If a trajectory plot is wanted for an oxidizer droplet instead of a fuel droplet, the size group desired should be entered.)	
211.	Trajectory start time	seconds
	(The trajectory is plotted for a fuel or oxidizer droplet which is injected at some chosen time. The time chosen should be entered.)	
212.	Injector Ring	(1.0 through 3.0)
	(The droplet trajectory originates at the injector ring specified.)	
	FLOW-RATE OVERRIDES	
	Flow-rate overrides are provided for propellant feed systems where the internal dimensions are inadequately known, but where experimental values are available for flow rate versus pressure drop. To define the feed system resistances, it is necessary to specify a propellant flow rate, and the associated pressure drop.	
213.	Fuel flow rate	pounds/second
214.	Total fuel feed system pressure drop	psi
	(Tank pressure - chamber pressure. An orifice diameter will be calculated equivalent in resistance to the entire feed system except valve and injector.)	

DDD1 Subscript	Description	Units
215.	Fuel valve pressure drop	psi
	(A valve port area will be calculated.)	
216.	Fuel injector pressure drop	psi
61	Percent of fuel through ring 1	
	(Percentage flow, fractional flow, or actual flow in same units as 62 and 63. Discharge coefficients will be calculated.)	
62	Percent of fuel through ring 2	
63	Percent of fuel through ring 3	
217	Oxidizer flow rate	pounds/second
218	Total oxidizer feed system pressure drop (tank pressure - chamber pressure)	psi
219	Oxid valve pressure drop	psi
220	Oxid injector pressure drop	psi
85	Percent of oxidizer through ring 1	
	(Percentage, fractional, or actual flows in units compatible with 86 and 87.)	
86	Percent of oxidizer through ring 2	
87	Percent of oxidizer through ring 3	
	OPERATING CONDITIONS	
13	Fuel tank pressure	psi
14	Fuel tank temperature	degrees K
15	Fuel tank pressure-rate of change	psi/sec
16	External pressure	psi
17	Oxidizer tank pressure	psi
18	Oxidizer tank temperature	degrees K

DDD1 Subscript	Description	<u>Units</u>
19	Oxidizer tank pressure-rate of change	psi/sec
21	Injector initial temperature	degrees K
22	Throat initial temperature	degrees K
25	Injector maximum temperature (equilibrium temperature, firing)	degrees K
26	Half-rise time (half-time for injector heatup)	seconds
27	Injector minimum temperature (equilibrium temperature, nonfiring)	degrees K
28	Half-fall time (half-time for injector cooling)	seconds
29	Throat maximum temperature (equilibrium temperature, firing)	degrees K
30	Half-rise time (half-time for throat heatup)	seconds
31	Throat minimum temperature (equilibrium temperature, nonfiring)	degrees K
32	Half-fall time (half-time for throat cooling)	seconds
	IGNITION DESCRIPTION	
33.	Assigned ignition delay	seconds
	(For sea-level hypergolic starts an experimental value for ignition delay time is entered)	
34.	Igniter port location	inches
	(If an external igniter is being used, the location, downstream of the injector, where the hot gases enter the chamber)	
35.	Igniter fuel flow rate	pounds/second
	(Flow rate of fuel through the igniter)	

DDD1 Subscript	Description	Units
36.	Igniter oxidizer flow rate	pounds/second
	(Flow rate of oxidizer through the igniter)	
37.	Activation energy	(calories/mole)
	(Activation energy of the global gas- phase ignition reaction)	
38.	Frequency factor X Q	(cubic cm/mole/sec) X
	(Molar collision frequency inultiplied by the heat of reaction of the initiating reaction)	(calories/mole)
3 9.	Perfect mixing option	(0.0 or 1.0)
	(If this flag is set to 1.0, ignition will be calculated only for the well-mixed free-stream gases, ignoring the pos- sibility of ignition in the boundary layers around droplets or on the wetted chamber walls)	
40.	No axial mixing option	(0.0 or 1.0)
	(If this is marked with a 1.0, ignition will be calculated only for the newly flashed fuel and oxidizer vapors presuming no exial mixing of vapor and ignoring ignition in the boundary layer)	
	FUEL FEED SYSTEM	
41	Fuel line length	inches
42	Fuel line diameter (I.D.)	inches
43	Restrictor diameter	inches
	(The Vena Contracta diameter of a real restrictor, or a diameter equivalent to the resistance of a complex feed system)	
44.	Fuel Venturi diameter (Cavitating Venturi throat diameter)	inches

DDD1 Subscript	Description	Units
45	Fuel valve port area (port area * discharge coefficient)	square inches
46	Fuel check valve option	(0.0 or 1.0)
	(If nonzero, no reverse flow permitted in the fuel feed line)	
47	Fuel valve opening time	seconds
	(Mechanical ramp duration, not electrical time)	
48	Fuel valve closing time	seconds
	(Mechanical ramp duration, not electrical time)	
57.	Initial void volume	cubic inches
	(Initial empty volume in the fuel feed system)	
59.	Transition volume	cubic inches
	(Volume of fuel that flows while the injection temperature decreases from the injector temperature to the tank temperature.) (See Figure A-3.)	
60.	Dribble Volume	cubic inches
	(Volume between the closed valve and the injector face)	
	OXIDIZER FEED SYSTEM	
	(65-72 and 81-84 are identical to entries 41-48 and 57-60 except that they describe the oxidizer feed system instead of the fuel feed system)	
	ATOMIZATION PARAMETERS	
89.	Fuel drop factor	
	(Correction factor for fuel droplet median diameter - not needed for tur- bulent flow through circular orifices flowing full)	

DDD1 Subscript	Description	Units
90.	Oxidizer drop factor	
	(Correction factor for oxidizer droplet median diameter)	
91.	Fuel fan length	orifice diameters
	(Distance between the impingement point and the point at which the fuel is atomized when the stream turns 45 degrees at impingement)	
92.	Oxidizer fan length	orifice diameters
93.	Hold at triple point option	(0.0 or 1.0)
	(When this flag is set to 1.0 a propel- lant stream flashing in a low pressure environment will not freeze, but will stop as triple-point liquid)	
94.	No initial dribble option	(0.0 or 1.0)
	(When this flag is set to 1.0, no liquid will be injected until the void volume is filled, even when the injector-temperature vapor pressure is higher than the chamber pressure)	
95.	Flash cone angle	degrees
	(This is the included apex angle of the cone of spray formed by a flashing stream)	
96.	Showerhead length/orifice diameter ratio	orifice diameters
	(The distance, in orifice diameters, that an unimpinged stream travels from the injection point to atomize completely)	
97-101.	Drop size 1 - drop size 5	
	(These are the diameter ratios of the droplet groups to the mass-median droplet)	

DDD1 Subscript	Description	Units
102.	No wall breakup option	(0.0 or 1.0)
	(When this flag is set to 1.0 an unatomized stream hitting the wall is not atomized by the impact)	
103.	Drop restitution coefficient	
	(The normal-velocity ratio for a drop- let which hits the wall and bounces off)	
104.	Fraction sticking	
	(The fraction of the droplets hitting the wall which stick to the wall instead of bouncing)	
105.	No fuel flash option	(0.0 or 1.0)
	(When this flag is set to 1.0, the injected fuel stream will not flash even when it is highly superheated)	
106.	No oxidizer flash option	(0.0 or 1.0)
	(When this flag is set to 1.0, the injected oxidizer stream will not flash even when it is highly superheated)	
107	No entrainment flag	(0.0 or 1.0)
	(A nonzero value will prevent any entrainment of wall film)	
108	Delete droplet means	(0.0 or 1.0)
	(A nonzero value will prevent calculation of the chamber droplet size means at every time iteration)	
	FUEL PROPERTIES	
109.	Fuel boiling point	degrees K
	(The normal boiling point of the fuel)	
110.	Fuel freezing point	degrees K
111.	Fuel critical temperature	degrees K
112.	Fuel critical pressure	psia

DDD1 Subscript	Description	<u>Units</u>
113.	Fuel vapor specific heat	calories/gram/ oK
	(Fuel vapor specific heat at a temperature midway between the combustion gas and the droplet surface)	•
114.	Fuel liquid specific heat	calories/gram/
	(Liquid fuel specific heat at the injection temperature)	I.
115.	Blank	
116.	Fuel molecular weight	grams/gram- mole
	(Molecular weight of fuel in the vapor phase)	more
117.	Fuel latent heat of vaporization	calories/gram
	(Latent heat of vaporization of fuel at the normal boiling point)	
118.	Fuel latent heat of fusion	calories/gram
	(Latent heat of fusion of the fuel at the freezing point)	
119.	Fuel liquid thermal conductivity	calories/sec/cm/ ^o K
120.	Fuel accommodation coefficient	
	(Coefficient for Langmuir-Knudsen vacuum evaporation rate)	
121.	Fuel reference temperature	degrees K
	(Temperature at which values for density, viscosity, and surface tension are given)	
122.	Fuel density	grams/cm ³
	(Density of liquid fuel at the reference temperature)	
123.	Fuel viscosity	poises
	(Viscosity of liquid fuel at the reference temperature)	

DDD1 Subscript	Description	<u>Units</u>
124	Fuel surface tension	dynes/cm
	(Surface tension of fuel at the reference temperature)	
125.	Fuel burning rate coefficient from droplet burning experiments	cm ² /sec
	(-dD ² /dt for a burning fuel droplet in pure oxidizer vapor)	
126.	Fuel monopropellant burning rate intercept	cm/sec
127.	Fuel monopropellant burning rate coefficient	cm/sec/(psia) ⁿ
128.	Fuel monopropellant burning rate exponent	
	(Monopropellant burning rate values from liquid-strand experiments correlated by $r = A + BP^n$ where r is in centimeters per second and P is in psia)	
	OXIDIZER PROPERTIES	
129-148	Identical to 109-128 except that they describe the oxidizer instead of the fuel	
	PRODUCT PROPERTIES	
	EQUILIBRIUM GAS TEMPERATURE	
149-159	Are the thermochemical equilibrium combustion product temperatures corresponding to fuel fractions of 0, 0.1, 0.2, 1.0	oK
	EQUILIBRIUM GAS MOLECULAR WEIGHT	
161-171	Are the mean molecular weights of the combustion products in thermochemical equilibrium at fuel fractions 0, 0.1, 0.2, 1.0	

DDD1 Subscript	Description	Units
	EQUILIBRIUM GAS GAMMA	
173-183	Are the values for frozen gamma of the equilibrium combustion products at fuel fractions 0, 0.1, 0.2, 1.0	
	THRUST COEFFICIENT TABLE	
	EQUILIBRIUM THRUST COEFFICIENT	
221-231	Vacuum thrust coefficients are entered for combustion product gases having fuel fractions of 0., 0.1, 0.2, 1.0	
232.	Nozzle expansion area ratio	
	ADDUCT PROPERTIES	
185.	Density of the contaminant material produced on the chamber wall	gm/cm ³
186.	Specific heat of the gases produced by the evaporation or pyrolysis of the wall-film material	cal/gm/ ^o K
187.	Latent heat of evaporation or heat of ablation of the wall-film material	cal/cm
188.	Surface temperature of the wall-film material during evaporation or pyrolysis	degrees K
+1	CONTAMINANT VISCOSITY	
189-199	Viscosity of the wall-film material at the temperature to be expected on the wall at fuel fractions 0, 0.1, 0.2, 1.0	
	MULTIPLE PULSE VALVE TIMING	
233-264	Valve timing for the first through eighth pulses of engine operation	
233	Fuel valve, time of first motion opening on first pulse	seconds
234.	Oxidizer valve, time of first motion opening on first pulse	seconds

DDD1 Subscript	Description	Units
234.	Fuel Valve, time of first motion closing on first pulse	seconds
236.	Oxidizer valve, time of first motion closing on first pulse	seconds
	MULTI-RING INJECTOR	
	FIRST RING	
	FUEL HOLES	
49.	Fuel hole diameter	inches
	(Injector hole size)	
50.	Fuel hole length	inches
	(Injector hole length)	
51.	Axial location	inches
	(Axial location of fuel injection point)	
52.	Radial location	inches
	(Radial location of fuel injection point)	
53.	Injection angle	degrees
	(Outward angles counted as positive)	
54.	Discharge coefficient	
55.	Number of fuel holes in this ring	
56.	Transverse angle	degrees
	(Used to impart swirl to film-coolant. Because of the two-dimensional drop-let calculations, transverse angle will not cause the propellant to approach the combustion chamber wall)	
	OXIDIZER HOLES	
73-80	Identical to entries 49-56 except that they describe the oxidizer holes instead of the fuel holes	

DDD1 Subscript	Description	Units
	SECOND RING	
265-272	Describe the second ring fuel holes	
273-280	Describe the second ring oxidizer holes	
	THIRD RING	
281-288	Describe the third ring fuel holes	
289-296	Describe the third ring oxidizer holes	
	COMBUSTION CHAMBER PROFILE	
297.	Axial location of injector face	inches
	(usually set to zero)	
298.	Injector face diameter	inches
299.	Axial location 2	inches
300.	Chamber diameter 2	inches
301-310	Pairs of axial stations and diameters describing the chamber contour	inches
311	Axial location of the throat plane	inches
312	Throat diameter	inches

(2) Namelist PRTDIR

Namelist PRTDIR directs the program to produce graphics of the time-varying parameters which are desired. The default value for the members at this array is zero, and graphics are produced only for members which are given non-zero values. The graphics which can be produced are:

DDD 2 Subscript

Variable to be plotted

- 1. Chamber pressure
- 2. Fuel valve trace (fraction of wide-open port area)
- 3. Oxidizer valve trace (fraction of wide-open port area)
- 4. Fuel flow rate leaving tank

Variable to be plotted

DDD 2 Subscript

- 5. Oxidizer flow rate leaving tank
- 6. Total propellant flow rate leaving tanks
- 7. Fuel injection rate
- 8. Oxidizer injection rate
- 9. Fuel (streams, fans and droplets) mass in chamber
- 10. Oxidizer (streams, fans and droplets) mass in chamber
- 11. Propellant (streams, fans and droplets) mass in chamber
- 12. Fuel mass on chamber wall
- 13. Oxidizer mass on chamber wall
- 14. Propellant mass on chamber wall
- 15. Gas mass in chamber
- 16. Mass fraction of the chamber gas which is fuel-derived
- 17. Chamber temperature (gas phase)
- 18. Fuel evaporation rate
- 19. Oxidizer evaporation rate
- 20. Propellant evaporation rate
- 21. Gas outflow rate
- 22. Fuel droplet outflow rate
- 23. Oxidizer droplet outflow rate
- 24. Total droplet outflow rate
- 25. Fuel film outflow rate
- 26. Oxidizer film outflow rate
- 27. Total film outflow rate
- 28. Mass fraction of total outflow which is gas phase
- 29. All chamber contents total explosion pressure capability
- 30. Mass of unatomized fuel streams and fans in chamber

DDD 2 Subscript

Variable to be plotted

- 31. Mass of unatomized oxidizer streams and fans in chamber
- 32. Mass of fuel droplets in chamber
- 33. Mass of oxidizer droplets in chamber
- 34. Gas and droplets only total explosion pressure capability
- 35. Chamber wall adduct only chamber explosion over-pressure capability
- 36. D30 of all fuel droplets in chamber
- 37. D31 of all fuel droplets in chamber
- 38. D32 of all fuel droplets in chamber
- 39. D30 of all oxidizer droplets in chamber
- 40. D31 of all oxidizer droplets in chamber
- 41. D32 of all oxidizer droplets in chamber
- 42. Fuel injector void volume
- 43. Oxidizer injector void volume
- 44. Thrust

(3) Namelists RESRT 1 and RESRT 2

These Namelists contain the amounts of fuel and oxidizer left deposited on the chamber wall at the conclusion of a previously computed engine firing. These Namelists are punched by the program at the end of a calculation if the "Punch for restart" option has been given a non-zero value.

b. Program Output

(1) Input Print

The first print which is produced by the program is a recapitulation of the program input

(2) Propellant Property Print, Derived Feed System Values and Chamber Description

The second print produced is a table of calculated propellant properties versus temperature. These values can be compared with known experimental values to assure that the program physical properties subroutines are giving an adequate representation of the true values. The values printed are liquid enthalpy, vapor enthalpy, vapor pressure, liquid density, liquid viscosity, and surface tension. These are given for evenly spaced intervals of reduced temperature, with the corresponding absolute temperature also given. The units are calories/gram, psia, grams/cc, poise, dyne/cm, and degree3 K.

If the flow overrides have been used, the values derived for effective valve port areas, orifice discharge coefficients and equivalent feed system orifice diameter are printed in square inches and inches.

The effective feed system component areas are given in square inches. The relative resistances to flow of the "line" (everything except valve and injector) and the injector are given in units of inches $^{-4}$ ($1/A^2$). The inertial aspect of the feed system geometry is expressed in inches $^{-1}$ (L/A). The effective injection area (geometrical area * discharge coefficient) is given for each injector ring, and the percentage of total fuel and oxidizer flow through each ring is given. The axial and radial location of the impingement point is given for each ring in inches.

The combustion chamber is categorized by giving the axial location, diameter, cross-sectional area, upstream volume, mass of fuel on the wall, mass of oxidizer on the wall, and wall temperature for each of 100 equally spaced axial locations. The units are inches, square inches, cubic inches, grams, and degrees Kelvin.

(3) Print of Time-Varying Rocket System Parameters

At each integrating time interval, values are printed for 48 of the more important system variables. This print consists of six lines containing eight values per line.

The first line contains the model time in milliseconds, the chamber pressure in psia, the ignition state of the chamber (1.0 for ignited, 0.0 for unignited), the mass fraction of the gas or vapor in the chamber which is fuel-derived, the chamber temperature in degrees K, the mean molecular weight of the gases or vapors in the chamber, the specific heat ratio for the gases or vapors in the chamber and the vacuum thrust coefficient for the gas or vapor phase flowing from the chamber.

The second line contains the fuel and oxidizer injection rates in pounds per second, and the fuel and oxidizer feed-line flow rates in pounds per second. The injection flowrate and feed-line flowrate will differ when the injector is dribbling into a vacuum after the propellant valves close, or when a partially empty dribble volume is being filled before the injector primes. The void volumes on fuel and oxidizer sides of the injector are given in cubic inches. The dribble injection rate will decrease to zero when the dribble volume is emptied. The fuel and oxidizer injection temperatures are given in degrees Kelvin. The injection temperatures are equal to the injector temperature, until a volume of propellant has flowed out of the tank equal to the dribble volume, then the temperature linearly decreases to the tank temperature, with the decrease spread out over the prescribed transition volume.

The third line contains the evaporation rates of the fuel drop ensemble and the oxidizer drop ensemble in pounds per second. The fuel flash rate and oxidizer flash rate are given in pounds per second of vapor produced by the vacuum flashing of the injected streams. The rates of propellant burnoff from the wall is given for fuel and oxidizer in pounds per second. The heat for wall burnoff comes from heat transferred from the

chamber gases. The rates of vacuum evaporation from the wall are given for fuel and oxidizer in pounds per second. The heat for vacuum evaporation is transferred from the chamber wall through the propellant film to the evaporating surface.

The fourth line contains the mass of fuel-derived combustion gas or vapor in the chamber in pounds. The mass of oxidizer-derived gas or vapor is given similarily. The masses of fuel droplets and oxidizer droplets in the chamber are given in pounds. The masses of injected, but not yet atomized fuel and oxidizer are given under the headings Fuel Streams and Oxidizer Streams. The mass of fuel and oxidizer deposited on the walls are the final entries of the fourth line.

The first and second entries of the fifth line give the mass rate of efflux for fuel and oxidizer-derived vapors or combustion-product gases in pounds per second, the third and fourth entries are the mass rate of ejection through the throat for incompletely burned fuel drops and oxidizer drops in pounds per second. The rate at which propellant film on the wall is being flowed through the throat is given next in pounds per second. "Gas Fraction" is the instantaneous value for the fraction of the total mass efflux which is in the gas phase. The final entry on this line is thrust, in pounds force. The thrust includes the axial momentum rate of the expelled droplets, and the thrust from the gas phase products.

The sixth line starts with the sauter mean droplet diameter for all the fuel and oxidizer spray contained in the chamber this time interval. The next two entries give fuel and oxidizer stream injection velocities. The maximum explosion pressure is the pressure which would be attained if all the chamber contents (gas, streams, fans, droplets, and wall film) were well mixed and converted to equilibrium combustion products. The mass out of tank is total mass of propellant flowed from the propellant tanks from the start of the run, in pounds. The integral PC x DT is the area under the curve of chamber-pressure-versus time, in psia x seconds. Total impulse is the time integral of thrust in pound-force x seconds.

(4) Summary Prints and Special Messages

Other prints which may occur interspersed with the normal six line prints are summary prints and special messages. The most frequent special messages specify when ignition or extinguishment occurs in the combustion chamber, or when the fuel or oxidizer manifolds fill. Other special messages indicate minor errors, such as failure to converge to within 2 percent of the correct surface temperature during vacuum evaporation, or catastrophic errors such as calculation of a negative chamber pressure, or filling up the droplet array locations.

The propellant disposition summary, whose frequency is specified in data item 203 consists of seven lines of numbers. The first four lines give a mass balance for all fuel and oxidizer injected up to the present time. The masses, in grams, are given for total injected fuel, oxidizer and sum of fuel plus oxidizer, and the amounts expelled as gas, as droplets and as wall film. Also given are the masses currently retained as gas, as droplets and as wall film. The last five lines gives the numbers of fuel drops and

oxidizer drops injected and ejected up to the present time. The summation of droplet diameters, diameters squared, diameters cubed and axial momentum is also given, which makes it possible to calculate the mean diameters and mean axial velocities of expelled droplets for the interval between any two summary prints. If KBUG has been given a nonzero value, large numbers of diagnostic data will be printed on the KBUGTH iteration. These values will be of interest for trouble-shooting only.

(5) Final Performance Summary

After the last of the time-steps is printed, a performance summary is printed. The mass flowed out of tank is given, the mass flowed through the injector is given and the mass flowed through the nozzle is given. These values are not generally the same, because of filling or voiding of the injector dribble volumes, and because of increase or depletion of propellant mass stored in the combustion chamber in the forms of gas, droplets or streams, and wall film. The mixture ratio, C-Star, and specific impulse are calculated on each of these propellant mass bases. The pressure time integral for the entire pulse and the total impulse are also printed. The final print also gives the disposition of injected propellant for the entire firing. The fractions of injected fuel, oxidizer and total propellant expelled in the form of gas, unburned vapor, droplets and as wall film are given, and the fraction retained in the chamber as gas, droplets, and wall film are given. The mean diameters for injected and ejected fuel and oxidizer are given in microns. The mean axial and radial velocities are given for the ejected droplets in feet per second. The calculated size distribution for injected and ejected fuel and oxidizer is given. When this is plotted, it shows signs of the five-group approximation for injected droplet size, which is a computational artifact; however, also apparent are the real effects of flashing versus impinging stream atomization, the preferential burning or small droplets, the segregation by drag versus momentum effects, etc.

(6) Final Chamber Condition

The axial distribution of fuel wall film, oxidizer wall film, and temperature are given to show the input which would be used for a restart of the chamber for the next pulse. This print also reveals the potential for chamber explosion or contamination based on the wall-film propellant remaining in the chamber after cutoff.

(7) Time-Averaged Interval Performance Values

Up to three prints can be called per run, giving aggregate values and rates for prescribed intervals of time. These intervals are often chosen to represent the start transient, steady-state operation, and the cutoff transient. Having values for total impulse and propellant consumption for the transients and for thrust and propellant consumption rate during steady state makes it possible to estimate performance for pulses having similar initial conditions but different pulse duration, simply by summing up the total impulse and propellant consumption for the two transients together with appropriate amounts of steady-state operation. The average performance or

contamination production for different pulse-widths or pulse-trains may be estimated in a similar way. Input values required for MULTRAN are printed for these same time-averaged interals.

(8) Computer Graphics

The computer graphics include the trajectory of the specified fuel or oxidizer drop, up to 44 time-varying system parameters plotted versus time, and a specified number of instantaneous profile plots of wall deposit thickness versus chamber length. The wall deposit plots may be eliminated by setting input item 204 to zero, motion picture frames are produced for the interval specified by input items 206 and 207. The desired time-varying system parameters are produced, by giving nonzero values to the corresponding subscripted flags in NAMELIST PRTDIR.

A. 5 SAMPLE CASE

Table A-III consists of a sample case for the TCC program. Illustrated in the following order are; a listing of the control cards, and a representative sample of the prints and graphics produced.

The printer output consists of the input data print, and representative samples of each of the types of output prints. A sampling of computer graphics includes a droplet trajectory plot, chamber pressure, fuel injection rate, oxidizer injection rate, one wall-film thickness plot, and one frame of motion picture, showing the distribution of propellant droplets and wall-film in the combustion chamber.

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### OPEN OXID VALVE OREN FUEL VALVE CLOSE ####################################	TEIRD PULSE TIRING	1
VALVE OPEN OXID VALVE OFEN FUEL VALVE CLOSE 4.00000 (274) FIFTH PULSE TIMING VALVE OPEN OXID VALVE GIEN FUEL VALVE CLOSE 6.00000 (250) 0.000000 (251) 0.00000 SIXTH PULSE TIMING VALVE OPEN OXID VALVE OFEN FUEL VALVE CLOSE 6.00000 (250) 0.000000 (299) 0.00000 SEVENTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE TIMING FIGHTH PULSE FIGHTH PULSE FIGHTH PULSE FIGHTH PULSE FIGHTH PULSE FIGHTH PULSE FIGHTH PULSE FIGHTH PULSE FIGHTH PULSE FIGHTH PULSE FIGHTH PULSE FIGHTH PULSE FIGHTH PULSE FIGHTH PULSE FIGHTH PULSE FIGHTH PULSE FIGHTH PULSE FIGHTH PULSE FIGHTH PULSE FIGHTH PULSE FIGHTH PULSE FIGHTH PULSE FIGHTH PULSE FIGHTH PULSE FIGHTH PULSE FIGHTH PULSE FIGHTH PULSE FIGHTH PULSE FI	D VALVE DEEN FUEL VALVE CLOSE DEED WAL 0.000000 (243) 0.000000 (244)	1
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### OPEN OXID VALVE CIEN FUEL VALVE CLOSE OXID ####################################	FIFTH PULSE TIMING	
SIXTH PULSE TIMING 0.00000 (254) 0.00000 (295) 0.00000 (SEVENTH PULSE TIMING VALVE OPEN OXID VALVE OF EN FUEL VALVE CLOSE 0.00000 (250) 0.00000 (205) 0.00000 (6.00000 (202) 0.00000 (203) 0.00000 (OXID VALVE GILN FUEL VALVE CLOSE ORID (250) 0.000000 (251) 0.000000 (252)	
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	COMPUSTION CHANNER PROFILE	
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AXIAL LOG. 3 CHAP	AXIAL LOG, 4 CHAMBE	R 514M, 4
AXIAL LDC. 9 CHAI	HEP DIAH; 5 AXIAL LGG, 6 CHAMBE	R DIAH, 6
AXIAL LOC. 7 CHAI	THEF DIAK, 7 THEOAT PLANE	THROAT DIAM.
INPUT UNITS ARE I PROPELLANT PROFE	ND DEGREES RELVIN. Polse, EVNE/GH,	

. 300000	178,260000	122,990000	345,809800	345,009000 .000001 .994670		,200413	76,710000
,325000	193.09000	133,204500	359,784750	.000026	. 980697	.127107	72,404632
35000	207,900000	145,485000	374,940300	100032	875990	.003407	40,44000
375000	222,750000	221,197500	389,336250	.002316	. 992733	.056426	64,973288
.40000	237,600000	231,444000	404,112000	.011017	.938751	.030139	60,069209
425,000	252,450000	241,600500	448,887780	.046309	4,684278	972380	98.978.88
. +50000	267,360080	251, 937000	433,063900	.140632	.910788	.020032	93,930744
475000	202.150000	262,183505	448,439250	,405244	.00000	.014739	90,00460
200000	287,000000	272,430000	463,215000	.070306	260500	110110	47,600346
,525000	311.050000	282,676500	477,990750	6.000.5	274090	*******	44,66130
.950000	326,700000	292,923000	492,766500	4,119665	.054861	.006400	41,055206
375969	341,550000	303,16050	567,54225	2,546824	940079	604076	80.101.08
. 600000	356.400000	313,416000	522,310000	12.00050	. 62659	.003989	36,641994
,625000	371,250000	323,662500	537,093750	21,203941	.012016	. 003433	34,225506
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.675000	400.950000	344,199900	966,645250	40.557001	.764053	.002694	20,753601
.700000	415.600000	354,402000	581,421000	72.756247	.770971	.002402	27,669749
225000	430,450000	364, 648500	584.194750	100.799827	28488	.002151	25,735089
.750000	445,500000	374,895469	610,972500	137.866477	.743007	.001933	23,00500
. 775000	460.350000	305,14150.	625,74825u	104,149020	.729026	.001743	22,136009
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.025006	490.050000	405,634900	052.299756	304.005004	.697662	.061432	18,443619
. 050000	564.930000	415.861000	670,07550	290.000007	.676221	.001303	16,150055
875000	518,750000	424, 427500	484,891250	484,656474	464035	181100	43,8774,2
. *************************************	534,600000	436.374006	609,027000	394,000405	.624088	.00100.	10,011953
,025000	549,450000	446.620501	714,402750	716,959165	.586626	066060.	6,619689
95660	564,300000	454,867000	228,1785ca	646.31507	.536334	95000	3.4634.4
.975000	579,150000	467,113560	743,994250	1518,780321	.470167	.0000.	1,029387
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.30060	129,300000	46,54800	194,959400	000000		.36990	76,696221
,325000	140.075000	50.427000	198,170350	. 00000	1.039170	,232006	72,473074
350800	150,850000	54,304000	201,381300	100000	1,062067	900887	50000100
.375000	161.625000	56,165000	204,592250	. 000012	1.700736	,102920	64,594710
. 400060	172,400000	62.064000	207,603200	.000133	1.760515	.071334	66,647856
425000	183,175080	45, 84300:	281,01415:	389080	1,734384	165030	87,36868
.450000	193,950000	69.822000	214,225100	.003206	1.768073	.036536	53,919301
475000	264,725000	73,7n160c	217,436050	. 012672	1,661052	,026677	90,480093
20000	215,500000	27.560000	220,447880	427244	1.685631	020084	47,50275
.525606	226,275000	61.499000	223,857950	.129105	1.629410	.019223	44,648992
, 550000	237.056000	65.336000	227,000900	339845	1.663189	.011686	41,043223
975980	247, 825080	88,237000	236,279850	111154	1,93444	***********	30,472678
. 400000	256.00000	93.096000	233,490800	1.774363	1,990747	.067130	36,631102
. 425000	269.375000	136.175000	236,701750	3,614636	1.924526	192900	34,219709
450000	280,150000	140.054005	238,912700	4,914437	1,448385	115500	32,696724
.675600	290,925000	143,933606	243,123090	12,519297	1.472084	.004913	29,745171
.760800	301,700000	147,812000	246,334000	21,999537	1,449863	.004381	27,661828
725000	312,475000	154.461000	240,545350	36,746346	2742443	100200	-86,767.98
.750000	323,250000	159,578000	252,756560	54.052766	1.363421	.003529	83,67000
.779006	334,025000	159,449000	255,967450	67,511463	1.367200	.003160	22,131669
1	344,400000	143, 35666	880,178400	-136,200800	4.300030	,00000	-20,408070
. 429660	355,575000	167.207000	262,309350	100,900223	1,300419	.082611	10,436334
. 050000	366.350080	171.086000	265,600300	260.400399	1.271023	.008377	16,145430
472600	377,625000	224,00000	846,444,846	178,44,868	1.004044	095600	\$308484E3
	387.90060	170.844000	272,022200	900.479868	1.170403	.001909	10,00000
,92900	396,675060	182.723000	275,233150	478,929785	1.186147	120100	0,617793
1	400-48000		878,444600	**********	2000007	100000	8440848
.07900	428,225000	190.401000	261,695650	1192,636631	. 881741	.001543	1,629992
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8	97829000	2.69426971	2,544412	02741230	0.0000000	000000000	294.0000000
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1	46.0000		1.000006-02		,0	CXID STREAMS			PARS DUT DE TANK	•	645 RETAINED 9.00000000 . C0000003	0. 0. 0.		PGL WT.	CXID VOID VOL	CXID MALL BURN	CXID STREAMS	CXID FILM NA	PASS DUT DE
	294,000		1,000000E-02	CHEA MALL MILES	0.	FLEL SPREAMS	- 17		MAX, EXPLOSION	•	MALL EXPELLED 0.00000000 0.00000000	UM D SOLABED S		CHANBER TENF K 247, 998	FUEL VCID VOL	₹]	FUEL STREAMS	FUEL FILM RATE	MAK, EXPLOSION
	0	24.0 M. 10.10	0	CATE CLASH BATE FIRE HALL		DAID DROPLETS	DATE DRAM BASE	.0	OXIC INTO VEL	•	DNOPS EXPELLED 0.00000000 0.000000000			FUEL FRACTION	OXID FLOW RATE	8	OXID DRCPLETS	OXID DROF RATE	OXID INJ. VEL.
	•	F1.51 F1.04 8.75	Ì	FLEL FLASH BATE		FUEL DROPLETS.	FIRE DROP DATE			•	GAS EXPELLED 0.00000000 0.000000000		0.	IGNITION	FUEL FLOW RATE	FUEL FLASH RATE	FUEL DPOPLETS	Jet.	FUEL 1NJ. VEL.
3443	1,300000E=06	CXID 1114 KATE		OXID DOOD BULL		7, 34341E-11	OXID GAS LEGILY		ON TO DROP UNE	•	17.1ECTFD TASP 0.0000000 0.00000000	INJECTED EJECTED	EJECTED	CHAMBER PRESE	OXID INJ KATE	OXID DRCP BUFA	DAID GAS HASE	6	OXID DROP D32
		FUEL IN GATE		FUEL DROF BLEN	•	FUEL GAS LASS	FUEL GAS CERTUX		FUEL DROP D32	•	FUEL GRAMS OXID BRAKS TOTAL GRAMS	FUEL CRUPS	מנאס	71ME HILLISEC .20080	FUEL INJ RATE	FUEL DROF BURN	FUEL BAS HASS	FUEL GAS EFFLUX	FUEL BROP 032

	ı						
FUEL DROP BURN 8.	N OXID DROP BUFN	PUEL PLASH NATE 1.5004676-03	0XID FLASH RATE 1,3651946-02	FUEL WALL BURN 0.	CXID WALL BURN	PUEL MALL BYAP	OXID WALL BYAP
FUEL 6AS PASS 5.172141E-07	0X12 9AS HASE 7	FUEL DRCPLFTS 1.7352646-06	DKID DROPLETS 3,198002E-09	FUEL STREAMS	CKID STREAMS	FUEL ON WALL	OXID ON WALL
TUEL GAS EFFLUX 1.099060E-04	UX OXID GAS EFFLUX 4 6.900836E-64	FUEL BROF RATE	ONID DROP RATE	FUEL FILM RATE	CXID FILM RATE	SAS FRACTION V.	VAC THRUST LOF
FUEL DADP D32 55,4369	0X1D DROP DUP 34.480	fuel 11.3. vel.	OXID INJ. VEL. 13.8314	MAX, EXPLOSION 2,98693	7.922416E#8	1.3021298-09	107AL IMPULSE 2,9615766-85
TIME MILLIBEC	CHAMBER PRESS	1011101	FUEL FRACTION	CHANGE TENE	11 104	Avnav	ee nee
•••••	9,037520E=02	•		7.004	40.0344	1.13397	1,96179
1.10409E-02	2 172605	FUEL FLOW RATE . 186206	163921	7.774754E-83	CAID VOID VOI	FUSL IN TONE	ONID IN TAME
FUEL DROP SURA	A DXID DROP BULL	27.4			-		
.0		3419E-03		0,0	0.	10	10
FUEL GAS PASS	OXID GAS MASS	FLSL DROPLETS	DAID DROPLETS	PUEL STREAMS	CAID STREAMS	Fuel Ox BALL	ATTE OF MALE
0.147709E-07	7,548460E-16	3,597978E-06	6,297277E=C9	93			0,0
2.4543195-04	A OXID GAS EFFLUX	SALE DRG BASE	DAID DROP BATE	FINEL FILE BATE	21 a a 16	4	ST STATES
		3	•	•	•	7.0000	\$622/2
Fuel phop pag 39,4369	34,4306	FUEL INJ. VEL. 2.53693	DXID INJ. VEL. 19,9492	HAX, EXPLOSION 5,05342	1,491694E-84 3,369633E-89	3, 3996338-05	0,3462736-05
TIPE HILLISEC	CHAMBEP PRESS	IGNITIO!	FUEL FRACTION	CHAMBER TEMP K	POL NT.	64 PHA 1-13000	CF VAC
FUEL INJ FATE	OXID INJ RATE	FUEL FLOW RATE	OXID FLOW RATE	FUEL VCID VOL	CXID VOID VOL	FUEL INJ TEMP	OXID INJ TEMP
FUEL DROP BURN	N OXID DROP BUFF.	FUEL FLASH RATE	OXID FLASH RATE FUEL WALL BURN 2,4754815-02 0,	FUEL WALL BURN		FUEL MALL EVAP	OXID HALL EVAP
FUEL GAS PASS	OXID GAS HASE	FUEL DROPLETS	OXID DROPLETS	FUEL STREAMS	CXID STREAMS	FUEL ON MALL	DXIB ON MALL
FUEL 6AS EFFLUX 3.9827476-64	OXID GAS EFFICK	FUEL DROP RATE	OXID DROP RATE	FUEL FILM RATE	CXID FILM MATE		VAC THRUST LBF
FUEL DROP D32	OXID DEGP D32	FUEL INJ. VEL.	OXID INJ, VEL.	MAX, EXPLOSION	PASS DUT OF TANK INTEGRAL PCODT	INTEGRAL PCOBT	TOTAL IMPULBE
SOUTE AT 1.00	HILISECONDS						
TIME MILLISEC	CHANBER RRESE	1611101	FUEL FRACTION	CHANDER TEMP K	P.01	Avnav	20 200
1.00000	3,78761	3.00000	9, 917499E-12		20,0317	1,24603	1,00900
1.707707E-02	0XID IN RATE	FUEL FLOW RATE.	OXID FLOW RATE	A ROARDOCAN	i.	FUEL INJ TRAF	0X10 14, 18p
		707/491	70.000	PIDALOK BERNS	\$104//02FeB2	100'102	274.010

FIRE GAS MASS	440						
2,530406E-06	2,29A363E-1.9	1.704896E-06	1.500449E-04	6,264390E-00	CAID STREAMS	FUEL ON MALL	
FUEL SAS EFFLU	SAS EFFLUX ONTO GAS SFELLY	SUEL DROP BATE	OXID DROP BATE	FUEL PARE BATE	CAID FILE BATE.	GAS FRACTION .	NC THRUST LBF
4, 7070245-04	2.064000E+C	•	•	-	•	1.0000	0.63736
FIEL DROP D32	OXIC 1808 032	FUEL 154, VEL- 3,92173	29,1070	13, 9555 13, 9555	3,667376E-04	K 155684L -BC+BT	2,139130fee3
TIME HILLISEC	CHAPPER PRESS	16×1710×	FUEL FRACTION 2,2856415-02	CHANBER TENP K	FOL b7.	64HHA 1-84678	CF VAC
FUEL INJ KATE	DXID IT.J HATE	FUEL PLON RATE	OXID FLOW RATE	FUEL VCID VOL.	CAID VOID VOL	FUEL INJ TEMP	OKET LAL SIXO
FUEL DROP FURN	OXID DPOP BUPA	FUEL FLASH RATE	OXID FLASH RATE	FUEL HALL BURN	CAID WALL BURN	FUEL HALL EYAP	OXID MALL EVAP
FUEL GAS FASS	ONID GAS PASS	FUEL DACPLETS	OXID DROPLETS	FUEL STREAMS	CXID STREAMS	FUEL ON HALL	OXID ON WALL
FUEL 648 EFFLUX 4,5045265-03	X CXID GAS EFFLUX PUEL	PUEL BRCP RATE	OXID DROP RATE	FUEL FILM RATE	CXID FILM MATE		VAC THRUST LOF
FUEL DROP D32	CKIE DPOP D32	FUEL INJ. VEL.	OXID INJ. VEL.	HAX, EXPLOSION	PASS DUT OF TANK	K INTEGRAL PC-DT	TOTAL IMPULSE
1.40000	CHAPPEL PRESS	16×1110×	FUEL CRACTION	CHANGER TENP K	10+	Canna	St VAC
FUEL INJ SATE	DXID IN BATE	FILES FICH BATE	OXID FLOW BATE	FUEL VOIS VOI	100 HED HEX	F141 141 1414	
6.83255gE-63	.270109	396525	.417984	•	0.713301E-03	204.002	294,002
2, 9319216+03	152171	CUSL FLASS RATE	OKIS FLASH RATE	1,2797106-04	3,396963E=84	FUEL MALL STAR	9,73443,6-63
2,751793E-06	0X10 6AS MASE 1,239369E=64	FUEL DROPLETS	0X10 DB0PLETS	FUEL STREAMS 0.197534E-06	CX10 STREAMS	FUEL ON MALL 1.5240406-07	0X10 00 Kell
THE GAS EFFEUR	TOXID GAS SPELLY FUEL	FUEL BROR PATE	OXID DROP BATE	25.00 mm. C. part	CKID FILE BATE	AAS FRAGTION V	ART SOMBLE OF
Cité pare pas	The section	3					21,000
32,6361	23,1105	1.94969	31,2270	24.7674	6,717107E-04	2,4561798-63	,,6000,40003
11ME MILL1880	CHAMBER PRESS	NO THE	FUEL FRACTION	CHATBER TERP K	70L KT.	OAMAA.	S AVE
FUEL INJ RATE	OXID INJ RATE . 274618	PUEL FLOW RATE	OXID FLOW RATE	FUEL VOID VOL.	4:	FUEL 111 787P	0x15 Ln. 48x0
FUEL DROP BURN	OXID DROP BUPK	FUEL FLASH RATE		OXID FLASH RATE FUEL MALL BURN	CATO NALL BURE	PUSE MALL BYAD	OXYD DALL DWAD

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TIME MILLISEC CHAMBER PRESS TRAITIES

6, 409324E=00	1,1409396-04		0.	2.9003706-05	CXID 87REAMS	FUEL ON MALL	OXID ON MALL
2,265911E-03	A OXID GAS EFFLUX FUEL . 101923 0.	FUEL DPOP RATE	E OXID DROP RATE	E FUEL FILM RATE	2	GAS FRACTION	15
FUEL D*0P D32	14,4801	FUEL 1KJ. VEL.	0x1D INJ. VEL. 31.6694	. MAX, EMPLOSION 101.290	8.910732E-04	AMK INTEGRAL PCODY	
THE MILLISSE	CHANNER RRESS	IGNITION	FUEL CRAPTICE	1		T	1
1.0000	2,04040		1.0802186-02	2 694.768	46.0093	1.16073	200
. 49090	OXID INJ RATE	PUSL FLOW RATE	OXID FLOW RASS	E FUEL VOID VOL	CXID VOID VOL	FUEL TR. PRES	
FUEL DROP SURN	OXIO DEGE SIST	24.06.		•	7.0967645=03	il.	204.062
ċ			O.	AND THE PART OF	7	1	OXID BALL BRAR
FUEL GAS MASS		PUEL BRCBI CYS	OXIN NBABISES	5003A//AE003		<i>:</i>	1,3093678081
				3.0221626-05	1.100796E-04	FUEL OR MALL	OXIO ON MALL
THE GAS SERLING	GAS EFFLUX OXID GAS SEFLUX	FUEL BROS 8416	OXID DROP BASE	FIRE CHE BATE	•		A POLOGODO
Charles		÷	•		10	1.00000	AND THRUSS LOS
79.7001	OXID DROP DES	FLEL IN VEL	DXID IN VEL	MAX, EXPLOSION.	PASS OUT OF TANK	=	
	:		35,3636	155.410	1,0456246+03	3,71763	
TIME MILLISEC	CHAMBER PRESS	16h1710h	FUEL FRACTION	CHAPBER TEMP K	POL VT.	BANKA	CF VAC
FUEL INJ RATE	OXID INJ RATE	FUEL FLOW RATE	0X10	2	CXTD VOID VOL	FUEL TALL TRUE	1000
FUEL DROP BURN		FUEL FLASS CATE			5,004460E=03		
			010 1.43H RATE	3,3874495-88	7.10 KALL BURN	FUEL WALL EVAP	OXID BALL BYAP
	0XID GAS MASS	FUEL DROPLETS	OXID DROPLETS	FUEL STREAMS	CXID STREAMS	FUEL ON WALL	OXID ON WALL
FUEL GAS EFFLUX 1,5045425-03	OXID GAS EFFLUX	FUEL DROP RATE	OXID DROP RATE	₹.	CXID FILM RATE	GAS FRACTION	VAC THRUST LBF
FUEL DROP 032	OXID DPC/ D32 F	FUEL 18-3. VEL.	OXID INJ. VEL.	Ž	PASS OUT OF TAME INTEGRAL		1 5
]	AAS EVELLIEN			1-3660-05-07-1	4-1666694-1	21.249000E=96
OXID GRANS	12011114	.03711138	0.0000000000000000000000000000000000000	0.0000000000000000000000000000000000000	. 00079299 . 03732948	. 11989433 . 12444299	. 00022401 . 00022401
FUEL CROP	FUEL TROPS INJECTED 31	46466404	•		i.	TOTAL HOMENTUP	
don's cape	1	9.905707E+c6	2.078261E-04	5.763653E-01	2.486879E-01	1,3006396-02	

	FUEL FLOS RATE	OXID FLOW RATE	FUEL VEID VOL.	4.76566E-83	204.002	204. DAS
THE STREET	FUSI STASE EASE	DXID ELASE BATE	FUEL MALL BUILD	CKER LALL BURN		
e.		0, 9,409536E=05	9,4695368+05	3,2529396-05	7.	1,3060738-03
1,107667E-64	FUEL DROPLETS 2,705000E-04	2,339684E-64	FUE, STREAMS 0,197534E-06	9, 862639E-05	3,5201335-09	0118 ON MALL
1,274402E+03 6,318611E-02	PUST DRCP RATE	OXID DROP BATE	FUEL FILE BATE	CAID FILE BATE	And Fraction w	26, 500 26, 504
080P 035	FUEL 15.1. VEL.	33,8709	MAX, EXPLOSION 250.061	1,475150E-03	14TESAAL PCAST	4,676796662
CHAMPER PRESE	IGNITION	FUEL FRACTION	CHAMBER TEMP K	+0L HT.	8474	7 A C
OXID INJ RATE	ON RATE	OXID FLOW RATE	VOL	CXID VOID VOL	FUEL INJ TEPP	OKID INJ TONE
i	ASH RATE	OXID FLASH RATE	FUEL WALL BURN	CAID WALL BUM	PUEL WALL EVAP	OXID PALL EVAP
OXID GAS HASS	FUEL DROPLETS	DAID DROPLETS	FUEL STREAMS	CX10 STREAMS	PUEL OK WALL	OKID ON MALL
3	OXID GAS EFFLUX FUEL DROP RATE	OXID DROP RATE	FUEL FILM RATE	CXID FILM RATE	2	VAC THRUST LOF
	FUEL INJ. VEL.	ONID INJ, VEL.	HAX, EMPLOSION	PASS OUT OF TANK	TANK INTEGRAL PC-DT	TOTAL IMPULSE
	19K1110N	FUEL FRACTION	CHANDER TEMP K 2064,35	FOL MT. 18,9918	1,2436	of VAC 1,03713
[FUEL FLOW RATE	OXID FLOW RATE	FUEL YOLD YOL	CAID VOID VOL	FUEL INJ TERP 204.004	CX10 1NJ 78HP
OXID DROP BUFN.	FUEL FLASH RATE	OXID FLASH RATE 0.	FUEL WALL BURN 4,004003E-03	CXID WALL BURN 0.	FUEL MALL BYAP	OXID HALL BYAP
0XID 645 HASE 1.14A536E-04	FUEL DROPLETS 3.4162666-04	OXID DROPLETS	FUEL STREAMS 0,1975345-06	CXID STREAMS 1,60010E-05	FUEL ON MALL 6,3746376-89	OXID GN WALL
3	FUEL GAS EFFLUX OXID GAS EFFLUX FUEL DROP RATE . 203514 . 301237 0.	OXID DROF RATE	FUEL FILM RATE	CAID FILM MATE	1.0000 V	VAC THRUST LBF 164.675
320	FUEL INJ. VEL. 112.044	OXID INJ. VEL.	MAK, EMPLOSION 189.978	1.936246E-03	1NTEGAAL PC-BT 3,0778426-82	107AL 1MPULSE 1100500
3	HILLISECONDS					
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FUEL DROP D32	0XID DR0P D35	FUFL IAJ. VEL.	OXID INJ. VEL.	MAX, EXPLCSION	PASS GUT OF TANK	INTEGRAL PC-DT	TOTAL EMPULSE
71KE HILLISEC 3.6666	CHAMBER PRESE 104,129	16N171Ch 1.06000	FUEL FRACTION	CHANDER TERP K 2050.15	FOL WT.	64MA 1.2350	CF VAC 3,040°2
FUEL INJ FATE	0x10 11.3 RATE , 722543	FUEL FLON RATE . 479615	OXID FLOW RATE . 722552	FUEL VEID VOL	CAID VAID VOL	FUEL INJ TENP 204.000	6x10 1nd 78m
FUEL DROP BURN , 354512	OXID DHOP BUFA.	FUEL FLASH RATE	DXID FLASH RATE	FUEL MALL BURN	CAID MALL BURN 1	FUEL MALL EVAP	GXID MALL EVAP
FUEL GAS KASS 1,316256E-04	0XID 645 MASE 1,9721876-04	F.EL DROPLETS 4,416498E-04	OMID DROPLETS 4,125125E-04	FUEL STREAMS 0,107534E-06	CAID STREAMS 1.011401E-04	FUEL ON WALL 1,390549E-04	CXID ON MALL
FUEL GAS EFFLUX ,318201	OXID GAS EFFICK FUEL B. 439973 5.98	FUEL DROP RATE 5.9801196-03	OXID DROP RATE	FUEL FILM RATE	CXID FILM MATE	BAS FRACTION V.	VAC THRUST LBF 277.219
FUEL DROP D32 107,454	0x1D DROP D32 50,5703	FUEL INJ. VEL. 100.225	0x1D 1NJ, VEL. 83,5187	MAX, EXPLOSION 414,328	PASS OUT OF TAME 3,139009E-63	INTEGRAL PC-D 1115318	107AL IMPULSE 136687
71ME MILLIEGE 3,80000	CHANGE PRESE 113,617	16N2 1.04	FUEL FAACTION	CHANGES 75HP R. 3096.92	20.2511	1.23310	1.05043
FIEL INJ. SATE. 465007	OXID INJ RATE	FUEL FLOW RATE	DXID FLOW BAYE.	FUEL VOID VOL	YOU	FUEL INJ TERE	Date in the
THE DROP BUSH . 301411	OXID DROP BUSA	FUEL SLASS RATE	OXID FLASH RATE	4,651040E-03	1,162611E+05	THE RALL STAF	CKIS PALL SYN
1.406647E+84	2,205752E=64	FUEL DROPLETS 4,333562E-84	0XID DROPLETS.	FUEL STREAMS 0,197534E-04	- 1	FLEL ON MALL 1,5450946-04	170 MM MALL
. 340050	. 520706	1,647394E-62	3,230609E-05	FUEL FILE BATE	2,9167266-05	100000 A	304.944
100,100	OXID DROP DAG 61.0634	FUEL 18-1 VGL	0X10 1NJ, VEL. 62,0000	435,291	1,375509E=63	1NTEGRAL PCASI ,130942	1014 January 1800
71ME MILLISEC 4.0000	CHAMBER PRESE	16h 1710h	FUEL FRACTION	CHANGER TENP K 3543.87	POL WT.	64 PM A - 1.93089	CF VAC
FUEL INJ KATE	OXID INJ RATE	FUEL FLOW RATE	OXID FLOW RATE	FUEL VOID VOL	VOL.	FUEL INJ TEMP	OXID INJ TEMP
FUEL DROF BURN	OXID DROF BUFA	FUEL PLASH RATE	OXIO	FUEL WALL BURK	CXID HALL BURK !	FUEL WALL EVAP	OXID HALL BYAP
FUEL BAS HASS	OXID GAS HASE	FUEL PROPLETS	OXID DROPLETS	FUEL STREAMS		FUEL DI: MALL	OXID ON WALL
SOBOOD STATUX	FUEL GAS EFFLUX OXID GAS EFFLUX FUEL DI	FUEL DROF RATE	OXID DROP RATE	FUEL FILM RATE	CXID FILM RATE	æ	VAC THRUST LBF
ruel page 532	OXID DROP DIS	PLIES TRAS. VES.	OXTO THE WE	MAY RUDI DRIBL		1	

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FUEL INJ RATE OXID	OXID INJ RATE	FUEL FLOW RATE	OXID FLOW RATE	FUEL VOID VOL	CXID VOID VOL	FUEL 1KJ TEMP 204.032	CXIE INJ 76HP 294,032
FUEL DROP BURN OXID	_	FUEL FLASH RATE	OXID FLASH RATE	FUEL WALL BURN 1,256022E=02	CXID WALL BURN 1,177026E=02	FUEL WALL EVAP	OXID HALL EVAP
1,223616E-64 2,	0X1D GAS MASS 2,466720E-04	FUEL DROPLFTS 3.250250E-04	0X1D DROPLETS	FUEL STREAMS 5,664330E-07	CAID STREAMS 1,3061846+04	FUEL ON WALL 5,165009E-04	3x16 GN WALL 5,915,016.0
Š	FFILX	FUEL DROP RATE 4.1501816-02	0XID DROP RATE 2,970564E=02	FUEL FILM 9ATE	CXID FILM RATE	18 FRACTION , 030639	VAC THRUST LBF 321,644
FUEL DROP D32 OXID	0x1D DROP D35	FUEL 14J. VEL.	OXID INJ, VEL. 75,4666	MAX, EXPLOSION 910,946	PASS CUT OF TANK 1	IN INTEGRAL PC-DT	1014, IMPULSE 2,26906

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MASS THROUGH INJECTOR - PAUNDS	.004196	690800'	1,940736	4076,036382	243,109040	
MARK THROUGH MOZZIE . POUNTS	003844	004458	1,824882	4883-150841	267,144663	
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15 S S S S S S S S S S S S S S S S S S S	TRACTION METAINED AS FILM .144072 .028307 .08844 TOTAL OF ABOVE VALUES 1.00080 1.08800 1.08900 TAS GREATER THAN 3.0 MAY INDICATE SMATTER FLOME MIT CLASSED AS STRONG-LIANGE ASS OF COLD VAPOR IN CHAMER BEFORE STARY, BET-ETION OF CHANGER GAS THEME HAY BE AN APPRECIABLE MASS OF COLD VAPOR IN CHAMER BEFORE STARY, BET-ETION OF MALL-FILL, WHEN A PIPING IS STARTED WITH PROPELLARY-COATED WALLS, BEALETION OF WALL-FILL, WHEN A PIPING IS STARTED WITH PROPELLARY-COATED WALLS, BEALETION OF WALL-FILL, WHEN A PIPING IS STARTED WITH PROPELLARY-COATED WALLS, BEALETION OF WALL-FILL, WHEN A PIPING IS STARTED WITH PROPELLARY-COATED WALLS, BEALETION OF WALL-FILL, WHEN A PIPING IS STARTED WITH PROPELLARY-COATED WALLS, BEALETION OF WALL-FILL, WHEN A PIPING IS STARTED WITH PROPELLARY-COATED WALLS, BEALETSON OF WALL-FILL, WHEN A PIPING IS STARTED WALLS, BEALETSON OF WALL-FILL, WHEN A PIPING IS STARTED WALLS, BEALETSON OF WALL-FILL, WHEN A PIPING IS STARTED WALLS, BEALETSON OF WALL-FILL, WHEN A PIPING IS STARTED WALLS, BEALETSON OF WALL-FILL, WHEN A PIPING IS STARTED WALLS, BEALETSON OF WALL-FILL, WHEN A PIPING IS STARTED WALLS, BEALETSON OF WALL-FILL FILLS WALL-FILL WALLDS AND AND THE SAIT PLANCE. 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69,3107 169,8367 49,4377	MEAN AXIAL VELCCITY FT/SEC 73,43 321,46 53,17 566,21 MEAN RABIAL VELCCITY FT/SEC 73,43 321,46 53,17 566,21 116,26 116,26 116,26 116,26 116,26 116,26 116,26 116,26 116,26 116,26 116,26 116,26 116,26 116,26	MEAN AXIAL VELCCITY FT/SEC 73,43 321,46 93,17 960,21 MEAN RABIAL VELCCITY F7/SEC 73,43 63.46 93,17 916,28 MEAN RABIAL VELCCITY F7/SEC 71.6 83.46 17 716 7M80AT PLANE, NOT THE SYIT PLANE, DIE - THE ABOVE EJECTED DROPLET AND EJECTED FILM VALUES ARE AT 716 7M80AT PLANE, NOT THE SYIT PLANE,	MEAN RADIAL VELCCITY FY/SEC 73,43 323,46 53,17 560,23 MEAN RADIAL VELCCITY FY/SEC 73,43 323,46 53,17 546,38 146,38 156 - 746 ABOVE EJECTED DROLLET ARE SILLE VALUES ARE AT THE THROAT PLANE, NOT THE EXIT PLANE,	MEAN RABIAL VELCCITY FT/SEC 73,43 MEAN RABIAL VELCCITY F7/SEC MEAN RABIAL VELCCITY F7/SEC MEAN RABIAL VELCCITY F7/SEC 13,43		45,4377	02,5420
74,48 822,46 94,27	MEAN RABIAL VELCCITY ET/SEC. 110.28 110.28 120.28 120.28 120.28 120.28 120.28 120.28 120.28 120.28 120.28 120.28 120.28	MEAN RABIAL VELCCITY F7/SEC. 13.6.36 13.6.36 14.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15.6.36 15	MEAN RABIAL VELCCITY ET/SEC 03.01 016.26 016.26 016.26 016.26 017 - THE ABOVE EJECTED DROWLET AND FILM VALUES AND AT THE THROAT PLANE, NOT THE EYET PLANE,	MEAN RADIAL VELCCITY ET/SEC. OTE - THE ABOVE EJECTED DROLLET AND EJECT	321,46	53,17	560.21
93.01	TS - THE ABOVE EJECTED DROFLET ARE STUDY AND ARE AT THE THROAT PLANE, NOT THE EXIT PLANE,	ITE - THE ABOVE EJECTED DROFLET AND 61212 VALUES ARE AT THE THROAT PLANE, ADT THE EXIT PLANE,	ITE - THE ABOVE EJECTED DROPLET JAC EJECTED FILM VALUES ARE AT THE THROAT PLANE, NOT THE EXIT PLANE,	ITE - THE ABOVE EJECTED DROLLET ALC EJECT	10.10		15. 4. 2. E. C. C. C. C. C. C. C. C. C. C. C. C. C.

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	e.rogeed seconds				
	OBBOOK SECONDS	8			
INTERVAL LENGTH	. OHEODE SECONDS	50			
TOTAL IMPULSE	1.699527 POUND-SEC	-SEC THRUST	212,446915 POUNDS	Sanga Fi	
STATES PRESENT INTERNAL		PPC CHANGE PRESSURE	ESSURE - 77,487278 PB14	ries a	
PUEL OUT OF TAPK	. PUJ226 POUNTE	E FUEL FLOW RATE		.483277 POUNDS/SEC	
GRID OUT OF TAKE	1004719 POUNCE	S CAID FLOW RATE		.989983 POUNDS/SEC	
PURE-ONTO OUT OF TANK	SUNDA SPEACE	5 101AL FLOW RATE		POSTOR POLNBASEC	
PIXTURE RATIO	1.442775 OXID/FUEL	FUEL			
GOSTAR	3614,3F5240 FEFF/SEC	SEC			
3979146 BLEEFE 141974	84-00-35-000-01-01-01-01-01-01-01-01-01-01-01-01				
	Phupillant Expe	PELLED THIS INTERVAL			
	1304 964714 1304	FUEL POUNDBYSES	THO SENDE OF TO	Ox15 Pounds/SEC	
PRODUCTS OF COPPUSTION	11177890	.222304		,405234	
UNDURNET WAPOR	\$10000Cu.	, 00001V	. 00000127	,000159	
Chauman Dropies	06793000	200200	03000001	\$200701	
HALL FILP	.00000925	. 663657	. 00000467	E-5060.	
3	1400134	DINETER AND VELOR	#		
	FUEL DROFLETS		OXID DPOPLETS	\$	
BS HICPCRS	100.071961		50.640490	01	
SOL NICHONS	190,0100				
-832 MICFONS	160,228731		96,27:095	•	
ANIAL VELOCITY FT/SEC	365,651774	1	650,333156	2	
Asplat vetocity flyste			100,02630	1	
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1.00							£75	91:391109 -0014087F 1003	584		PHACTION	CUPULATIVE	**2000*	***************************************	,619876	.021200	***************************************	,022984	1029100	103601	1110949	+9200+	194800	. 966613	. 080613	\$1909a*	, 985845	50686	6006001	
							OXID DEOFLEYS	166116	.015965		Oxic rass Phaction	INCREMENTAL	,00026	.007646	,011962	. 001403	1002904	0.00000	,006204	900300'	.087873	5790624	*863931	. 417915	000000	3000000	. 005032	000000	00000019	
LUFS FOR PULTHAN	V 1	•	**2/SEC**2 TEG.A	*2/SEC**2 Ftu.R		ats/it, sec.		hespt roos		EXFILLED DRCPLET SIZE DISTRIBUTION	PRACTION	CLPULATIVL	.060321	.000145	£9749.	.016984	.0294c4	. 022355	.034372	7,6960.	.073743	.120074	240042.	.148081	.19760	116922'	.380605	46855	CC6800'	
IVELA NAL	77,447272 PS1	3048 3:3821 DEC 9	15A3,433A37 FT.	12486.472133 Ffee2/SECee2 FEE.K	\$660 6a. I	, 24.9092 POUNTS/FT, SEC.	FULL DAGILETS	45.2F4509 POURTS/FT445	.052085	EXFILLED DAC	PIEL HASS	TEGI SPIETAL	. 344621	***************************************	. 54455	. rusous	0306460	. 003951	.01201	339310	.036791	. 346331	. 02000	0.00000.0	125670'	404824	.154000	,067990	00000010	
	SUF t.	Phatter	•		4			ŧ	ASE GAS			RADIUS FELT	1.640:-05	3.2421.035	4.5211-65	6.5621-65	\$3= 1202°3	9.843t-05	1.1466+04	1.3121-04	1.476t-04	1.640104	1.004k=04	1.9691-64	2.133k-04	\$0-1462°2	2.4011-64	2.625+-64	2.789Emps	
	CHANGER PRESSUFF.	CHAMBER TEPPERATURE	GAS CORSTANT	SAS SPECIFIC HEAT	OAS BENTA	GAS VISCOSITY		BROPLET DENSITY	MASS DROFS/PASS GAS			BIAM MICHON	10	2	36	07	*	•0	70		0.6	100	813	120	130	043	150	360	848	

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100	**	210	220	***	240	250	0.02	270	286	003	300	310	986	336	340	**	360	370	98	390	20+	**	420	400	

INTERVAL LENGTH	STRONG SECONTS			
SHPERVAL LERGTE.	\$2.40.36 000.15°			
	. LUZODO SECONDS			
TOTAL IPPULSE	. 0:3454 PCLRE-SET	THUST	311,727191 POLNDS	POLNES
Charles Farestry 11 1:0141.	- 234084 PS18948C	CHAPTER PRESEUPE	112-010031-091v	
PUEL OUT OF TANK	. ALJESS POLLYPS	FUEL FLOW RATE	.467491	.407401 POLNDS/SEC
GKID OUT OF TALK	.CJ1307 FOLNES	CAID FLOW PATE	.053717	.053717 PCLNDS/SFC
		turat Fton Reif	1,765.09	1,001409 POUNTS/SEC
MINTURE HATTO	1.0"3462 "XIE/! UEL			
G-87AR	4849,143899 FEFT/SEC			
APC SAFEIN IS INDICAS				
	FROPILLANT EXPELLED THIS INTERVAL	THIS INTERVAL		
£+4	file - frends - Puer - Pource/Sie		0915 - POUNDS - CX18 - POLNDS/SEC	P0th05/5EC
PRODUCTS OF CCPPUSTION	. 3 3061451	,307255	.00125099	,625403
ENBURNED WAPCH	3,6366888	00.0 0000001	0.0000000	9,00000
CABUTALD BROTLLTS		**********		4+44381
balt FILM	. 9: 600263	. 001313	. 60000232	.001809
**************************************	EXPELLEPT PICPLET MEAN ETAN	tat Elanethic and Velocity		
FUF	FUFL BHOFLETS	0	OXID DROPLETS	
839 MICHONS	114.345511	•	52,797727	
Set Hickory	149,174490		961049-06	
B3g HICRONS	171.959618		99.559510	
ANIAL VELOCITY FT/SEC	296,672866		997,102629	
ABBIAL VELOCITY PT/SEC	90,320704		133,003077	

### COLING THE TANK TANK TANK TANK TANK TANK TANK TANK	GMAMBER PRESSULE	RESSULE	111.C10831 PSIA	10731 PSIA			
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12233,345982 TropySECock Ded.R	046 CO.ST.	ANT	1000,470037	Toop/SECoo2 DEL.R			
#### 111 Disoler POLNES/FT.SEC, ###################################	THE BELLEVIE		12233, 345582	T**2/SEC**2 DEU.R			
## ### ### ###########################	648 VISCUS	1114	4 860090.	OLNES/FT.SEC.			
### ### ##############################	BROPLET DE	***	FIFE DESILETS		OXID DROPLI	118	
##ETUS FEET INCREMENTAL CUMLATIVE INFRENTAL 1.64GI-05 0.4GD00 0.000000 1.000000 4.521-05 0.4GD00 0.000002 0.000000 6.5026-05 0.400000 0.000002 0.000000 1.148-04 0.4G0000 0.011712 0.000000 1.148-04 0.4G0000 0.011712 0.000000 1.148-04 0.4G0000 0.011712 0.000000 1.148-04 0.4G0000 0.011712 0.000000 1.148-04 0.4G0000 0.011712 0.000000 1.148-04 0.4G0000 0.011712 0.000000 1.148-04 0.4G0000 0.011712 0.000000 1.148-04 0.4G0000 0.011712 0.000000 1.148-04 0.4G0000 0.011712 0.000000 1.148-04 0.4G0000 0.011712 0.000000 1.148-04 0.000000 0.01712 0.000000 1.148-04 0.000000 0.01712 0.000000 1.148-04 0.000000 0.01712 0.000000 1.148-04 0.000000 0.01712 0.000000 1.148-04 0.000000 0.01712 0.000000 1.148-04 0.000000 0.01712 0.000000 1.148-04 0.000000 0.01712 0.000000 1.148-04 0.000000 0.01712 0.000000 1.148-04 0.000000 0.01712 0.000000 1.148-04 0.000000 0.01712 0.000000 1.148-04 0.000000 0.01712 0.000000 1.148-04 0.000000 0.01712 0.000000 1.148-04 0.000000 0.01712 0.000000 1.148-04 0.000000 0.000000 0.0000000 1.148-04 0.000000 0.000000 0.0000000 1.148-04 0.000000 0.000000 0.0000000 1.148-04 0.000000 0.000000 0.0000000 1.148-04 0.000000 0.000000 0.0000000000000000	PASS DAPPS	/"ASE CAS	. 344534	CURT Syl ton 3	196:14	· T	
##FIUS FEET INCREMENTAL CUMLATIVE INFERENTAL 3.000000 .000000 .000000 .000000 4.52159			t xrillED EF	IOPLET SIZE DISTRIBU		50	
10 1.6451-05 0.46000 0.000000 1.000000 1.000000 1.000000 0.0000000 1.000000 1.000000 0.000000 0.000000 1.000000 0.000000 0.0000000 0.000000 0.000000	BIAP MICHON		3	Patetica	Ţ	FRAST 104	
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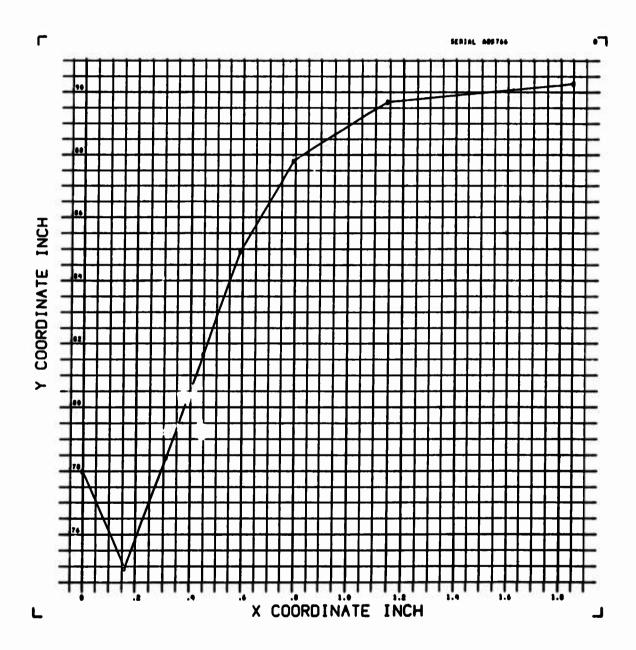
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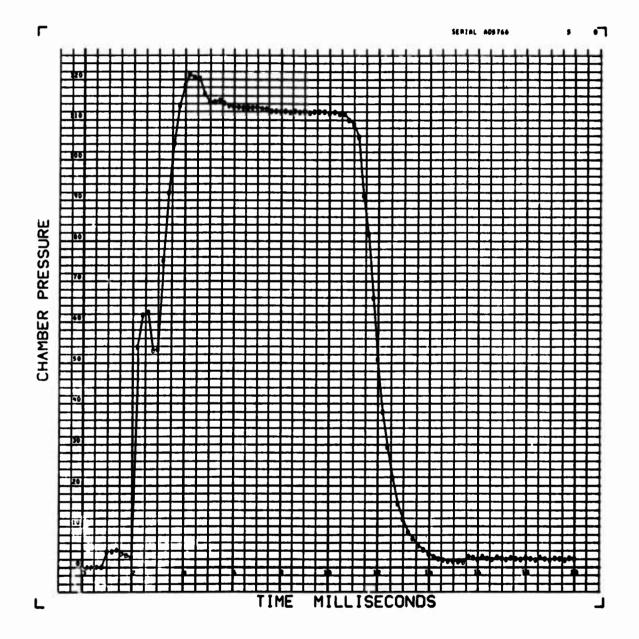
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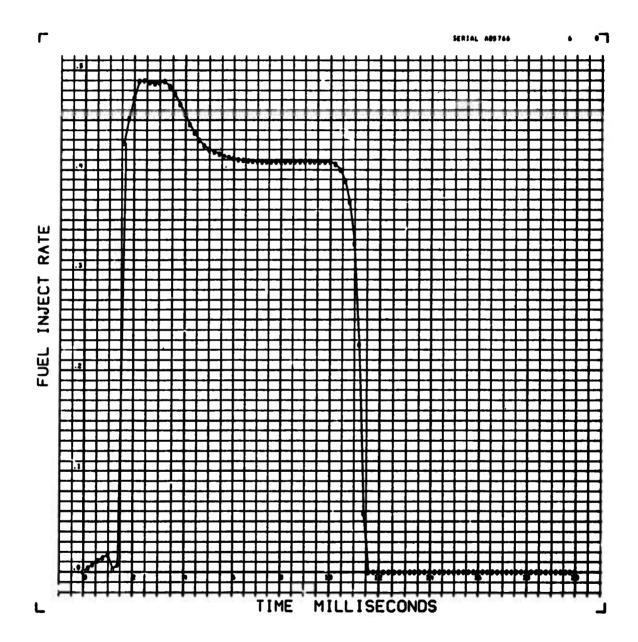
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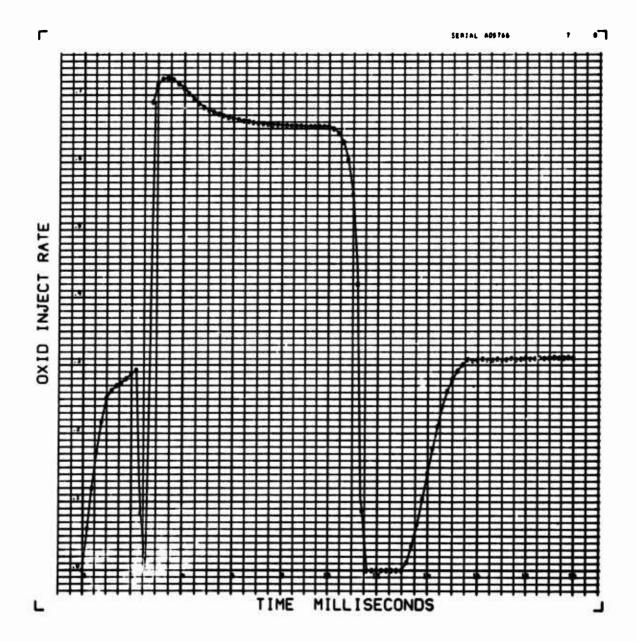
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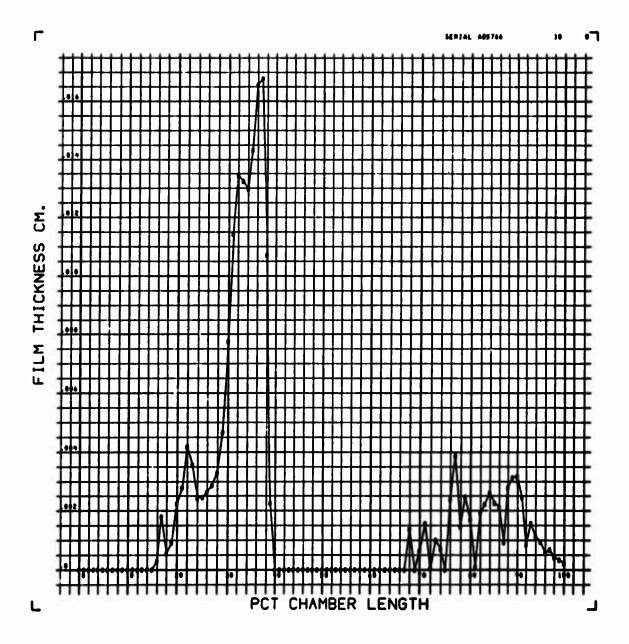
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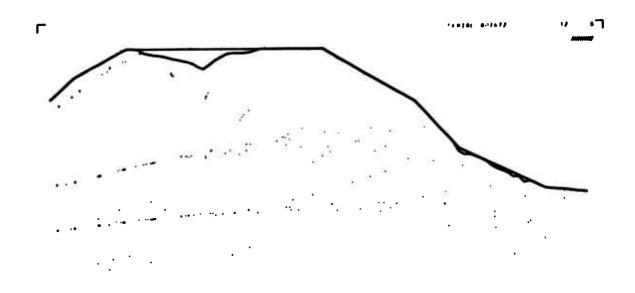


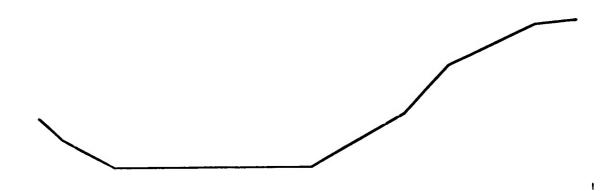












Appendix B

MULTRAN

MULTIPHASE NOZZLE AND PLUME TRANSPORT COMPUTER PROGRAM

A Multiphase Nozzle and Plume Flow Field Characterization Model

Program Number H612

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Appendix B

MULTRAN

MULTIPHASE NOZZLE AND PLUME TRANSPORT COMPUTER PROGRAM

A Multiphase Nozzle and Plume Flow Field Characterization Model

B. 1 INTRODUCTION

The computer program described in this appendix is a subprogram to the Plume Contamination Effects Prediction Computer Program, CONTAM, and performs the subsonic, transonic, and supersonic computations required to define the steady-state multiphase flow field within a rocket nozzle and exhaust plume. The TCC program (Appendix A) provides the input to MULTRAN in terms of quasi-steady values of droplet distributions, and gas properties, averaged over specified portions of a transient engine pulse. The nozzle and plume flow field defined by MULTRAN provides the input data for the kinetics and condensation computation, KINCON (Appendix C), and subsequently for the deposition and surface efforts computations, SURFACE (Appendix D). MULTRAN may also be used as an independent computer program on any third generation computer with a core exceeding 120,000g words and a Fortran IV processor.

The MULTRAN program combines three previously independent programs:

- a. <u>TD2</u>, Axisymmetric Two-Phase Perfect Gas Performance Computer Program (developed by TRW for NASA/MSFC, reference B-1)
- b. <u>TD2P</u>, Axisymmetric Two-Phase Perfect Gas Plume Analysis Computer Program (developed by Dynamic Science and MDAC)
- c. <u>SLINES</u>, Streamline Generation Computer Program (developed by MDAC)
- B.2 ANALYSIS, INTEGRATION METHOD, AND SUBROUTINE STRUCTURE

The axisymmetric two-phase analysis, numerical methods, and subroutine structure used in TD2, which also forms the basis for TD2P, is discussed in detail in Reference B-1; however, a very brief summary of the procedures given in the reference (as well as modifications necessary to treat contaminant transport) will be presented. The conservation equations of gas continuity, particle continuity, momentum, and energy in conjunction with the equations of the perfect gas law, particle drag and particle heat transfer

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which govern an inviscid axisymmetric, two-phase mixture flow have been derived under the following assumptions:

- a. There are no mass or energy losses from the system
- b. The gas is inviscid except for its interactions with the condensed particles
 - c. The volume occupied by the condensed particles is negligible
- d. The thermal (Brownian) motion of the condensed particles is negligible
 - e. The condensed particles do not interact
- f. The condensed particle size distribution may be approximated by groups of different size spheres
 - g. The internal temperature of the condensed particles is uniform
- h. Energy exchange between the gas and the condensed particles occurs only by convection
 - i. The only forces on the condensed particles are viscous drag forces
 - j. There is no mass transfer between the gas and the condensed phases.

The equations were put into the form of characteristic relationships in the reference where the characteristic directions are along gas streamlines, along gas mach lines, and along particle streamlines. It is found that the characteristics are real if the flow is supersonic (M > 1). Thus, the method-of-characteristics is used to compute the supersonic flow of a gas-particle mixture.

In the transonic flow region, however, solution for a gas-particle mixture is an extremely formidable task because the throat conditions are determined by the nozzle inlet geometry. Thus, to obtain initial conditions to start a characteristic calculation for a gas-particle system, the equations for the complete subsonic and transonic flow field in the nozzle inlet and throat regions must be solved. Qualitatively, it is found that the particle and gas streamlines diverge in the throat region, leading to the concentration of particles along the nozzle axis, greatly complicating the calculation of gas-particle flow properties in the nozzle throat. Mach number is found to be

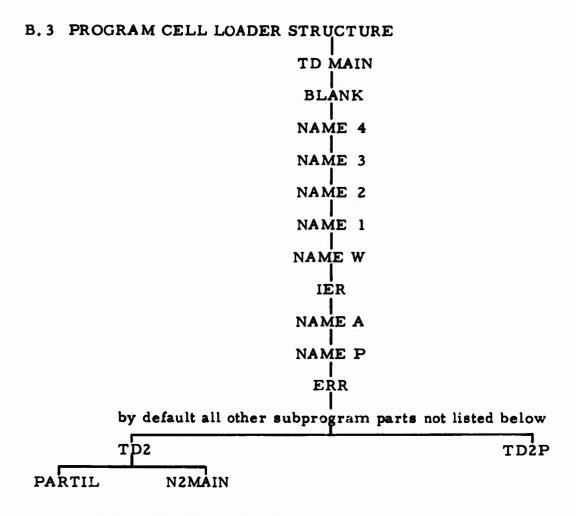
less than one at the nozzle throat; hence, the transonic zone in a gas-particle flow extends downstream of the throat.

In view of the complexities involved, only approximate solutions to the equations governing the transonic flow of a gas-particle system can be obtained. In order to simplify the calculations, the nozzle inlet and throat geometry is assumed to consist of a conical inlet section joined smoothly to a constant radius of curvature throat section.

The following method as discussed in Reference B-1 is used to obtain approximate initial conditions for the characteristic calculations. In the conical inlet section, the flow is assumed to be a one-dimensional sink flow. The equations governing the one-dimensional flow of a gas-particle system are solved to obtain the flow properties on the sink line. The gas properties in the throat region are approximated by the perfect gas relationships, and particle trajectories are calculated through the throat region to determine the particle properties along the initial line. An average expansion coefficient is used which approximates the gas-particle expansion, including the effects of gas-particle nonequilibrium.

When adapting this method to contaminant transport predictions, certain modifications are necessary to better approximate the physical processes involved. Specifically, as discussed in Reference B-1, the particles in the nozzle inlet region are assumed to be directed with the gas in a one-dimensional sink flow (converging toward the axis). This assumption is reasonable for typical solid propellant engines where particle radii are usually less than 15μ and the gas-particle flow is near equilibrium. However, in the case of bipropellant exhausts, the unburned fuel and oxidizer droplets can range from 15μ to 100μ in radius and be far from equilibrium. The particle trajectories as computed up to the nozzle throat by the transient combustion chamber model TCC indicate that these large droplets travel in approximately straight lines parallel to the nozzle axis.

In order to better model bipropellant contaminant transport in the nozzle, the particle distributions and velocities are specified at the geometric throat from the TCC output. Particle trajectories are then computed from the throat to the supersonic start line assuming the velocity at the throat is parallel to the axis. The gas-phase properties in the transonic region are computed as before. An option is available for solid rocket exhausts which computes particles trajectories in the original manner as outlined in Reference B-1.



B. 4 PROGRAM USER'S MANUAL

This program was developed on the CDC 6500 computer using the FORTRAN IV language. Conversion to another computer system should be straightforward provided sufficient core storage (120,000 words) is available. The Program Cell Structure extends three levels deep including the executive level, when used as a subprogram to CONTAM: the MULTRAN driver program, TDMAIN, is used to define the overlay structure for common blocks that are the same in TD2 and TD2P.

The description of the input to and output from MULTRAN is divided into the following four subsections:

- B. 4. a TD2 INPUT
- B. 4. b TD2P INPUT
- B. 4. c TD2/TD2P OUTPUT DESCRIPTION
- B. 4. d SLINES Subprogram

A card listing for the complete input for the sample case is given in Subsection B.5

a. TD2 Program Input

The program input for the Axisymmetric Two-Phase Perfect Gas Performance Subprogram <u>TD2</u> conforms to the I.B.M. NAMELIST format. All input items are read under control of the name \$DATA. The input items are divided into five types 1) Propellant Data, 2) Particle Data, 3) Inlet and Throat Parameters, 4) Characteristics mesh control data, 5) Nozzle Wall Contour Data.

For some input items, default values are assumed by the program. These items need not be input to the program if default values are desired.

\$ DATA

(1) Propellant Data

Item Name	Input Quantity	Unit s
CAPN =	N, Viscosity temperature exponent	none
CPG =	C _{pg} , specific heat of gas at constant pressure	ft ² /sec ² •R
CPL =	C_{p_l} , particle heat capacity $(T_p > T_{p_m})$.	ft ² /sec ² •R
CPS =	C_{p_s} , particle heat capacity $(T_p < T_{p_m})$.	ft ² /sec ² °R
GAMMA =	Y, specific heat ratio, C_{p_g}/C_{g_V} .	none
GMGO =	μ_{g_0} , chamber gas viscosity coefficient	1 b/ft sec
HPL =	h_{p_f} , liquid particle enthalpy $(T_p = T_{p_m})$.	ft ² /sec ²
HPS =	h_{p_s} , solid particle enthalpy $(T_p = T_{p_m})$.	ft ² /sec ²
ISØLID =	l, solid rocket exhaust	none
	0 , contaminant transport (default value)	

Item Name	Input Quantity	Units
PC =	Pg, chamber pressure.	PSIA
PR =	P _r , Prandtl number.	none
RCAP =	R, gas constant.	ft ² /sec ² •R
SMP =	m _p , particle density.	lb/ft ³
TG0 =	Tg, chamber temperature	•R
TPM =	T _p , particle solidification temperature.	•R

(2) Particle Data

Item	Input Quantity	Units
R(1) =	r_{p_j} , the radius of each of n particles is to be input so that $r_{p_1} < r_{p_2} \dots < r_{p_n}$ set $r_{p_{n+1}} = 0$. $n < 10$ is required.	ft
WPWGT =	$\Sigma \dot{w}_{\alpha}$ / \dot{w}_{α} , ratio of particle to gas weight flow.	none
W PW T (1) =	$\dot{w}_{p_j}/\Sigma\dot{w}_{p_j}$, particle weight flow fractions corresponding to each of the above particle p_j .	none

(3) Inlet and Throat Parameters (Figure B-1)

Item Name	Input Quantity	Units	Assumed Value
DZI =	Δz , particle trajectory integration step size.	none	0.002
DZMIN =	Δz_{\min} , inlet step size parameter.	none	0.002
NILP =	N _i , number of initial line points.	none	15
RRT =	R_c , throat radius of curvature. A value $R_c \ge 2$ is required.	none	

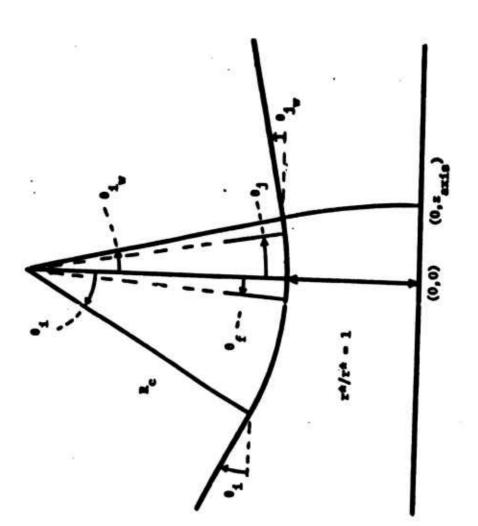


Figure B-1. Inlet and Throat Parameters

Item Name	Input Quantity	Units	Assumed Value
RT =	r*, throat radius	ft	
SAUR(1) =	First estimates of x_0 , u_0 , α , β , and Y for the special throat expansion. Required only if $\theta_i > \theta_f$.	none	-0.15, 1, 0.5, 0.3, -1
THFD =	$\theta_{\mathbf{f}}$, faring angle $(\theta_{\mathbf{f}} > \theta_{\mathbf{i}} = \text{no faring})$.	degrees	5.0
THID =	θ_{i} , inlet angle	degrees	
THIW =	θ, intersection of initial line and wall.	degrees	12.0
THJD =	θ, angle defining the zone farthest downstream	degrees	9. 0
VAR(1) =	First estimates of x_0 , u_0 , α , β , and Y for the zone farthest upstream	none	0.3, 0, 0, 0.1, 0.1
ZAX =	zaxis, intersection of initial line and axis	none	
ZI =	n; number of upstream zones.	none	3.0
ZJ =	n, number of downstream zones.	none	2.0
(4)	Characteristics Mesh Control Data		Assumed
Item Name	Input Quantity	Units	Value
DL =	Δℓ, maximum LRC mesh width.	none	0.2
DTWI =	$\Delta\theta_{\mathbf{w}}$, maximum flow angle change along the wall.	degrees	3.0
DR =	Δr, maximum RRC mesh width.	none	0.2
EW =	εw, end of nozzle wall criterion.	none	0.001
IMAX =	i max, maximum number of iterations per mesh point.	none	5.0
N1 =	n ₁ , select each n ₁ th LRC for print.	none	1,000
N2 =	n ₂ , print each n ₂ th point on selected characteristics.	none	1.0

(5) Nozzle Wall Contour Data

Item Name	Input Quantity	Units
IWALL =	Option flag.	none
	0=> tabular input	
	l=> cone	
	2=> parabola	
	3=> circular arc	

If the wall is to be input in tabular form (IWALL = 0):

$$PW(1) = (r_i, z_i), \text{ wall coordinates } i = 1, 2, \ldots, n \text{ points.}$$

$$viz.: PW(1) = r_1, z_1, r_2, z_2, \ldots, r_n, z_n, 0, 0.$$

$$Note:$$

- (a) always mark the end of the table with two zeros.
- (b) $n \le 79$ is required.

If a cone, parabola, or circular arc contour is to be specified then:

THJW =	θ_{j} , attachment angle for the contour; e.g., for a cone, the conical half angle.	degrees
EPS =	ϵ , nozzle expansion ratio (cone only).	none
RWMAX =	rmax' nozzle exit radius (parabola or arc only.	none
ZWMAX =	zmax, nozzle length from throat to exit (parabola or arc only).	none
\$	END OF CASE.	

NOTE:

A case is defined as the data included or implied between the \$DATA card and the \$ signifying the end of case. Whenever a value is input as data for a case, that value will remain set for succeeding cases until a new value is input. The values indicated as assumed by the program hold only until a different value is input. If more than one value is input for a given quantity within a case, the last value given will be used.

b. TD2P Program Input

The program input for the Axisymmetric Two-Phase Perfect Gas Plume Analysis computer subprogram is an extension of the input to the <u>TD2</u> computer subprogram. Input to the <u>TD2P</u> computer subprogram consists of a binary start tape ((TAPE 8) or an extend disc storage file (TAPE 8) generated by the <u>TD2</u> computer subprogram. The same NAMELIST data deck used for the <u>TD2</u> calculation must be used for the <u>TD2P</u> calculation and is read from TAPE 8 automatically. The following input items must be inserted into the data deck following the <u>TD2</u> data. Additional data for the TD2P subprogram is read under control of the name DATAP.

\$ DATAP

Item Na	me	Input Quantity		Units
PMA	=	θ _{PM} , The Prandtl-Meyer e angle to be used at the nozz		degrees
NPM	=	The number of Prandtl-Mey be generated.	er points to	none
		NPM < 0 The maximum num of points will be g		
		NPM = 0 Points will be ger degree apart unle than 5 points would	ss less	
		NPM > 0 This number of possible generated unless mum would be extended, points one degree apart would again exceed mum or would rest than 5 points.	the maxi- ceeded. If ald be are placed unless this ed the maxi-	
ZMAX	=	This item defines a cutoff p calculations. The run will ated when an axis point is c located downstream of $Z = Z$	be termin- alculated	none
PCUT	=	If a Pressure, Pg, is calcuthat the ratio Pg/Pgo drops value, no more points will ton the characteristic.	lated such below this be calculated	none

Item Nam	<u>ne</u>	Input Quantity	Units
NPLOT	=	Flag for auxiliary radiation output on tape unit 9, (see Description of Program Output).	none
		NPLOT = 0 deletes this output.	
		NPLOT = 1 requests this output.	
AREA	=	Flag for auxiliary force field output on tape unit 9. This item is also used to represent the frontal surface area of a body in the plume, (see Description of Program Output).	in. ²
		AREA = 0 deletes this output.	
		AREA > 0 requests this output e.g., AREA = 1.	
EMIS :	=1	ep. particle emissivity used for the radiation printout.	none
C1 =	=	C ₁ , force coefficient used for the force printout.	none
C2 :	=	C ₂ , force coefficient used for the force printout.	none

REMARKS

If the mesh width control items DL and DR have been input as part of the NAMELIST DATA deck, these items should be deleted for the plume input. The <u>TD2P</u> computer program will then assume values for these variables, minimizing the mesh points.

The last item printed at the end of a complete <u>TD2</u> calculation is the item IP = nm, where nm is number of points used along the last running characteristic. Reference should be made to this number to determine if sufficient space is available (98-IP) for the Prandtl-Meyer expansion.

The TD2P computer subprogram is not suitable for running consecutive cases.

c. TD2/TD2P Output Description

() The state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of t

Program output may be viewed in detail by examining the sample case in Section B. 5. The program output follows the sequence listed below:

For the one-dimensional inlet and the axisymmetric transonic throat calculations

- (1) The values k and mg for gas-particle equilibrium.
- (2) Initial and final conditions for the one-dimensional inlet integration.
- (3) Converged values for x_0 , u_0 , α , β , γ and f_1 , f_2 , f_3 , f_4 , f_5 for each transonic flow zone.
- (4) The corrected estimate for k, corrected values for x_0 , u_0 , α , β , and γ in the transonic zone containing the initial supersonic data line, corrected values for \dot{m}_{α} and \dot{m}_{p_i} .
 - (5) Items 2, 3, and 4 are iterated twice.
- (6) The gas-particle flow properties P_g , ρ_g , u_g , v_g , r, z, h_{p_j} , ρ_{p_j} , u_{p_j} and v_{p_j} along the initial supersonic data line.

For the supersonic method of characteristics calculations for the nozzle and plume, printout may occur after the completion of each mesh point calculation. Points for print are selected as follows:

The following points are always printed:

axis points.

Kth particle boundary points.

initial line points.

wall points.

Interior points are selected for print only along every n_1^{th} left running characteristic and only at every n_2^{th} position along these characteristics.

Inserted points are printed if all points are to be printed $(n_1 = n_2 = 1)$.

The items printed are listed below in the order they appear, left to right, on the output sheet. A header is printed for identification purposes above each characteristic.

Row One:

Item	Header	Meaning	Units
LRC number	LRC	Left running characteristic number.	none
Ident, number	ID	Type of point (see below).	none
r	R	r position coordinate,	none
z	Z	z position coordinate.	none
М	MACH	Mach number,	none
Tg	TG	Gas temperature.	• R
$\mathbf{v}_{\mathbf{g}}$	VG	Gas velocity (scalar).	ft/sec
$\theta_{\mathbf{g}}$	THETA-G	Gas streamline angle	degrees
T _g /T _{g_o}	TG/TG0	Ratio of gas temperature to chamber temperature.	none
P _g /P _g o	PG/PG0	Patio of gas pressure to chamber pressure.	none
ρ _g /ρ _{go}	DG/DG0	Ratio of gas density to chamber density.	none
$\sum_{p_j}^{k_{p_j}/\rho_g}$	SDK/DG	Patio of total particle density to gas density.	none
c _F	CF	Thrust coefficient,	none
I _{sp}	ISP	Specific impulse.	sec.
iteration no.	IT	Number of iterations required	none
		•	

Rows Two through K+1

A row is printed for each particle size, k=1, ..., K.

<u>Item</u>	Header	Meaning	Units
k	К	Particle size number,	none
Re _k	R EK	Particle Reynolds number	none
v_{p_k}	V PK	Particle velocity (scalar).	ft/sec

Item	Header	Meaning	Units
$\theta_{\mathbf{p_k}}$	THETA-K	Particle streamline angle.	degrees
T_{p_k}	TPK	Particle temperature,	•R
$^{ ho}_{ m p_k}/^{ ho}_{ m g}$	DPK/DG	Ratio of particle density to gas density.	none
$^{ ho_{ m p_k}/ ho_{ m p_o}}$	DPK/DP0	Ratio of particle density to chamber particle density.	none
$^{\mathrm{r}}\mathrm{p}_{\mathbf{k}}$	RPK	Particle radius	ft ^{et}

d. Additional Output

Table B-1 illustrates additional output options as discussed above.

(1) Force Field Print

If the force field auxiliary output is requested (AREA > 0) the following items will be computed and written on tape unit 9.

Item	Units
$F_g = \frac{C_1 A}{144g} \frac{\gamma}{2} P_g M^2$	${ m lb}_{f f}$
$F_{p_k} = \frac{C_2 A}{144g} \rho_{p_k} V_{p_k}^2$	$^{\mathbf{1b}}\mathbf{_{f}}$
$F_p = \sum_{k=1}^{k} F_{p_k}$	$^{ m lb}_{ m f}$
$F = F_g + F_p$	$^{1\mathbf{b}}\mathbf{f}$
F_{p}/F_{g}	None

The quantities C_1 and C_2 (which are nondimensional force coefficients) and A (which is surface area in square inches) are input as C_1 , C_2 , and AREA.

(2) Radiation Print

If the radiation auxiliary output is requested (NPLQT = 1) the following items will be computed and written on tape unit 9.

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Table B-1. OPTIONAL OUT PUT - RADIATION AND FORCEFIELD PROPERTIES

<u>.</u> .	*	*	*
&	11.4:2445:1	12.07%	10
76.15 19.7° 18	54.107405E-ul	20°-32'15'56'-1	10-11-60-65-56
# 12.0c5e47f-01 > 27.531e14f-01 3 34.7e9e4ef-01 4 73.21773ff-01 5 54.671431f-01 7 33.234346f-01 7 33.64436f-01	1 75.217006F-02 2 17.973304F-01 3 25.514093F-01 5 55.240016F-01 5 73.47496F-01 7 34.97392F-01 7 34.97392F-01 27.446106F-00	1 73.341.404F-72 2 17.444655F-01 3 25.042447F-01 4 54.476353F-01 5 79.015371F-12 4 54.550009F-01 23.741284F-00	1 71.120683f-02 2 16.947033f-01 3 24.372346f-01 4 53.345172f-11 5 78.140420f-01
12.0cf7066.09 +2.n154226.08 17.779176.08 13.1111526.08 51.7737536.07 14.9017866.07 7.584356.07	26.0374196.03 45.4033496.03 45.4033496.03 77.8749956.03 43.7148496.03 43.7148496.03 43.7148496.03 43.7148496.03	25.5315446.08 60.0131536.08 72.0855366.07 73.5865126.07 75.5808 60.00 15.537826.07	32.407745E.nb 18.432293E.nr 07.117495E.n7 98.666206E.97
71.4689255-11 26.4378-71-10 69.1884945-10 13.5311115-19 35.5728775-09 50.4475435-09	44.6378265-11 24.6211205-10 60.1906855-10 10.9460085-09 14.6992575-09 24.7235725-09 34.4195745-09	40.6229316-11 26.4211205-10 60.1906456-10 10.5450085-09 18.4925575-09 24.0616005-09 17.9119116-01	34.515105E-11 24.621120F-10 60.1906M5E-10 10.64600AF-09 14.699257E-09
##1 - 144 PK ##1.5336-7F-00 13.6270-3F-00 24.0534-4F-00 30.2454-3E-00 14.73954-9E-00 16.190230E-00 12.7714-02	25.40A8A5F-01 55.4ch69RF-01 10.304635F-00 13.005381E-00 83.355136F-01 53.853806F-01	22.559563E-n1 53.27.7255F-n1 55.424917E-n1 99.266053E-n1 12.614034F-n0 79.4354.7E-n1 FV/FG	14.134761E-01 50.399887E-01 52.434918E-01 94.374451E-01 12.109540E+00
79. 4 29. 3761995.1 39. 7561695.1 37. 3377415.1 60. 5867555.1 43. 4366725.1 44. 6436655.1	16.301#39£+1 29.376589£+1 29.376589£+1 29.376589£+1 30.184785£+1 32.738927£+1	14.6557025+1 29.3765806+1 29.3765806+1 29.3765806+1 29.3765806+1 29.3765806+1	14.971463E+1 29.376589E+1 29.376589E+1 29.376589E+1

Item	Header	Meaning	Units
Row 1:			
LRC number	LRC	Left characteristic number.	none
Ident, number	ID	Identification code.	none
r	R	r, radial coordinate	none
z	Z	z, axial coordinate	none
Rows 1 through	k _{max} (i.e., for	each particle size k = 1,	k _{max}):
N _{Pk}	NPK	Particle number density	number of particles/ft ³
q_1	RAD PK		BTU/sec
^q 2	RAD BAR PK		BTU-number of particles/sec
T ⁴ _{p_k}	TPK 4		•R ⁴
Row k + 1:	TBAR 4	Effective particle temperature	*R ⁴
where: $N_{P_k} = \frac{1}{\frac{4}{3}}$	ρ _{p_k} π r ³ m _p		
$q_1 = \epsilon_p$	$\sigma T_{p_k}^4 r_{p_k}^2$		
$q_2 = \epsilon_p$	$\begin{smallmatrix} \sigma & T^4_{p_k} & r^2_{p_k} & N_{p_k} \end{smallmatrix}$		
$\sigma = 0$.	475834 x 10 ⁻¹²	BTU/ft ² -sec-°R ⁴	
$\overline{T}_{p}^{4} = \frac{\sum_{k=1}^{k}}{\sum_{k=1}^{m}}$	$\sum_{k=1}^{\text{pax}} \sum_{k=1}^{1} N_{p_k} \sum_{k=1}^{2} N_{p_k}$	i.	

d. SLINES Subprogram

This computer subprogram, <u>SLINES</u>, was developed to provide the necessary interface between the <u>TD2</u> and <u>TD2P</u> subprograms and the <u>KINCON</u> subprogram. Basically, the program interpolates from data points on the characteristic lines (<u>TD2</u> and <u>TD2P</u> results) to determine points of constant percentage of mass flow (bounded by axis and point of interest) running through the throat, nozzle, and plume.

Subsection (1) lists the program variables FORTRAN names, symbols, and definitions. Subsection (2) discusses the program logic interpolation equations and editing technique. Subsection (3) provides the user with the necessary operating information. Subsection B.5 provides input and output for a sample case of the MULTRAN program, which includes the output from SLINES (last portion).

(1) FORTRAN Variables

The FORTRAN names given are the names used throughout the program. The symbols are used in the text of this document.

FORTRAN NAME	SYMBOL	DEFINITION
Z	Z	The axial distance of a point on a stream- line.
R	R	The radial distance of a point on a streamline.
P	P	The pressure at a point on a streamline.
S	S	The distance from the initial point on a streamline to a given point on a stream-line.
T	T	The temperature at a point on a stream- line.
v	v	The velocity of the gas at a point on a streamline.
KD	i	The i th point along a streamline. $(1 \le i \le n)$
JD	n	The total number of points along a stream- line before editing.
ID	j	The streamline identification number. $(1 \le j \le m)$
NSL	m	The total number of streamlines.
K	k	A given point on a characteristic line. $(1 \le k \le NPTS)$

FORTRAN NAME	SYMBOL	DEFINITION
NPTS	-	Total number of points on a given characteristic line.
N	N	The number of points on a streamline in the throat only.
PCM	η	The percentage of normalized integrated total mass flow pertaining to the point on a characteristic line.
PCS	•	The percentage of normalized integrated total mass flow criteria for a streamline.
Z, R, P, S, T, V ZMGKS, RMGKS, PMGKS, TMGKS, VMGKS	х	The class of parameters pertaining to the properties of a point on a streamline or characteristic line. Each parameter is defined elsewhere in this glossary.
DSK	ΔS	The streamline distance criteria used in editing.
sx	S ^{np}	The distance of the streamline from the supersonic start line through the nozzle and plume.
SID	-	The name of the namelist through which the user input data is provided.
ТФТМ	-	The integrated total mass flow along a characteristic line.
ZMGKS	-	The axial distance at a point on a characteristic line.
RMGKS	-	The radial distance at a point on a characteristic line.
PMGKS	•	The pressure at a point on a characteristic line.
TMGKS	-	The temperature at a point on a characteristic line.
ZINIT	-	The distance of the initial point on a streamline (ZINIT = $S(j, 1) = 0$).
PINIT	-	The pressure at the initial point of a streamline.

FORTRAN NAME	SYMBOL	DEFINITION
TINIT	-	The temperature at the initial point of a streamline.
VINIT	-	The gas velocity at the initial point of a streamline.
MGKSK	-	The total number of points on a stream- line after editing.
EXIT	-	The streamline distance of the last point on a streamline [EXIT = S(j, MGKSK)].

(2) Program Description

The program runs on the CDC 6500 computer system. It is written in FORTRAN IV and requires a field length less than 40,000g. A nominal case providing ten streamlines will require less than 7 central processor seconds and less than 20 peripheral processor seconds to execute.

The program includes five distinct functions:

- 1. Input
- 2. Streamline Location
- 3. Data Interpolation
- 4. Data Editing
- 5. Output

Each function is discussed in the subsequent paragraphs.

(a) Data Input

Two sources are required for data input: cards via NAME-LIST which is user data allowing the program to know how many streamlines are desired and the percent mass flowing within a cylinder bounded by each streamline; and logical file TAPE 12 (tape or disc) which provides the program with data from the <u>TD2</u> and <u>TD2P</u> subprograms on which to operate.

The data from the TD2 and TD2P subprograms consists of two records for each characteristic line. The first record tells how many points are on the characteristic line and the total integrated mass associated with the characteristic line. The second record contains the data designating the location, pressure and percentage of the normalized integrated total mass flow for each point on the characteristic line. The first requirements of the program is to provide initial temperature and velocity properties of each streamline. Therefore, the initial characteristic line has an additional record of data associated with it which gives the temperature and velocity properties associated with each point on the initial characteristic line. This

record follows the first two records, in order. The last record of both the TD2 and the TD2P subprograms contain a one word, negative number, record. This permits the program to know when there are no more characteristic lines provided by the respective programs.

(b) Streamline Location

The definition of a streamline is that line which runs through the throat, nozzle, and plume bounding a given constant percentage of the mass flow between it and the nozzle axis.

There are a discrete number of points on a characteristic line. The properties at any point usually will not meet the criteria for a point on the desired streamline, therefore, interpolation must be performed between the two input data points which bound the desired point on the streamline in order to obtain the necessary properties (location and pressure) on the streamline.

The initial point, with regard to distance along a streamline, is designated as the interpolated point on the initial TD2 subsonic start line. Each subsequent streamline point has a distance corresponding to:

$$S_i = S_{i-1} + \left[(Z_i - Z_{i-1})^2 + (R_i - R_{i-1})^2 \right]^{1/2}$$

where

S = the distance along the streamline

Z = axial distance along the streamline

R = radial distance along the streamline

 $i = the i^{th}$ point along the streamline $(1 \le i \le n)$

n = the total number of points along the streamline

As previously stated, $S_1 = 0$.

(c) Data Interpolation

Linear interpolation is used to locate a streamline point as follows:

$$X_{ji} = X_{k-1} + \left[\frac{\epsilon_{j} - \eta_{k-1}}{\eta_{k} - \eta_{k-1}}\right] \left[X_{k} - X_{k-1}\right]$$

where:

- X = the required streamline property (location, pressure, mass, etc.)
- the percentage of normalized integrated total mass criteria for the streamline
- The percentage of normalized integrated total mass pertaining to the points on the characteristic line
- k = the point on the characteristic line having the greater value and bounding the streamline value
- k-l = the point on the characteristic line having the lesser value and bounding the streamline value
 - $i = the i^{th}$ point along the streamline $(1 \le i \le n)$
 - n = the total number of points along the streamline
 - $j = the streamline identification number (<math>1 \le j \le m$)
 - m = the total number of streamlines
 - (d) Data Editing

There exists two reasons for editing the streamline data before output:

- l. Due to the change in the nature of the characteristic lines at the supersonic start line in conjunction with the logic used previously in the program, there exists a redundant point along the axis streamline at the supersonic start line. This redundant point must be removed.
- 2. The <u>KINCON</u> program will accept a maximum of 101 data points for each streamline.

To locate the redundant point along the axial streamline, the points along that streamline are searched until two points are found which have no separation distance. The redundant point and its associated properties are removed from that streamline. In addition, the location of that point provides information as to the location of the supersonic start line and the number of points in the throat, along the streamline, which will be required for use in subsequent editing.

It is desirable that each point and its relevant properties along each streamline in the throat be provided as input to the <u>KINCON</u> program. Therefore these data are not edited. The remaining points along

the streamline, in the nozzle and plume, must be edited by some criteria which will limit the total number of points along the streamline to not more than 101. The criteria chosen is such that each point (exclusive of the last two points) in the nozzle and plume shall be separated by at least some distance determined by:

$$\Delta S_{j} = \frac{S_{j}^{np}}{100 - N_{j}}$$

where

 ΔS = the distance criteria for editing

S^{np} = the distance of the streamline from the supersonic start line through the nozzle and plume

N = the number of points along the streamline through the throat to the supersonic start line

j = the streamline identification number

The last point along the streamline is always made available for output regardless of its distance from the previous point.

(e) Output

The output of this program is both printed and written onto a logical file (TAPE 8). The printed output is provided for the scrutiny of the user. The TAPE 8 file output is for use as input to the <u>KINCON</u> program. The written data contains only that information which is necessary for the operation of the <u>KINCON</u> program. The printed output provides the same data as the TAPE 8 file output but also contains other pertinent data which allows the user to determine the validity of the data. A sample of the printed output may be found in Subsection B. 5.

(3) Program User's Manual

Input to the computer program is divided into two subsections:

- (a) NAMELIST/SID
- (b) Logical File

Written output from the computer program is on logical file unit number 8. The sample case provided in Subsection B. 5 details the loadsheet for the NAMELIST card, and computer output.

(a) NAMELIST/SID

The NAMELIST feature permits the input of parameters without a format specification. The NAMELIST/SID contains two parameters:

- 1. NSL; the number of streamlines for which the program is to provide data.
- 2. PCS; an array of up to ten mass % values which provide the criteria for the streamline definition. The variable NSL determines the number of values which must be input.

The two permissible formats for PCS are:

- 1. PCS(1) = a, b, c,, j where a through j are the quantities corresponding to the PCS criteria.
- 2. PCS(2) = b, PCS(1) = a, PCS(4) = d, PCS(5) = e, PCS(7) = g, PCS(3) = c, and so on.

(b) Logical Files

The logical file input may be either tape or disc file. The unit number for the logical file is 12. This means that the logical file name used on the REQUEST or ATTACH card must be TAPE 12. The data on the logical file is provided from the <u>TD2</u> and <u>TD2P</u> programs. The data is binary (unformatted).

B. 5 SAMPLE CASE

Included in the section are the computer input and output for a sample case using the <u>MULTRAN</u> program as an independent program. The subprograms to <u>MULTRAN</u>, which are called sequentially are <u>TD2</u>, <u>TD2P</u> and <u>SLINES</u>. The use of MULTRAN as a subprogram to <u>CONTAM</u> for the analysis of contaminant transport is illustrated in Section 4 at the main text. To illustrated the use of <u>MULTRAN</u> as an independent program for other than bipropellant contaminant transport analysis, the sample case in this section is for a solid rocket motor nozzle and plume.

The calculation is performed for a typical solid propellant engine operating at a chamber pressure of 2,250 psia whose exhaust contains 13.6% Al₂O₃ by weight. The particle size distribution was obtained from experimental data of Radke, Delaney and Smith (Reference B-2). Six particle sizes are assumed with radius yielding weight flows as follows:

Particle Size (it x 10 ⁶)	Weight Fraction of Total Particles
4. 92	0.0214
9.84	0.1180
16. 32	0.2390
22. 90	0.3260
29. 40	0.2344
36. 20	0.0612
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A complete set of Control Cards and Data Cards for the sample case are listed preceding the sample case.

B.6 REFERENCES

- B-1. G. R. Nickerson and J. R. Kliegel. Axisymmetric Two-Phase
 Perfect Gas Performance Program, TRW Systems Report No. 028746006-R000, Vol. I and II, April 1967.
- B-2. H. H. Radke, L. J. Delaney, and Lt. P. Smith. Exhaust Particle
 Size Data from Small and Large Solid Rocket Motors. San Bernardino
 Operations, Aerospace Corporation, Report No. TOR-1001 (S295118)-3, July, 1967.

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VAN(1) *4.6136713E*81 1.0467436681 6.8.8386684	-3.7409339E-02		**************************************	1.34370406-00	-3, 73118696-62 -4, 40611696-61	34.69881108.76289148,39483106.46991	STEP SIZE . 00200u	5	78175	-1,42990205-01	1, 8108788161 1, 8108786161 16, 888788161	VAM(1)	3 1392656-01 1 9279496-01	VAR(1)	3,5175176E-01 3,4483819E-01	1, 9243389E=01	-7.1001031E-02 7.7901907E-01	4,4991254E-01 -9,2723006E-01	VAR(1)	4 949498 00 1 1397087E=01	VAR(1)	1.21345276.00
	•			400	***	WDP(1)* 9.73860	-2.66987	97	4197,775 .03237		n 12 4 10		, a o e i		- 20 10	- 45	•		•			~ ~
THEATION NUMBER		8 BARES.1924469	ITERATION NUMBER			WB6- 3346-137 ND	20s -4.7250 MAX		1109,190	4301		STERATION NUMBER		TERATION NUMBER					ITERATION NUMBER		TERATION NUMBER	

		**	4.670110E=02 -5.6361672E=01	6.39528996-06	::			
Bourn wollywood	1	+		444				
		-400	1.6130396F-01 1.3437642E-00	2,1792357	E-08			
		7 - 5	-1.7311990E-02 -1.7311990E-02 -4.4001199E-01	-1.3543399E-07 -2.3199911E-07 -4.7519091E-07	E-07 E-07 E-07			
TERATION NUMBER	IER 1		VARCES	r(1)				
		10.79	1,34376396000	-1,3464616E-09 -4,496616E-08	6-06 6-06 6-06			
		• •	4, 40 0 1 1 4 2 E - 0 1	-3,3171460	E-07			
DAS DIMENSIONAL	Ī		58148887188788378148139798188147188-27.83 117	910613979810	6.47 <u>200</u> 27.69;			
	40490 · Zo oXVX		3129 Size . 002000	95	458,346 - 19	1910- 6243,979	40= 4.1582	1
1364,530	Ī:	1000	1023.00	1336.906	0100,540	.93092	101	•
1277.610	3:33	. 6323		1225.162	0109.417	54800	. 75267	
ITERATION NUMBER	2 2	-	VARCES	503				
			1.000000E-01 1.000000E-03 4.0390504E-01	-1.5662005E-07	E-07			
		•	1.356766E-01	**************************************	6000 E-06			
	-		**************************************	4,18923,3	000			;
			1,920930E-01 4,9100639E-01	9,97627906=09				
ITERATION NUMBER) E 3		VAR(1)	F(1)	E-27			
			5,51,550,66-01 3,44431,386-01 3,78769888-01	-2: 904061 9E-07 9.3436098E-10	E-17 E-10			
TERATION NUMBER	JER 3	n° →	AAR(1)					
		474 173	7.791546E-01 7.791546E-01 4.0309226E-01	2.65529796-07 -1.6587886-07 3.64783696-03	E-03			
		•	.9;2719362E-01					
			VAR(1) 3.0195097E-01	1,1477189	E-07			
		7	1,1307054E-01 1,1307054E-01 -7,057056E-01	-4.12034416-07 -7.5102996-07 -1.05279566-06	E C 7			
[TERATION NUMBER	16A 3		VAA(1)	F(1)	E-07			
			9. 14Entess.		4			

	6	1141477645	U ->.U=03/04E	700
	3	4,9539044E=0		-06
	4	2,89577436-0	2 -6,6923970E	-08
	- 5		1 0,39901298	
ITERATION NUMBER 4	1	VARCE	F(1)	
THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY O				
	Ž	1,3437841E+0	0 -1,49533036	-07
	3	4,8115801E-0	1 -1.3543488E	-07
		-3,7311429Eu0	2 -2,31904908	-07
V -10 4 4004700	5	-4,4081143E-0	1 -4,7520591E	-07
K BAR=1,1924398	-			
ITERATION NUMBER G	1	VAR(1)	F(1)	
	1	-4.6130382E-0		-08
	<u>\$</u>	1:3457841E+0	0 -9,00102178	
	3	4.81156016-0		
	4	-3,7311429E-0		
	,	-4140471498-0	7 5124048075	-00
MBQ= 3346.204 HDP(1)=	9.73479	53,69968108,76	500148,35720106	67162 27,8511
DOGGOOGO CHANGE ZONES O	*****	**		
· z		ш	MBC	_
	,	K	MPC	P
=1,2570 =1,2500		104503	0.00000	1972,0469
-1,2500		09206	,00549	1773.0764
-1,2900			101532	1993,9179
-1,2500		118413	.02194	1994,1199
-1,2500		123016	03424	1994,9138
-1,2500 -1,2500		135553	.04928	1997,1363
-1.2500		,34826	.08736	1998.6056
-1,250è			11037	2000,3379
-1,2500		,46032	,1359+	2002,3970
-1,2500	-	,50436	116417	2004,6881
1,2500		133524	117487	2007.3549
-1,2500 -1,250c	-	,57842 ,64445	,22 4 03 ,26358	2010.3 0 13 2013.7 9 01
-1,25ch		109040	130147	2017, 6020
-1.2508		,73652	134159	2021,8347
-1.2500		.70255	, 38385	2024.5018
115900		195939	45,622	2001.0117
-1,2500	0	.87461	,47436	2037,1659
-1,2500 -1,2500	•	,92065	,52234	2043,1576
-1,2500	0	1.01271	, 97192 , 62293	2049,5704
-1,2500	Š	1,09874	67516	2043,5313
=1,2900		1,10476	,7284U	2070,9785
-1,2500	0	1.15081	,78240	2078,6400
-1,2500		1,19684	,83487	2006,4171
-1,29¢0		1,24267	189192	2074,1869
-1,2500 -1,2500		1,33494	,94602 1,00000	2101.79899
TOTAL MASS FLOH	- 3,55	44769E+03	NO. POINTS	. 30
MP				
1918.2912	4	6128,73463		
1918,2912 1917,9669		6128,77583		
1910,9870		8128,87787		
1915,3306		6129,10884		
1912,9634	8	6129,40749		
1909.8377		6754-80752		
1905,8932		6130,29724		
1901.0581	2	4130,90383		

70A7.1 5444A	6121100425		
1888,37457	4132,48771		
1880,33248	4133,49646		
1871,01980			
1860.31110	4135,95450		
1848,10221	6137,44651		
1824:52515	6239.28477		
1818,70556	6140,99867		
1801,29216	4143,07597		
1781193018	6149,36226		
1760:53096	6147.86036		
1737,02426	6150,56970		
1711,36497	6323,48297		
1683,54102	6176,57838		
1453,58349	6159,89263		
1621,98012	6203,34076		
1587,68906	6166,93121		
1552,16054	6170,60736		
1919,36013	6274,32739		
4477 7424			
1477,79873	6176,03231		
1440,16790	6181,65083		
1403,38026	1189,09809		
	·		
2	R	HPC	•
*1,16151	0.0000	0.00000	1968,1976
-1,16151	104435	100133	1968,2729
	,08870		
-1:16151		100732	1966,4997
1,16151	12200	101176	1400.0014
-1.16151	117741	02125	1707.4252
-1,16151	,22176	,03318	1970.13792
71,16191	124611	,04773	1971.03020
-1,16151	31046	06490	1972,1141
-1,16151	39461	08466	1973.4035
		100700	177314037
-1116191	139917	110498	1974.9132
-1,16151	,44352	13186	1776.6593
-1.16151	,45787	115724	1770,6583
-1116191	159222	118910	1900.92083
-1,16151	.57657	.22140	1983,48104
-1.16151	62093	25607	1906,3362
*1,16151	185728	,29306	
			1989, 50581
-1,16151	,70963	, 33235	1773.0011
-1,16151	,75398	137381	1996,82982
~1:10131	179833	141738	2000.9930
-1,16151	.84269	146297	2005.49784
-1.16151	.88704	51048	2010.3263
-1120171 	195139		
			2019,4722
-1,16151	197574	61075	2020.7060
-1,16151	1.02006	,66324	2026,5956
-1119131	2100444	17170	2038,4993
-1,16151	1.13880	177213	2038,5458
-1,16191	1,19315	82817	2044.67263
-1,16151	1119790	- 100494	2090,70370
-1,16151	1,24105	94235	2056,7660
-1,16151			
-1110131	1,24620	1:00000	2062.47262
TOTAL HASS FLOW =	3,5393880E+03	NO. POINTS	• 30
\ -			
	- 100 mg	HPG	
-1.07302	0.0000	0.0000	1939,71774
-1,07302	104283	100129	1939.70092
-1:07302	1,09999	100217	1139.97127
-1,07302	,12848		
		01163	1740.27120
-1.07302	117131	02067	1940,74492
-1.07302	121414	103227	5742.33776
-1.07302	,25696	,04644	1942.07675
₹ 1:07352	129979	06315	1942.9699
-4 077.0	74040	,,,,,	
	VA-18-1		

-+10.504	107606	. 49490	1779.06022
-1,07302	138544	10414	1745,25651
-1,07302	.42427	112942	1746.47034
-1107302	147210	179970	1948,27879
-1,07302	,51373	, 18433	1750.07236
-1,07302	,53675	, 21592	1772,12127
-2107308	197798	154484	2994197409
-1,07302		• 59950	1776.80012
-1,07302	, 60923	, 32479	1999,90345
-1,07302	172000	, 36962	1902,94779
-1,07302	177089	,40862	1965,75206
-1,07302	,81372	,45374	1969,19469
-1,07302 -1,07302	,05054	,50089	1972,00423
-1,07302	,94220	.54999 .60096	1976,74703
-1,07502	190502	19567	1909,00019
-1,07302	1,02705	70807	1909,42426
-1,07302	1,07048	76397	1993,06693
-1,07302	1,11091	182127	1998,32971
-1,07302	1,19633	87982	2002,74175
-1,07302	1,19916	93945	2007,01840
	1,24199	1,00000	2011,09989
TOTAL HASS FLOH =	3,52869478+03	NO. POINTS a	30
	-		
2	R	MPC	P
-,98453	0.00000	0.00000	1906,87539
-170495	104145	100120	1400.45524
-,98453	,05270	.00505	1907,06392
-, 78453	,12435	.01135	1707.30154
-178493	120900	102070	1707,03781
-,78453 -,98453	, 20725 , 24869	03152	1708.07617
	129014	, 04534	1708,62103 2709,27734
-,98453	,33159	· 00170 · 08072	1910,05076
-,98493	37304	10182	1916.94728
- 178493	142449	12990	1911,97311
• 98493	, 45594	19178	1913,13441
-,98453	, 49739	18040	1914,43704
-,78493	158884	121145	1919,00024
-, 98453	,50029	24402	1917,40633
-,98453	, 62174	28056	1919,24030
-, 78493	100317	131861	2982124944
-,96453	,70463	, 35894	1923,21287
-,90453	174608	40150	1925.42705
-170493	493 ,74608 ,40190 1929.425 493 ,70793 ,44429 1927.705 453 ,02090 ,49314 1930.275 453 ,07043 ,54211 1932.005 493 ,9100 ,99315 1939.996 493 ,95333 ,64604 1930.375		1927,78928
7,98453			1930.27707
-,98453			1932,88758
0,98493	,72100	126920	1499 34047
-,98453	. 7>333	,64004	1936,37937
- 98453	,99478	70084	1941.20278
98493	1,00023	75748	1944,02761
- 98453 - 98453	1,07766 1,11913	81580	1946,80616
1,78493	1,14097	, 87973 , 93717	1949,48168
-,98453	1,20202	1,00000	1991,90748
-1.0400	014444	TIANADA	
TOTAL HASS FLON .	3.5826275E+03	NO. POINTS .	- 30
		4, , 4,,,,,	43/E
2	R	HPC	P
*19A94	0.00000	0.00000	1009,19097
-,89604	+04021	.00124	1009,22601
-,69604	,08042	,00495	1869,30742
-, 898g4	12003	101113	7809,44403
-,89604	,14004	,01978	1869,63730
-,69604	.20105	,03089	1869,80904
****	-44.44		

	147160	. 4997	10/4,24170
-,89604	26147	06050	1870,57748
-,89604	32168	0789R	1871,01961
0,89004	136189	10990	
	10010	10700	1871,53084
-,89604	40210	12329	1872,11398
-,89404	,44231	14901	1872,77156
4,89904	140525	117719	1873,90968
-,89604	52273	20775	1874,31768
-,89604	54294	24069	1875,20801
0,89504	,60319	27598	
		18/240	1876,17584
-,59004	,64336	,31360	1877,21884
-,89604	, 68357	, 35394	1878,33279
7,89634	172378	139776	1879,55124
7,89604	,76399	,44023	1880,74513
-,89604	.80420	, 48692	1882,02234
4,89604	184441	,53586	1003,32720
-,89604			
7 1 9 9 9 9 9	, 85463	,58682	1884,64038
-,89604	192484	, 63995	1885,93759
194604	149203	169714	1007,10966
-,89604	1.00526	,75233	1888,36258
-,89604	1,04547	,81144	1889,41500
■,898 04	1,00518	187251	1890.29962
-,89604	1,12589	,93536	1890.96163
-,89604			
-10400+	1,10016	1:00000	1871,33825
TOTAL HASS FLOW =	3.5186764E+03	NO. POINTS .	30
	0102001010		
t		ирс- ·	
-,80756	0.00000	G • 0000C	1826,20444
-,80756	,03910		
-140/20		.00122	1826.20791
-,80756	107821	100486	1959 . 57944
-,60756	11781	.01094	1626.23631
-,80756	19642	.01945	1026,26199
▼,8 <u>0</u> 790	19992	103039	1826,29608
- 60756	, 23463	,04375	1826,33926
-,80756	,27373	05953	1826,39219
180758	131203	107773	
	131243		1826,49947
-,80756	135194	, 09435	1826,52949
80756	139104	•12137	1826,61433
-,0 ₀ 756	,45019	14679	1820,70761
-,80756	,46925	•1746a	1826.81436
-,80756	,50835	20480	1826.92678
4,00750	194746	123736	1827,04413
	159656		
-,0756	• 10, -10	27232	1827,16244
-,80756	,62567	.30961	1827.27633
V; 90796	,66477	,34924	1827,37874
-,60756	,70388	.39119	1827,46064
- 80756	,74298	43545	1827,51080
₹10 0796	170200	48200	1927, 51940
-,80756	,02119	,53082	1827, 45776
-,80756			
	,86029	,58189	1827,31800
-189796	189940	163514	1954 04597
-,80756	,93850	,69044	1826,69415
•,80756	,97780	,74432	1826,15083
-10 ₁₁ 790	2102072	190912	1925, 40015
*.80756	1,05581	87002	1824.41824
	1,07472		
":80756		, 93399	1823.13998
4180730	7:23405	1.00000	1851.27804
TOTAL HASS FLOW =	3.5174243E+03	NO. POINTS .	30
14149 1466 1564	-1/1/40444	MAT LANKIA	~~
Z	H	HPG F	
-,71907	0+00000		1777,40965
		0.0000	
4,71707	103012	100126	1777,36600
-,71907	.07625	.00450	1777.31500
▼ ":71907	11438	.01086	1777.19649
- 7.0.7			

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0176 0136 0156					֡
# 10 # 10 # 10 # 10 # 10 # 10 # 10 # 10	6127963FP-4 S.	S. 7842524Feat		4 64437376000	
10 17 10 10 10 10 10 10 10 10 10 10 10 10 10	1.0504046-0	\$,592314F-63	20-342194Z+4		0 4 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5
49764018907 4978404894444444444444444444444444444444	2		1.0847898E+03	9,7428587E-01	5.6084909E-01
124	7 6.965FACE=03	5.245389E+03	0.579170E-02		
. 924124E+0	ě	4	1.00873085+03	P, 3112202E=01	5,61969986un
	7 1.095GGGEEGG	5,471136E-03	8.073476E-02		
Probaggiert?	İ	į	4.444467892		
3,17304106-06 3,		6747136F-63	9. AFAT2876-02	0.000AB00Fees	R 0402446aa
1.9333725.0				70-4940099014	10-3-0-3-1-1
4,985620E+07		5.221266E-03	5.939962E+02		
51045592E-0	1.697666E-62	- 1	i		
		- 3			
56-56 3.	36		ċ	8,8153400E-01	0.0421592E-01
4.0044000404			7.4476908-02		
5.049462E+07	7 1.6936756-02				
Syndone - Co		1	311301902012		
5,116643E+37	7 2-0131216-02	4,595521E+03	150		
26036 3.	.257428\$£-a4	.0443960E+#3	9. n1507175enp	0.64440301-001	4.444708500
235E+		5.4212566+03	7.245885E+02	STREET, SALV.	
4.097419E-0					
5.091497E017		4.7267495			
5,110942E+J7			2.348979E+02		
5,1431936+07			The second		
.3510296k+r6 3.	3,3590570E-01 5	. 9809939E+03	7.7023584E+02	7.73396076-01	6.49303506=01
1		1	1		- I
5,0165485+07					
5,069919E+C7			200		
7113770450					
54154679E+07	7 5.7379476-03	6.4401.68E-03	1.600251E-02		
4E+26 3.	.4237071E-01 5.	i		7.0308733E-01	6.7545744Fans
3456+	il R		5.5300295-02		
51629109E+3			41300420E+02		
5.1165016-07	1.0720001102	4.6437500.03	3.150167E-02		
94144986417		į	ZO SOLITABLE DE NO		
5.104004E+U7	7 5.620423E-03		1.4510196-02		
36+26 3.47	*		•	6,32770606-01	6.09120536-01
4.9944796.97					
5.3396426-37	1		ı		
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THOUPHASE NOZZLE PROGRAM MAS BREN COMPLETED	CP TIME # 205,407 PP TIME # 121,921			
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		0,1266+01, 0,10016+01, 9,12666+01, 0,10026+01, 0,19826+01, 0,19826+01, 0,19826+01, 0,1246+01, 0,12246+01,
		0,010E+01, 0,1019E+01, 0,000E+01, 0,1012E+01, 0,501E+01, 0,1517E+01, 0,631E+01, 0,1629E+01, 0,795E+01, 0,1749E+01,
	-	0.16+02, 0.18746+01, 0.1266+02, 0.20266+01, 0.15826+02, 0.21868+01, 0.19956+02, 0.23846+01, 0.2916+02, 0.29996+01,
		0,316E+02, 0,276E+01, 0,398E+02, 0,3E+Ō1, 0,501E+Ō2, 0,3292E+01, 0,631E+02, 0,3934E+01, 0,795E+02, 0,3829E+01,
		0.1001E+04, 0.2002E+02, 0.1E+04, 0.2E+04, 0.0, 0.0, 0.0, 0.0, 0.0
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TPM = 0,417E+04, GPG = 0,1575E+05, NILP = 15, THID = 0,3E+02, MPWT = 0,214E-C1, 0,118E+00, 0,239E+00,	0,3268+00, 0,23446+00,
TPH = 0.417E+04, QPG = 0.1575E+05, NILP = 15, THID = 0.3E+02, WPWT = 0.214E-01, 0.118E+00, 0.239E+00, 0.612E-01, 0.0, 0.0, 0.0,	0,3268+00. 0,23446+00.
TPM = 0.417E+04, GPG = 0.1575E+05, N1LP = 15, TMID = 0.3E+02, MPHT = 0.214E-01, 0.118E+00, 0.239E+00, 0.612E-01, 0.0, 0.0, 0.0, 0.0, RT = 0.697E+00,	0,326 8 +80, 0, 23 44 £ +00,
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9,62280 0.00000 0.00000 16,51373 10,18421 .11892 .00034 15,49992 10,60690 .26062 .26062 .20470 AASS FLOH = 6.82280016+03 NO, POINTS = 3 Z HACH TG VQ THETA-G TG/TGO PG/PGO 10,55917 3,2774 3119,3 8815,6 0,000 ,49909 6,4689936-03 10806-01 6.7578626-03 0, 3,1539476-03 3,1617456-03	9,62280 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,18421 10,1
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MACH TG VG THETA-G TG/TGO PG/PGO VPK THETA-K TPK DPK/DG 7 3,2774 3119,3 8815,8 0,000 ,49909 6,4688938-03 8.7578626-03 0. 3,1539478-03 3,1617498-03	MACH TG VG THETA-S TG/TGO PG/PGO Va THETA-S TG/TGO PG/PGO Va THETA-K TPK DPK/DG 7 3.2774 3119.3 8815.8 0.000 .49909 6.46893E=03 8.153947E=03 3.161745E=03 8.454232E=03 4.745512E=02
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	0.464624E003 0: 3.454282E003 4.7454091E-02

THOFFHASE PLUNE PROGRAM HAS BEEN COMPLETED	CP TIME & 825,132 PP TIME # 131,990		

	CP TIME = 325	,153 PP	TIME = 133,690	
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0505 AM				
ZINIT =	INE(1) = 0,0000(CHONDINENS		1 1 0 M
PINIT =	1772.764306	(PSIA)		
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ZF INAL.		(NOND ! HENS	IDNAL - Z/RC)	
	12	INO, OF PR	ESSURE TABLE PO	11NT81
POINT	AKTAL	HADIAL	STREAMLINE	PRESSURE
,00	DISTANCE	DISTANCE	DISTANCE	(PSIA)
	(Z/RC)	(R/RC)	(S/RC)	
1) •	0.	1,992764E+03
	-1.161511E+00 () •	8,848896E-02	1,968198E-03
)	2,694669E-C1	1,906875E-03
غ -	-8.960442E-01	; ;	3,539559E-01	1,869199E-03
,	#01079592E-01	71	4.4244485-01	1,8262042-03
í	_4 0.00000) () (5.309338E-01 6.194227E-01	1.777410E+03 1.727201E+03
	-9,420005E-01	7	7,0791176-01	11032414E+03
10	-4.535994E-01	•	7.964007E-01	1,652289E+03
11) 	8,848896E-01 9,733786E-01	1,604440E*03
ร์งั	-1.801325E-01) (1.061568E+00	1.480200E+03
1.9	-9.964350E-02	•	1.150357E+00	1.430930E+03
16		,	1,327334E+00	1,317779E+03
17		,	1,415823E+00	1,246342E+03
		T	1,5043126+00	1,106698E+03
17 20	4) () (1,592A01E+00 1,681290E+00	1,116877E+03 1,057315E+03
	01000000E-01 0	11	2.0200005-00	8,084789E*02
22	8.626364E-31	•	2.132636E+0U	7.709337E+02
2ð		•	2.217302E*00 2.304699E*03	7,325796E+02 6,933135E+02
25	I IADABKELI I		2.395485E+0U	6,525191E-02
29	1.240607E+00 0	10	2.490607E+00	6.111341E-02
26			2,698053E+00	5,260718E+02
27	1.563166E+00 0		2.813166E+00	4.8276556+02
	1, 688363E+00	•	2,958363E+00	+;392225E+02
31 32		le io	3.076226E.00 3.230156E.00	3,954678E+02 3,517774E+02
35		7	2.4615435400.	51339095g-05
34	2,284704E+30 0	1	3.534704E+00	2.731407E+02
35		10	3.650110E+00	2,560348E*02
37 37		1	3.777314E*00 3.909475E*00	2.0590356+02
36	2.7980746+30 0		4.048070E+00	1.820713E+02
39	2,941874E-00 0	•	4,1710742700	110441045-03
40 41		0 0	4.346856E+00 4.510932E+00	1,441947E+02 1,276404E+02
42	3:434967E+30 0	1	4,6849678-00	1.1130518-02
43	3,413755E+00 0	•	4.863755E+00	9.816917E-01

/ . 66/40/18 + 67/ 6 . 73/00/18 + 61/ 8 . 96/16 66 + 61/	4,1401306+01 4,1401306+01 8,2275606+01	2,574049E=01 2,192847E=01 2,106970E=01		2,269369E+01 2,269369E+01 2,196363E+01	7 04234E+01 1 049940E+01 1 049940E+01		
5,42007E+02 5,420095E+00 5,627391E+00	6,268630E+00 6,739338E+00	7,674562E+00 0,00447E+00	6,412996E+00 6,412996E+00 6,627977E+00	9.179992E+0U 9.534540E+00	v, 97//36E+UU 1,046114E+D1 1,107280E+O1		
178995E+38 377391E+88	018630E+00 0	6,424582E+00 0, 6,75447E+00 0,	162996E+00 377977E+00	3			
;; ;	.	188 188 189	. W.W.	96	9 7 6		

		INE(2)= 20.000			FLOW
	ZINIT =	3.0000		ONAL - S/RC1	
	TINIT .	2007:82343 6136:18548			
	VINIT .	1858.42102			
		11,13200	1 (NUNDIHENSI	DNAL - Z/RC1	
	HGKSK =	55	(NO, OF PRE	BSURE TABLE PO	INTS)
	POINT	AXIAL	RADIAL	STREAMLINE	PRESSURE
	NO.	DISTANCE	DISTANCE	DISTANCE	(PSIA)
		12/RC)	(H/RC)	(S/RC)	
	1	-1.250000E+36	5,995139E-01	0.	2,0078236+03
	3	-1.101311E-38 -1.073022E-00	5,371440E-01	1,786454E-01	1,9517898-03
	Ă	-9.845331E-01	5,235767E-01	2,676898E-01	1,915353E+03
		-019404456-01	9,129879E-01	3,570647E-01	1,0741128-03
	7	-8.075552E-01	5.021362E-01	4,461629E-01	1.8269096-03
		-7,170662E-01	4,924377E-01	5,351817E-01 6,292196E-01	1,773265E+03 1,7490732+03
	9	-5.420883E-01	4.704441E-01	7,138592E-01	1,7046926-03
	10	-4.535994E-01	4,655492E-01	8,024947E-01	1.4934106+03
	- 11	-3.651194E-01	4,007004E-C1	0,911100E-01	1,5990148-03
	12	-2,766214E-01	4,562891E-01	9,797152E-01	1,5325858+03
	13	-1,881325E-01	4,914948E-01	1,068334E+00	1,477028E+03
	15	-1.115454E-02	4.476713E-01	1,245354E+00	1,351434±03
	10	7,733442E-02	4.467961E-01	1,3338486+00	1,2839905+03
		-11450234E-01	41449979E-01	11422337E+00	1,2130006+03
	1.5	2.543124E-01	4.474593E-01	1,510830E+00	1,1449938-03
	19	3.428013E-01	4,500540E-01	1,599346E+00	1.0751176-03
	20	7,488261E-01	4,929000E-01	1,0078712-00	7 8400475403
	22	8.266563E=31	4.714112E-01	2.005423E+00 2.067308E+00	7.812043E+02 7.514840E+02
		- 91457896E-01	91100407E-01	2,2143n4E+p0	6,7812398+02
	24	1.081492E+00	5.609717E-01	2.356396E+00	6.086027E-02
	25	1.215880E+00	5,996573E-01	2,496241E+00	5,426303E+02
	20	113470295+00	8:3398365-AI	5.0252225.00	4.0000115.05
	27	1.4780188-30	6.64470 E-01	2.7662935+00	4,256136E-02
		1.605764E+00	6.903317E-01	2,896631E+00 3,027842E+00	3,769050E+02 3,334399E+u2
	30	1.870469E+00	7.414044E-01	3.166220E+00	2,937517E+02
	31	2.013737E+00	7,710184E-01	3.312517E+00	2.572703E+02
	32	5.1084846+00	8.0403908-01	3,4087982900	2.2363838+05
	33	2.330585E+00	8,411768E-01	3.637043E+00	1.9272536+02
	34	2.508448E+00	8.831555E-01	3.819793E+00	1,651280E-02
	39 36	2.7029292+00 2.860634E+00	9,308437E-01 9,934297E-01	4,0199882-00	1,424400E*02 1,279978E+02
	37	3,104043E+C0	1,0005836+00	4,199147E+00 4,427479E+00	1,142637E+02
		3,3703472+00	1,038358600	4,6092772+00	1,0170092002
	39	3.646197E+00	1.113960E+00	4,981361E+00	8 9689036+01
	40	4,103559E+C0	1,200693E.00	5,448431E+00	7,426068E+01
	41	4,2615352+00	1,2413992.00	9,6097982+00	8.970832E801
	42 43	4.488760E+C0 4.776931E+C0	1.286976E.00 1.34622JE.00	5,841509E+00	6.396294E+01
		5,051874E+00	1,4009345400	6,135706E+00	5,7375396+01 5,2004518+01
	45	5,369631E+30	1,463867E+00	6,739969E+00	4,6671516-01
	46	5,691198E+GO	1.527228E+00	7.067720E+00	4,200858E+01
	47	9,0003745+00	1.9998295.00	7,4438092+00	3,7380042401
	48	6,445995E+CO	1,674527E+00	7.836756E+00	3,327434E+01
7	49	6,868851E+CO	1.755874E+00	8,267366E+00	2,945093E-01

•					
	1,653636-01 1,653636-01 1,476056-01				
9,164284E+00 2 9,630842E+00 2	1,0116718-01 1 1,0614186-01 1 1,1132606-01 1				
1,922456E.00 2,00760E.00	2,2788586.00 2,1940376.00 2,1940376.00				
7.798164E-00 6.206034E-00	9.45534E-18				
	7.3.8				

	ILINE(3)= 40.000	44 644 644 644	THE TOTAL MASS	FLON
ZINIT			IONAL - S/RC)	
PINIT				
TINIT				
VINIT 2FINAL				
HGKSK			IONAL - Z/RC? Essure table Po	ITHTE
HUNUN	- ,,	ING! WE FAI	ENGONE INDIE	14497
POINT	AXIAL	RADIAL	STHEAMLINE	PRESSURE
NOT	DISTANCE	DISTANCE	DISTANCE	(PSIA)
	12/RC)		19/RC)	
1	-1.250000E+06	7,993263E-01	0.	2,028344E+03
	-111013115.00	7.0003012-01	9.0440842402	
Ī	-1.073022E+00	7.023021E-01	1.808096E-01	1.965110E+03
•	-9.845331E-01	7,446223E-01	2.710474E-01	1,925349E+03
		7.2742226-01	3,6119466-81	
•	-8.075552E-01	7.116973E-01	4.510722E-01	1.8274716.03
7	-7.190662E-01	6.969157E-01	5.407407E-01	1,768371E+03
-	-613037732-31	017372462-01	0:31/110E-07.	2,7078992003
9	-5,420883E-01	6.668657E-01	7.207029E-01	1,712268E+03
10	-4,535794E-01	6.385182E-01	8.095848E-01	1,650323E-03
11	-31651104E-01	4479049E-01	8,963955E-01	1,5025546+03
12	-2,766214E-01	6,443928E-01	9.871282E-01	1,509524E+03
13	-1,841325E-31	6,381041E-01	1.075840E+00	1,464486E+03
		- 6,847973E-01	11104393E+00	-11392030E-03
15	•1.115454E-02	6.325974E-01	1.2529082.00	1,319565k*03
19	7.733442E-02	6,317235E-01	1.341401E+00	1,2420746-03
	11650254E-01	-6,5252326-01-	114290946+00	1,1700012003
10	2.543124E-01	6,344533E-01	1,5184046+00	1.094832E+03
14	3.428013E-01	6,3826856-01	1.606975E+00	1,023362E+03
20 21	4,3129032-U1 6,958348E-G1	0,420011E-U1	119333975400	7 5205045002
22		6,425414E-01	1,9601265+00	7,520504E+02
	7.157167E-01	6,494937E-01	1,9511896+00 2,1268696 +00	7,413017E+02
24	9,923979E-01	7.40691E-01	2.272506E+00	5,864850E+02
25	1.1309756+30	7.852728E-01		
	115,035,5-00	- 8.300935g-01	2.41005500 2.904660E+00	5.093910E*02
27	1.411687E-00	8.757255E-01	2.713010£*00	3.766447E+02
žė	1,5549246+00	9,223460E-01	2.863643E+00	
	1.7000475-00	-9:479427E-01	-3.017140E+00	3,152831E*02
30	1.8496086+00	1.0183145+00	3.173560E+00	2,298456E+02
31	2.001063E+00	1.065874E+00	3.332307E+00	2.0437684-02
- 35	211972602*00	1,1133602-00	3.4755712700	1,0773178-02
33	2.322368E+00	1.163310E+00	3.668062E+00	1.7278105-02
34	2.497280E-00	1.215889E+00	3.850704E+00	1,5851916+02
	21484490E+00	1.271702E+00	4.0460882+00	1,4479728+02
36	2.865954E+30	1.331547E+00	4.2>6222E+00	1.316152E+02
37	3.104899E-00	1.3960216-00	4,484462£+0C	1,109184E+02
30	3.3447622-00	1,4001428000	4.7343655-00	1,0000035-03
34	3.551928E+00	1.496188E-00	4,943698E+00	9,730349E+01
40	3.824086E+00	1.5632406-00	5.223945E+00	8,4581896+01
41	4.1333802*00	1.6399986-00	9,9425738-00	7.0223938*01
42	4,490515E+00	1.727859E+00	5,910455E+00	6,620898E+01
43	5,058952E+00	1.009191E.00	6,496197E.00	5,354862E.01
44	9,2979178400	119190208000	8.700920E+00	4,9848988601
45	5.5407716-00	1.988783E-00	6.992638E+00	4,522423E*01
44	5.703051E+00	2.079678E-00	7.366147E+00	4.0037726+01
47	6.2472342-00	2.1050335.00	7.7227162-00	3,5850928-01
40	6,651287E+00	2.265316E-00	8,137068£+00	3,1652566+01
v 47	7.055885E+00	2.365294E+00	8.553836E+00	2,8115206+01
▼ 31	4 30-009-11		A	

4.4272348*01 2.1180086*01 1.7497736*01 1.4062948*01				
7.0294666.02 1.0096066.03 1.0096046.03 1.0096046.03				
2.003063E-00 2.740616E-00 2.740616E-00				
/.52320096-00 6.0120096-00 8.5517136-00 9.1649748-00			**************************************	
> 4 10 10 11 10 10				

		(4)= 60,000		THE TOTAL HAS	FLOH
	ZINIT =	0.00000 2053:317 02		ONAL - B/RC)	
	TINIT .	6158 - 41207	4 (DEG, R)		
	VINIT .	1667.04861	5 (FT/8EC)	ale de la colonia	
	FENAL =	9.07204		ONAL - Z/RC1	111761
	naner -	51	THU! OF PRE	SSURE TABLE PO	JATO/
	POINT	AXIAL	RADIAL	STREAMLINE	PRESSURE
	NO.	DISTANCE	DISTANCE	DISTANCE	(PSIA)
		12/RC)	(R/RC)	18/90)	
	1 •1	250000E+30	9,420203E-01	0.	2,0533176+03
		1919115-00	\$1003\$00E-01	4.5150025.05	5.0141.004.02
	3 -1	073022E+00 845331E-01	9,4138896-01	1,840790E-01 2,757930E-01	1,700745603 1,935760603
		740442E-01	- 0,743767E+01	-3,471435E-01	-1,004968e03
		075552E-G1	8,735829E-01	4,500934E-01	1,8272356-03
		,170662E=01 ,305778E =01	8,543783E-01	5,486423E-01	1,7422056+03
		4208836-01	8,141964E-01	7,2 ⁹ 6032E-01	1,7147426+03
	10 -4	335994E-01	8.047424E-01	8,1882796-01	1,6420795+03
	11 -3	451104E-01	7,907938E-01	9:070011E-01	112022556-03
	12 •2 10 •1	766214E-01 881325E-01	7,863383E-01 7,811676E-01	9,967693E-01 1,085548E-00	1,440001 0 03 1,442231 6 +03
		744890E-32	7,7699686-01	1,1741376+00	1,3408936+03
	17 41	115454E-02	7.749157E-01	1.262660E+00	1,2774408-03
		733442E-02	7,739161E-01	1.351151E+00	1,1737076-03
		 450234E-01 543124E-01	7,754934E-01 7,787151E-01	1,4 39654 6+ 00	1,1184716+03 1,0355356+03
	19 3	428013E-C1	7.842467E-01	1,616863E+00	9,610086E+02
	20 4	13127n3E-01	719119418-01	1.7030515.00	8,81899889B
		404510E-01 831918E-01	7,948469E-01 8,089020E-01	1,914815E+00 1,959684E+00	7,1717056+02 4,9389816+02
		203237E-01	-0,519107E-01	21103555E+00	+,094299E+02
	24 9	384595E-01	8,971486E-01	2.249067E+00	5,219971E+02
	25 1	098846E+00	9,449184E-01	2.397169E+00	4,3654008-02
		3871198400	1.0501216-00	2.764047E+00	3,0309406+02 3,0112356+02
	28 1	936511E+00	1.1081246+00	2.864304E+00	2,583525E+02
	29 1	409195E+30	1,1007296-00	3,027810E+00	2,3023212+02
		845097E+00 004881E+00	1.224722E+00 1.282237E+00	3,194188E+00	2,20040\$E+02
	32 2	1944955-38	71228g46g+00	3.364000E+00	2.028544E*02
	33 2	339102E+00	1.397222E-00	3.717460E+00	1.7150426+02
	34 20	515892E+00	1.4558296-00	3.903712E+00	1.571420E+02
	36 2	703440E+00	1.563449E+00	4.311067E+00	1.3025476+02
1	37 3	116761E*00	1.653752E+00	4.536339E+00	1.1776335-02
·	30 31	3477992-00	117294305+80	41777478500	210348648405
		599726E+00	1.6117208+00	5.044482E+00	9,475904E+01
		876663E+00	1.901948E-00	5.335747E+00 5.371143E+00	8.4111168*01 7.6498718*01
	42 4	421066E+38	2.028198E+00	5.895324E+00	6.749101E+01
	43 41	776991E+00	2.127592E+00	6.264866E+00	5.4843228-01
	45 5	189426E+30 849836E+00	2.2434978490	7. 779945-00	3,0300138-01
		181545E+00	2.431164E+00 2.497727E+00	7.379841E+00 7.620915E+00	4,019394E+01 3,722404E+01
	47 61	4109792400	2.3911378000	7,9633382+00	3,3941058-01
	48 61	835618E+00	2.714446E+00	8.405517E+00	2,8557396+01
	49 7,	245205E+60	2.030023E+00	8,833339E+00	2,293517601

91 98	7.7307X>E-UD 8.230201E+00	3.1479576-00	9,872043K+08	1.3294146-01
ZINIT 4		O CHONDINENS!	THE TOTAL MASS	FLOW
TINIT	6171.8094	(DEG, R)		
HOKSK .	8:5274	7 INONDIMENS	ONAL - Z/RG) SSURE TABLE PO	INTS)
POINT NO ₄	AXIAL DISTANCE (2/RC)	RADIAL DISTANCE (R/RC)	STREAMLINE DISTANCE (S/RC)	PRESSURE (PSIA)
<u>.</u>	-1.250000E+06	1,1494025-00	0.	2,0411536+03
3 4	-1:073022E-00 -9:049331E-01	1.130922-00 1.097606E-00 1.066449E-00	1.876254E-01 2.434392E-01	2,0417738*03 1,9766736*03 1,9460946*03
• • • • • • • • • • • • • • • • • • • •	-0,909428-01 -0,079592E-01 -7,190662E-01	1,037444200 1,0114026+00 9,8773846-01	3,764929R+01 4,607962E-01 5,603946E-01	1,8892118+03 1,8295076+03 1,7942496+03
•	-5.420663E-01	9,463754E-01	7.423069k=01	1.7115766+03
10 11 12	-4.539994E-01 -3.691184E-81 -2.766214E-01	9.317907E-01 9.173739E-01 9.073322E-01	0.319962E-01 7.213469E-01 1.010403E-00	1,6279076°03 1,9392698°03 1,4496636°03
13	-1.881325E-01 -1.115454E-02	9.0193976-01 8.9471918-01 8.9431376-01	1,099239E+00 1,187899E+00 1,276377E+00	1,4099736°03 1,310938°03 1,2279996°03
16	7.733442E-02 11 99234E- 01	8,9435376-01 8,9704226-01	1,364866E+00	1,1363806*03 1,0961408*03
19	2,543124E-01 3,425013E-01 4,3127032-01	9.0202716-01 9.0972396-01 9.1997012-01	1,542025E+00 1,630848E+00 1,714863E+00	9,662164E*02 8,871731E*02 8,019944E*02
25 52 51	5.822534E-01 6.339078E-01 6.784233E-01	9,294423E-01 9,435143E-01 +,9442982-01	1.871182E+00 1.924744E+00 1.909719E+00	6,7510316°02 6,4118986°02 6,1512478°02
24 25	7.059375E-01 8.849062E-01	9,441490E-01 1,021289E-00	1.999595E+00 2.167464E+00	9,933811E+02 4,6762996+02
27	1.170395E+00 1.317004E+00	1.1263076.00 1.1263076.00	2:33/1342*00 2:491642E*00 2:450235E*00	3,8287168002 3,104979602 2,860143602
30 31	1.4877472-00 1.420744E-00 1.777414E-00	1.2434978 · UU 1.3027148 • 00 1.36266 • 600	2.975370E+00 3.143107E+00	2,447127E+02 2,447127E+02 2,256445E+02
32 33 34	2.104198E+00 2.275866E+00	1,4239496+00 1,4894296+00 1,9486426+00	3,3192192 + 00 3,492132E + 00 3,675188E + 00	2.0797998-02 1.906349E-02 1.747092E-02
39	2,454364E+00 2,440553E+00	1,614045E+00	3,8692232+00 4,0631712+00	1,4940338+02
37 30 39	2.836097E+00 3:0440962+00 3:265661E+00	1,751344E+00 1,8258y02+00 1,905023E+00	4,270897E+00 4,471823E+00 4,727123E+00	1,322361802 1,197027802 1,080707802
40 48 42	3.904190E+00 3.762350E+00	2:0422722-00	4.980414E+00 5.254498E+00	9.7002 758 +01
48	4.044566E+00 4.355551E+00 4.000107E+00	2,293656E+00 2,293656E+00	5,554105E+00 5,884230E+00	7,672922E-01 6,749932E-01 -,133253E-01
49	4,957402E+00 5,354126E+00 5,819216E+00	2,4483986+00 2,5713736+00 2,7149838+00	6,904607E+00 6,923866E+00 7,408707E+00	5,3746186*01 4,6519686*01 3,9632608*01
40	6.584686E+00 6.868650E+00	2,984964E+00 3,101444E+00	8,220397E+00 8,927407E+00	1.981126E+01 1.990489E+01

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Appendix C

KINCON

NONEQUILIBRIUM CHEMICAL KINETICS AND CONDENSATION COMPUTER PROGRAM

A Multiphase Reacting Gas Streamtube Model

Program Number H360

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LIST OF ABBREVIATIONS AND SYMBOLS

A, a	Cross-sectional area of the streamtube
aj, b	Reaction-rate parameters
C _p	Specific heat at constant pressure
c	Mass fraction of i th species
$\mathbf{f_i}$	Free energy of ith species
g	Mass fraction of condensed phase
н	Energy addition rate, per unit normalized length, per unit initial streamtube mass flux
h	Enthalpy
h'	Enthalpy of condensed phase
J	Nucleation rate, critical-size nuclei formed per unit volume, per unit time
k	Boltzmann constant
К _j	Equilibrium constant
k, k	Reaction-rate parameters
M	Momentum flux, per unit normalized length, per unit initial streamtube mass flux
Mi	Third-body reaction term
\overline{M}_{i}	Chemical symbol for the ith species
m	Mass addition rate, per unit normalized length per unit initial streamtube mass flux
m _{ji}	Reaction-rate ratio
m _i	Molecular weight of ith species
Na	Avogadro's number
n	Total number of species

Reaction-rate parameters 'nj Pressure and partial pressure of ith species, respectively P, P, R Gas constant of the mixture Universal gas constant R Normalization factor for x Critical droplet radius P** ith species mass addition rate, per unit normalized length, per unit initial streamtube mass flux Š, T Temperature V Flow velocity Species production-rate term X_{i} Axial distance Y Ratio of specific heats Surface tension Condensation coefficient Stoichiometric coefficients ν_{ii}, ν'_{ii} Density Production rate of ith species

SUBSCRIPT

D Droplet

L Liquid

V Vapor

VS Saturated vapor

i i i species

j th reaction

Appendix C

KINCON

NONEQUILIBRIUM CHEMICAL KINETICS AND CONDENSATION COMPUTER PROGRAM

A Multiphase Reacting Gas Streamtube Model

C. 1 INTRODUCTION

w s

The computer program described in this appendix is a subprogram to the Plume Contamination Effects Prediction Computer Program, CONTAM, and performs chemical-kinetic and single-species condensation calculations along gas-phase streamlines as computed by the Multiphase Nozzle and Plume Transport Computer Program, MULTRAN. KINCON may also be used as an independent computer program on any third-generation computer with a core exceeding 120,000 words and a Fortran IV processor.

The present computer program is based on the ICRPG One-Dimensional Kinetic Nozzle Analysis Computer Program, ODK (References C-1 and C-2), and on modifications to the ODK by Dynamic Science Corporation under contract to McDonnell Douglas Astronautics Company (Reference C-3). The purpose of the original program was to provide an automated engineering tool for the kinetic analysis of one-dimensional chemically reacting gas systems. To this end, a number of options were included in the program to aid the user. These included a mass, momentum, and energy streamtube addition option, generalized oblique shock calculation, normal shock-stagnation streamline calculation, area-defined streamtube option, and a reaction screening option. Modifications performed under the present study include the addition of a thermodynamic nonequilibrium condensation model and the automation of the program to perform successive kinetics and condensation calculations for a series of streamlines.

Species and reactions are input to the program in symbolic form. The user may input arbitrary species (up to 40) and arbitrary gas phase reactions (up to 150). Specified third-body reaction rate ratios may be employed. A comprehensive library of thermochemical data is available as part of the computer program. This data may be expanded by input of tables punched directly from the JANAF format. Automatic plotting of temperature, density, and species concentration is available. A unique feature of the program is its ability to integrate—with complete numerical stability—the differential equations governing the kinetic system.

Section C. 2 contains a discussion of physical assumptions. The equations governing the inviscid, one-dimensional flow of a chemically reacting gas mixture are given in the form in which they are integrated in the computer program.

Section C. 3 contains a discussion of the integration method used in the computer program.

Section C. 4 contains a description of the program overlay structure.

Section C. 5 contains a detailed engineering and programming description of the logic and the calculations performed in the computer program.

Section C. 6 contains a program user's manual describing the use of the computer program with an explanation of the program input and output.

Section C. 7 contains input and output for a sample case.

C. 2 ANALYSIS

The method of solution consists of integrating the conservation equations for the chemical system in such a form that the chemistry is generalized for binary exchange and dissociation-recombination reactions. Condensed phase products are considered by a single-species nonequilibrium condensation option.

a. Conservation Equations

The <u>KINCON</u> computer program integrates a set of simultaneous differential equations along a pressure-defined streamtube (i.e., P(x) and dP(x)/dx are known). These differential equations represent the conservation of species, mass, momentum, and energy for the system as expressed by Equations (C-5), (C-6), (C-7), and (C-8) below.

The conservation equations governing the inviscid flow of reacting gas mixtures have been given by Hirschfelder, Curtiss, and Bird (Reference C-4), Penner (Reference C-5), and others. The following basic assumptions are made in the derivation of these equations.

- 1. Mass (m, si), momentum (M), and energy (H) addition rates are defined for the system.
- 2. The gas is inviscid.
- 3. Each component of the gas is a perfect gas.
- 4. The internal degrees of freedom of each component of the gas are in equilibrium.

In one-dimensional flow, the conservation equations have the form 1

species
$$\frac{d}{dx} \left[(1 + \overline{m}) c_i \right] = \dot{s}_i + (1 + m) \frac{\omega_i r^*}{\rho V}$$
 (C-1)

mass
$$\frac{d}{dx}(1+\overline{m}) = \dot{m}$$
 (C-2)

momentum
$$\frac{d}{dx} [(1 + \vec{m}) V] = \dot{M} - \frac{(1 + \vec{m})}{\rho V} \frac{dP}{dx}$$
 (C-3)

The independent variable, x, is taken as unitless with r* as the conversion factor to units. The quantity $1 + \overline{m}$ represents the streamtube mass flux normalized by the initial streamtube mass flux; i.e., $1 + \overline{m} = (\rho Va)/(\rho Va)$

$$\frac{d}{dx}\left[(1+\overline{m})h_{T}\right] = \dot{H},$$

$$h_T = \sum_{i=1}^{n} c_i h_i + \frac{v^2}{2}$$
 (C-4)

If the expansion process is specified by the pressure distribution as a function of distance, Equations (C-1 through C-4) can be written as

$$\frac{dc_i}{dx} = \frac{\dot{s}_i - \dot{m}c_i}{1 + \overline{m}} + \frac{\dot{u}_i r^*}{\rho V}$$
 (C-5)

$$\frac{\mathrm{dV}}{\mathrm{dx}} = \frac{\dot{M} - \dot{m}V}{1 + \overline{m}} - \frac{1}{\rho V} \frac{\mathrm{dP}}{\mathrm{dx}} \tag{C-6}$$

$$\frac{dT}{dx} = \frac{1}{C_p} \left[\frac{\dot{H} - \dot{m} h_T}{1 + \overline{m}} - \frac{V (\dot{M} - \dot{m}V)}{1 + \overline{m}} + \frac{1}{\rho} \frac{dP}{dx} - \sum_{i=1}^{n} h_i \frac{dc_i}{dx} \right] (C-7)$$

$$\frac{d\rho}{dx} = \left[\frac{1}{P}\frac{dP}{dx} - \frac{1}{T}\frac{dT}{dx} - \frac{1}{R}\left(\sum_{i=1}^{n} R_{i}\frac{dc_{i}}{dx}\right)\right]\rho \tag{C-8}$$

where

$$C_{\mathbf{p}} = \sum_{i=1}^{n} c_{i} C_{\mathbf{p}i}$$
 (C-9)

$$Y = \frac{C_p}{(C_p - R)} \tag{C-10}$$

$$h_i = \int_0^T C_{pi} dT + h_{i0}$$
 (C-11)

For each component of the gas, the equation of state is

$$P_{i} = \rho_{i}R_{i}T \qquad (C-12)$$

Summing over all the components of the mixture, the overall equation of state is obtained

$$P = \rho RT = \rho T \sum_{i=1}^{n} c_i R_i \qquad (C-13)$$

The net species production rate ω_i for each species (component) is calculated from

$$\omega_{i} = \overline{m}_{i} \rho^{2} \sum_{j=1}^{n} (\nu'_{ij} - \nu_{ij}) X_{j}$$
 (C-14)

where

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$$X_{j} = \left[K_{j} \prod_{i=1}^{n} c_{i}^{\nu_{ij}} - \rho^{\lambda} \prod_{i=1}^{n} c_{i}^{\nu'_{ij}}\right] k_{j} M_{j}$$
 (C-15)

and λ depends on the order of the reaction and M, is calculated only for dissociation-recombination reactions.

The equilibrium constant, K, is

$$K_{j} = e^{-\Delta F/RT}$$
(C-16)

$$\Delta F = \sum_{i=1}^{n} f_{i} v_{ij} - \sum_{i=1}^{n} f_{i} v'_{ij}$$

The computer program considers chemical reactions defined by the generalized chemical reaction equation

$$\sum_{i=1}^{n} \nu_{ij} \overline{M}_{i} = \sum_{i=1}^{n} \nu'_{ij} \overline{M}_{i}$$
 (C-17)

where v_{ij} and v_{ij} are the stoichiometric coefficients to be used in Equation (C-15) while \overline{M}_i represents the symbol for the i^{th} chemical species.

The reaction rates, k_j , for the j^{th} reaction appearing in Equation (C-15) are represented in the Arrhenius form

$$k_{j} = a_{j}T^{-n}_{j} e^{(-b_{j}/\mathcal{R}T)}$$
 (C-18)

where

a; is the pre-exponential coefficient

n; is temperature dependence of the pre-exponential factor

b, is the activation energy

Since each dissociation-recombination reaction has a distinct reaction rate associated with each third body, the net production rate for each dissociation-recombination reaction should be calculated from

$$X_{j} = \sum_{k=1}^{n} \left[K_{j} \prod_{i=1}^{n} c_{i}^{\nu_{ij}} - \rho \prod_{i=1}^{n} c_{i}^{\nu_{i}}^{ij} \right] c_{k}^{k_{kj}}$$
 (C-19)

rather than Equation (C-15). However, Benson and Fueno (Reference C-6) have shown theoretically that the temperature-dependence of recombination rates is approximately independent of the third body. Assuming that the temperature dependence of recombination rates is independent of the third body, the recombination rate associated with the kth species (third body) can be represented as

$$k_{kj} = a_{kj}^{-n} T^{-b_j/RT}$$
 (C-20)

where only the constants a_{kj} are different for different species (third bodies). From Equation (C-19) it can be shown that

$$X_{j} = \left[K_{j} \prod_{i=1}^{n} c_{i}^{\nu_{ij}} - \rho \prod_{i=1}^{n} c_{i}^{\nu'_{ij}}\right] \left[\sum_{i=1}^{n} \frac{a_{ij}}{a_{kj}} c_{i}\right] a_{kj}^{-n_{j}} e^{-b_{j}/RT} \quad (C-21)$$

Thus, the recombination rates associated with each third body can be considered as in Equation (C-15) by calculating the general third body term $M_{\dot{j}}$ as

$$M_{j} = \sum_{i=1}^{n} m_{ji} c_{i}$$
 (C-22)

where m_{ji} is the ratio a_{ij}/a_{kj} .

In order to numerically integrate Equations (C-1), (C-5), (C-6), and (C-7), it is necessary to input the following type of information concerning the chemical system:

Boundary Conditions:

x initial axial position

P initial pressure

To initial temperature

V Initial velocity

x max final axial position

P(x) table of pressure versus axial position

dP(x)/dx table of pressure derivatives versus axial position

Species Information:²

 \overline{M}_i species name

m, species molecular weight

 $C_{pi}(T)$ species specific heat

h;(T) species enthalpy

 $f_i(T)$ species free energy

The items C_{pi} , h_i , and f_i are not available directly in the appropriate units. The computer program calculates these items for each species from the JANAF data for: C_p vs T

$$-\frac{{\rm H}^{\circ} - {\rm H}^{\circ} {\rm 298}}{{\rm T}} {\rm vs} {\rm T}$$

$$-\frac{{\rm H}^{\circ} - {\rm H}^{\circ} {\rm 298}}{{\rm T}} {\rm vs} {\rm T}$$

$$+{\rm H}^{\circ}_{f}$$

and

Reaction Information:

$$\sum_{i=1}^{n} \nu_{ij} \overline{M}_{i} \ \rightleftharpoons \ \sum_{i=1}^{n} \nu_{ij}^{\dagger} \overline{M}_{i} \quad \text{each reaction must be input in terms of its stoichiometric coefficients and constituent species}$$

$$a_{j}, \ n_{j}, \ b_{j} \quad \text{constants defining } k_{j}, \ \text{the reaction rates}$$

$$m_{ij} \quad \text{third body reaction-rate ratios}$$

Miscellaneous Information:

r* normalization factor for x

Mass, Momentum, Energy, and Species Addition Functions:

- m (x) mass addition rate, per unit normalized length, per unit initial streamtube mass flux
- M (x) Momentum flux, per unit normalized length, per unit initial streamtube mass flux
- H (x) energy addition rate, per unit normalized length, per unit initial streamtube mass flux
- ith species mass addition rate, per unit normalized length, per unit initial streamtube mass flux

A considerable amount of data (such as JANAF tables defining $C_{\mathbf{pi}}$, h_i , and f_i , reaction rate parameters and cards defining chemical reactions) are available with the computer program. Details concerning input to the computer program are given in Subsection C.6.

For the area defined option including mass, energy, and momentum addition, the pressure profile is obtained by an iteration to obtain the pressure profile such that A(x) obtained A(x) input A(x) input A(x) input

b. Condensation Equations

Dropwise condensation of a single gaseous species is computed from classical liquid drop theory. In addition to the assumptions noted above, the condensation analysis assumes the following:

- 1. Condensed phase mass is uniformly distributed.
- 2. Droplets are spherical.
- 3. Droplets are small and follow gas streamlines.
- 4. Volume occupied by condensed phase is small compared to gas volume.

Two distinct processes are treated, nucleation and droplet growth. The nucleation process (spontaneous self-nucleation) occurs in the expanding supersaturated vapor and involves the clustering of vapor molecules to give rise to very small nuclei (radius of 10 to 100 Å). Only nuclei reaching the critical drop radius r** can exist and grow. Critical drop radius is given by Frenkel (Reference C-7)

$$r^{**} = \frac{2\sigma}{\rho_L R_v T \ln\left(\frac{p_v}{p_{vs}}\right)}$$
 (C-23)

The nucleation rate, J, represents the number of critical-size nuclei formed per unit volume per unit time and is calculated from the expression by Stever (Reference C-8)

$$J = \left(\frac{p_{V}}{kT}\right)^{2} \frac{1}{\rho_{L}} \left(\frac{2\sigma m_{i}}{\pi Na}\right)^{1/2} exp \left(-\frac{4\pi\sigma r^{**2}}{3 kT}\right)$$
 (C-24)

Once a suitable number of nuclei are formed in the vapor, the process of droplet growth accounts for the actual condensation. Droplet growth occurs through the collision of vapor molecules and stable liquid droplets.

The net flux of vapor to the droplet surface is computed from kinetic theory considerations where droplets are typically smaller than the mean free path of the gas. The droplet growth equation of Hill (Reference C-9) is utilized

$$\frac{d\mathbf{r}}{d\mathbf{x}} = \frac{\eta}{\rho_L V} \frac{1}{(2\pi R_V)} 1/2 (\beta - \beta_D)$$
 (C-25)

where

$$\beta = \frac{P_v}{T^{1/2}}$$

$$\beta_{\rm D} = \frac{p_{\rm D}}{T_{\rm D}^{1/2}}$$

Droplet temperature is assumed to be that of the saturated vapor.

The appropriate mass, momentum, and energy addition rates are computed internally from the following expressions

$$\dot{m} = -\frac{dg}{dx}$$

$$\dot{N} = \dot{m}V + g \frac{(1 - \overline{m})}{\rho V} \frac{dp}{dx}$$
Momentum
$$\dot{H} = (1 - 2g) \dot{m}h' - (1 + \overline{m}) g \frac{dh'}{dx}$$
Energy
$$\dot{S}_{i} = \dot{m}, i = \text{condensing species}$$

$$= 0 \text{ for other species}$$

The rate-of-change of condensed-phase mass fraction, dg/dx, is evaluated by summing the mass of all droplets formed upstream of a specific location, x, as follows

$$\frac{\mathrm{d}\mathbf{g}}{\mathrm{d}\mathbf{x}} = \frac{4\pi\rho_{\mathbf{L}}}{\rho V} \left[\frac{1}{3} \mathbf{r}^{**}(\mathbf{x}) J(\mathbf{x}) \frac{A(\mathbf{x})}{A_{o}} + \frac{\mathrm{d}\mathbf{r}}{\mathrm{d}\mathbf{x}} \int_{\mathbf{x}_{o}}^{\mathbf{x}} \mathbf{r}(\mathbf{x}, \xi)^{2} J(\xi) \frac{A(\xi)}{A_{o}} \mathrm{d}\xi \right]$$
(C-26)

where the integral is replaced by a summation for numerical evaluation.

C.3 NUMERICAL METHOD

It has been shown (e.g., Reference C-10) that explicit methods of numerical integration are unstable when applied to relaxation equations [such as Equations (C-1), (C-5), (C-6), and (C-7)], unless the integration step size is of the order of the characteristic relaxation distance. Since in the near equilibrium flow regime the characteristic relaxation distance is typically many orders of magnitude smaller than characteristic physical dimensions of the system of interest, the use of explicit methods to integrate relaxation equations often results in excessively long computation times. An implicit integration method which is inherently stable in all flow situations (whether near equilibrium or frozen) is therefore used by the computer program. With this method, step sizes which are of the order of the physical dimensions of the system of interest can be used, reducing the computation time per case by several orders of magnitude when compared with conventional explicit integration methods.

Consider N first-order simultaneous differential equations

$$\frac{dy_i}{dx} = f_i(X, y_i, ..., y_N)$$
 $i = 1, 2, ..., N$ (C-27)

with known partial derivatives (i. e. the Jacobian for the system)³

$$a_{i} = \frac{\partial f_{i}}{\partial x}$$
 (C-28)

$$\beta_{i,j} = \frac{\partial f_i}{\partial y_i}$$
 (C-29)

The following implicit difference equations are used by the computer program to determine the $y_{i, n+1}$, (the subscript n denotes the n^{th} integration step)

$$y_{i, n+1} = y_{i, n} + k_{i, n+1}$$
 $h = x_{n+1} - x_n$ (C-30)

where

$$k_{i, n+1} = \left[f_{i, n} + \alpha_{i, n} h + \sum_{j=1}^{N} \beta_{i, j, n} k_{j, n+1} \right] \cdot h$$
 (C-31)

for the initial step and for restart (first order).

$$k_{i, n+1} = \frac{1}{3} \left[k_{i, n} + 2 \left(f_{i, n} + \alpha_{i, n} h + \sum_{j=1}^{N} \beta_{i, j, n} k_{j, n+1} \right) \cdot h \right]$$
 (C-32)

 $^{^3}$ The computer program uses analytic expressions for calculation of the partial derivatives, $\alpha_{\underline{i}},~\beta_{\underline{i}\underline{j}}.$

for equal steps (second order with h = previous h)

$$k_{i, n+1} = \frac{h_{n+1}^{2}}{(2h_{n+1} + h_{n}) \cdot h_{n}}$$

$$\left[k_{i,n} + \left[f_{i,n} + \alpha_{i,n} h_{n+1} + \sum_{j=1}^{N} \beta_{i,j,n} k_{j,n+1} \right] \cdot \frac{h_{n}}{h_{n+1}} (h_{n+1} + h_{n}) \right]^{(C-33)}$$

for unequal steps (2nd order with h # previous h)

A derivation of these equations is given in Reference C-2.

If the flow is frozen, the explicit form of the above equations can be used ($_{i}$ = 0, β_{ij} = 0); i.e., Equations (C-31), (C-32), and (C-33) are each reduced from an NXN system of linear simultaneous equations to N explicit equations (N = 3 + number of species).

Control of the integration step size, h, is provided by calculating estimates for the truncation error and comparing these to an input criterion, δ .

The step size is halved if for any i = 1, 2, ..., N

$$E_i > \delta$$

The step size is doubled if for all i = 1, 2, ..., N

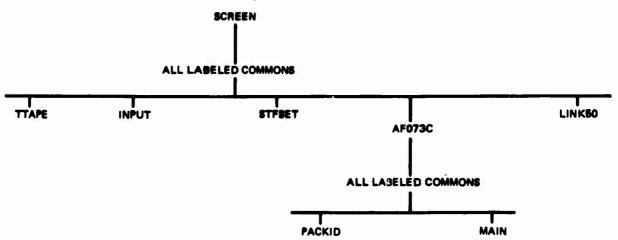
$$E_i < \frac{\delta}{10}$$

where

$$E_{i} = \left| \frac{k_{i, n+1} - 2k_{i, n} + k_{i, n-1}}{3k_{i, n+1} - k_{i, n}} \right|$$
 (C-34)

The above expression for E; is derived in Reference C-2.

C.4 CELL LOADER STRUCTURE



C. 5 PROGRAM SUBROUTINES

This section contains a description of the program subroutines, and these commentaries are given in the order in which the subroutines appear.

The main program sets up master limits for the chemistry, initializes certain logical control variables, and calls subroutine DRIVER. This subroutine provides the overall logic control for the subprogram.

After the program is loaded, a master JANAF Thermochemical Data tape is prepared either from card input or by summary of the current master file in use.

a. Initialization Subroutines Description

(1) Program SCREEN

This subroutine provides overlay communication, defines the labeled common blocks, sets maximum limits for the chemical tables, and initializes certain logical control variables. It calls subroutine DRIVER.

(2) Subroutine DRIVER

This subroutine performs the overall logic for the program.

(3) Subroutine FIND

Provides indices of the table entries which bracket the value of a current variable. The subroutine saves its place in the table.

(4) Subroutine STOICC

Provides up to ten reactants indices and ten product indices from the master stoichiometric coefficient table.

(5) Subroutine AFØ73C

This subroutine provides linkage between the kinetic packing link and the kinetic expansion computations.

(6) Subroutine GTF

This subroutine computes the effective gas constant, gaseous heat capacity, Y, $\partial Y/\partial T$, $\partial Y/\partial C_i$ from the following formulae:

$$R = \sum_{i=1}^{NSP} C_i \cdot R_i$$

$$C_p = \sum_{i=1}^{NSP} C_i \cdot C_{p_i}$$

$$Y = \frac{Cp}{Cp-R}$$

$$\frac{\partial \gamma}{\partial T} = -\frac{\gamma \cdot (\gamma - 1)}{CP} \cdot \sum_{i=1}^{NSP} C_i \cdot \frac{\partial Cp_i}{\partial T}$$

$$\frac{\partial \gamma}{\partial C_i} = \gamma \cdot (\gamma - 1) \cdot \left[\frac{R_i}{R} - \frac{Cp_i}{Cp} \right] \qquad i = 1, \dots, NSP$$

(7) Subroutine SPLN

Performs cubic interpolation for a function and its first two derivatives. Given function values y_n and y_{n+1} and first derivative values y'_n and y'_{n+1} at x_n and x_{n+1} , this subroutine evaluates y(x), y'(x), and y''(x) for $x_n \le x < x_{n+1}$ using:

$$y = A(x - x_n)^3 + B(x - x_n)^2 + C(x - x_n) + D$$

 $y' = 3A(x - x_n)^2 + 2B(x - x_n) + C$
 $y'' = 6A(x - x_n) + 2B$

where:

$$A = \frac{1}{h^3} \cdot [(y'_{n+1} + y'_n) h - 2k]$$

$$B = -\frac{1}{h^2} \cdot \left[(y_{n+1}^{\dagger} + 2y_{n}^{\dagger}) h - 3k \right]$$

$$C = y_n^1$$

$$D = y_n$$

$$h = x_{n+1} - x_n$$

$$k = y_{n+1} - y_n$$

(8) Subroutine STF

Using the SPLN interpolation subroutine, this subroutine computes the heat capacity and its temperature derivatives, enthalpy, and free energy, at the current temperature for all gaseous chemical species.

(9) Subroutine APPROX

This subroutine provides extension of the thermochemical data between the temperatures 9,000 and 20,000 $^{\circ}$ R. A message is provided each time an approximation is calculated. The approximation formulae are given below with X = 9,000 $^{\circ}$ R:

$$C_{\mathbf{P}_{\mathbf{T}}} = C_{\mathbf{P}_{\mathbf{X}}}$$

$$H_T = H_X + C_{P_X} * \Delta T$$

$$F_{T} = F_{X} - \left[\frac{H_{T}^{\circ} - H_{298}^{\circ}}{T} - \frac{H_{T}^{\circ} - H_{298}^{\circ}}{X} - C_{P_{T}} \log \left(\frac{T}{X} \right) \right]$$

(10) Subroutine TTAPE

This subroutine generates a master JANAF thermochemical tape which is subsequently utilized by Subroutine STFSET. The tape is written in the binary mode with the thermodynamic functions in caloric units. The thermodynamic functions for each species include:

Function	Units
c ^b .	cal/mole-°K
H° -H° 298	k-cal/mole
- (F° - H°298)	cal/mole-° K

given at 100 °K temperature increments over the range 100 °K to 5, 000 °K, inclusive. Reference may be made to Subsection C. 6, the Program Users Manual, for a complete description of thermodynamic input format and output options.

(11) Subroutine COLOUT

Provides columnar output of species names for those species residing on the master thermo file.

(12) Subroutine ECNV

This subroutine translates a BCD string of characters, into one floating point numeric value. E, I, and F formats are permitted with the result always a floating point number. It is called by subroutine SPRXIN to decode numeric fields in the species and reactions cards.

(13) Subroutine INPUT

This subroutine performs specific case input for the program. It performs the following functions:

- 1. Variable initialization to nominal values.
- 2. Read title card.
- 3. Call subroutine SPRXIN to input the species and reactions cards.

- 4. Read \$PROPEL namelist for case input data.
- 5. Check input mole or mass fractions for unity (±1.0E-4).
- 6. Read initial conditions and pressure table from Tape 8 when operated in automatic mode.

(14) Subroutine NUMBER

This subroutine converts a one-character BCD number to a FORTRAN integer number. It is called by subroutine ECNV to decode free field numeric data.

(15) Subroutine SPRXIN

This subroutine processes the species and reactions cards. Species symbols, numeric mass or mole fractions, symbolic reactions, and rate parameters are processed. Reference may be made to Subsection C. 6, the Program Users Manual, for a complete description of input requirements.

(16) Subroutine STFSET

This subroutine uses the master JANAF tape written by subroutine TTAPE, to generate a species thermal-function tape (KSTF) in blocked form for the kinetic calculations. The tabulated functions on the master tape are:

<u>Function</u>	Units
$Cp_{i}^{i} = Cp_{i}^{o}$	cal/mole-° K
$H_{i}^{\prime} = [H^{\circ} - H^{\circ}_{298}]_{i}$	kcal/mole
$F'_{i} = -\left[\frac{F^{\circ} - H^{\circ} 298}{T}\right]_{i}$	cal/mole-° K

For the kinetic calculations the above functions must be converted to the ft/sec*R units system by the following:

Function		Conversion	Internal Units
$\mathtt{Cp}_{\mathbf{i}}$	=	[Cp _i '· R _i]/1.98726	ft ² /sec ² -°R
H _i	=	$\left(H'_{i} + \Delta H^{\bullet}_{F_{i}}\right) R_{i} \cdot 905.770$	ft ² /sec ²
$\mathbf{F_i}$	=	F' ₁ /1.98726	unitless

These converted functions are then written in 900°R temperature blocks. The free energy and heat capacity derivatives are computed using the formulae below with $\Delta T = 180.0$.

The function derivatives are computed according to the following formulae:

$$\frac{d\eta}{dT}(i, T_1) = \frac{4 \cdot \eta (i, T_2)^{-3 \cdot \eta} (i, T_1)^{-\eta} (i, T_3)}{2 \cdot \Delta T}$$

$$\frac{d\eta}{dT}(i, T_j) = \frac{\eta (i, T_j + \Delta T)^{-\eta} (i, T_j - \Delta T)}{2 \cdot \Delta T}$$

$$\frac{d\eta}{dT}(i, T_{50}) = \frac{3 \cdot \eta (i, T_{50})^{-4 \cdot \eta} (i, T_{49})^{+\eta} (i, T_{48})}{2 \cdot \Delta T}$$

where η_i may be species heat capacity Cp' or free energy F'.

(17) Subroutine ADDFIT

For mass, energy, momentum and species addition functions, this subroutine calculates addition function tables and their derivatives from the input tables using one of the following options:

- 1. Simple differencing.
- 2. Spline fit.
- 3. Input derivative tables.
- 4. Parabolic fit.

The addition functions are normalized, modified by the appropriate multiplicative constants if required, and output in tabular form.

If single species condensation option is utilized, addition functions are not input but computed internally. No operations are performed by ADDFIT.

(18) Subroutine CONVRT

This subroutine converts input data from the externally input units to internally used computation units. In order to conserve computation time during the kinetic expansion, parameters such as molecular weights are included in these conversions. Primed numbers are input quantities.

(a) Dissociation-Recombination Reaction Rate Ratio

Input units: unitless

Internal units: (lbs-mass/lb-mole)-1

Formula: $XMM_{j, i} = XMM_{j, i}/MW_{i}$

(b) Pre-exponential Reaction Rate Ratio

Dissociation-recombination reactions

Input units: cm, *K, g-mole, sec

Internal units: ft³, *R, lb-mole, sec

$$A_{j} = \frac{A'_{j} \cdot (.0160183)^{\eta} \cdot 1.8^{XN_{j}}}{\prod_{i=1}^{n} MW_{i}^{\nu'} ij}$$

Where η depends on the order of the reaction.

and

$$0.0160183 = \frac{3.531 \cdot 10^{-5} \text{ft}^3}{1 \text{ cm}^3} \cdot \frac{1 \text{ g-mass}}{2.2 \cdot 10^{-3} \text{lb-mass}}$$

(c) Exponential Term

Input units: kcal/mole

Internal units: *R

Formula: $B_{i} = B'_{i} \cdot 905.770$

where

905. 770 =
$$\frac{1000 \text{ cal}}{1 \text{ kcal}} \cdot \frac{1}{1.98726 \text{ cal/mole-}^{\circ}\text{K}} \cdot \frac{1.8^{\circ}\text{R}}{1.0^{\circ}\text{K}}$$

(d) Equilibrium Constant Multiplicative Factor

Input units: not input

Internal units: (lb-mass) - *R/ft3

Formula:

DATEF(J) =
$$\frac{\prod_{i=1}^{n} MW_{i}^{v_{ij}}}{\prod_{i=1}^{n} MW_{i}^{v_{ij}} \cdot 0.73034}$$

where

(e) Heats of Reaction

Input units: Kcal/mole (via heats of formation)

Internal units: *R

Formula: $DELH(J) = DELH(J)' \cdot 905.770$

(f) Pressure

Input units: PSIA

Internal units: poundals/ft²

Formula: $P = P' \cdot 4633.056$

where

$$4633.056 = \frac{144 \text{ in}^2}{1 \text{ ft}^2} \cdot 32.174 \frac{\text{ft}}{\text{sec}^2}$$

(19) Subroutine PACKIP

On the basis of those species currently being considered, this subroutine packs species and reaction information from the master tables into those control sections utilized by the kinetic calculation links.

The following is a description of the subroutine functions:

- a. The symbolic reactions and their input rate parameters are printed.
- b. Reaction mass balance is checked for a tolerance of ±1.0 E-10.
- c. Heats of reaction are computed.

(20) Subroutine PRES

This subroutine provides a pressure table and its derivatives suitable for processing by the kinetic calculation links. For a normal shock stagnation streamline, velocity table and its derivatives are provided in a form suitable for processing by the kinetic calculation links.

(21) Subroutine SLP (X, Y, N, MFLAG, YP, W1, W2, W3, IFLAG)

The purpose of this subroutine is to supply derivatives for a tabulated function. The end point derivatives may be specified or are calculated internally by parabolic interpolation. Interior point derivatives may be found by a cubic spline fit procedure.

Calling Sequence:

- X is a table of independent variables, x
- Y is a table of the dependent variables, y,
- N is the number of entries in each of the tables X, Y, and YP. i = 1, ... N

MFLAG this entry is a flag, m, such that

m > 0 implies x is equally spaced

m < 0 implies x is not equally spaced

m = 1 y' will be continuous

|m| = 2 y' and y'' will be continuous

YP is a table of the derivative, y

W1 working storage of length N

W2 working storage of length N

W3 working storage of length N

IFLAC this entry is a flag, i, such that

i = o implies value for YP(1) and YP(N) will be calculated internally by parabolic differencing

i = 1 implies values for YP(1) and YP(N) will be input

Method:

The cubic spline fit procedure utilizes the interpolation formula given within the description of subroutine SPLN, i.e.:

$$y = A(x - x_0)^3 + B(x - x_0)^2 + C(x - x_0) + D$$

 $y' = 3A(x - x_0)^2 + 2B(x - x_0) + C$
 $y'' = 6A(x - x_0) + 2B$

The piecewise cubic fit to a tubular function by the above relations will yield a discontinuity in the second derivative y", between adjacent fits of:

$$y_{101}^{i} - y_{112}^{i} = \frac{1}{h_{01}} \left(2y_{0}^{i} + 4y_{1}^{i} - 6\frac{k_{01}}{h_{01}} \right) - \frac{1}{h_{12}} \left(6\frac{k_{12}}{h_{12}} - 4y_{1}^{i} - 2y_{2}^{i} \right)$$

where

$$h_{01} = x_1 - x_0$$
 $h_{12} = x_2 - x_1$
 $k_{01} = y_1 - y_0$
 $k_{12} = y_2 - y_1$

The method consists of setting the left-hand side of the above relation equal to zero so that the second derivative is continuous across juncture points. As applied to a tabular function, the above procedure results in a set of linear simultaneous equations (tri-diagonal) to be solved for the y₁, provided that values for y' at the end points are known.

(22) Subroutine TCALC

This is a dummy subroutine to permit the user to generate the addition function tables using his own supplied subroutine. It must be replaced with the appropriate TCALC routine and used in conjunction with the IADD \emptyset P = 2 option.

b. One-Dimensional Kinetic Expansion Subroutines Description

This link contains the one dimensional kinetic expansion subroutines.

The implicit integration method, used to integrate the fluid dynamic and chemical relaxation equations, requires the values of the partial derivatives of all total derivatives with respect to every variable. The program will generate a matrix of partial derivatives such that the entry in the nth row and the mth column is the partial derivative of d [n]/dx with respect to m. This matrix is called BETA(I, J). The velocity, density, and temperature-fluid dynamic variables considered for every case reside in rows 1, 2, and 3 respectively. The chemical species occupy rows 4 through the number of species plus 3. The following notation will be used to denote partial derivatives:

$$\beta(A, B) = \frac{\partial \left[\frac{\partial A}{\partial x}\right]}{\partial B}$$

To facilitate the identification of program variables with the engineering notation, the following format will be used where applicable:

engineering notation-program variable-equation

The program will also generate a matrix which will be solved for the variable increments for each integration step. This matrix will expand or contract, depending on the number of chemical species to be considered.

The total derivatives f_i and partial derivatives β_{ij} have been separated into two components: (1) adiabatic component with no mass, momentum, or species addition, and (2) addition component due to mass, momentum, energy, or species addition. Subroutines DERIV and FLU calculate the adiabatic components and subroutine ADDXXX calculates the addition component. When no mass, momentum, energy, or species addition functions are input, the addition component calculations are bypassed.

For the single-species condensation option, subroutines CØNAD, VAPØR and DRØPS compute the mass, momentum, and energy addition functions used by ADDXXX to calculate addition components.

(1) Subroutine DERIV

This subroutine computes the adiabatic components of total derivatives f_i and the partial derivatives $\beta(i,j)$ for the chemical relaxation equations.

Notation: i = Species subscript

j = Reaction subscript

t = Total number of chemical reactions

m = Number of dissociation-recombination reactions

n = Total number of gaseous species

The generalized chemical reaction which is handled by this subroutine is defined by:

$$\sum_{i=1}^{n} v_{ij} C_{i} = \sum_{i=1}^{n} v'_{ij} C_{i}$$

with:

$$\Psi_{ij} = \nu'_{ij} - \nu_{ij}$$

The reverse reaction rate constant is defined by the equation:

$$k_j$$
 $SK(J) = A_j \cdot T^{-XN_j} \cdot exp(-B_j/T)$

The net production rate for a reaction is given by:

$$X_{j} \qquad X(J) = \left[K_{j} \cdot \prod_{i=1}^{n} C_{i}^{\nu_{ij}} - \rho^{\lambda} \cdot \prod_{i=1}^{n} C_{i}^{\nu'_{ij}}\right] \cdot k_{j} \cdot M_{j}$$

where:

 $\lambda = 1$ for a dissociation-recombination reaction

= 0 for a binary exchange reaction

and

$$M_{j} = \sum_{i=1}^{n} XMM_{j,i}$$
 C for a dissociation-recombination reaction
$$= 1 \qquad \text{for a binary exchange reaction}$$

The net individual species production rate is given by the equation:

$$\frac{dC_i}{dx} \qquad FN(I) = \overline{K}_i \cdot \sum_{j=1}^{\ell} \psi_{ij} \cdot X_j$$

where:

$$\overline{K}_i = (MW_i \cdot \rho \cdot r^*)/V$$

The partial derivatives of the net species production rate with respect to the chemical species, the gas velocity, the gas density, and the gas temperature are:

$$\beta(C_k, C_i)$$
 BT(I, K) = $\overline{K}_i \cdot \sum_{j=1}^{\ell} \frac{\partial X_j}{\partial C_i}$ i = 1, ..., NSP

$$\beta(C_i, V)$$
 PHI(I, 1) = $-\frac{1}{V} \frac{dC_i}{dx}$ i = 1, ..., NSP

$$\beta(C_{i},\rho) = \frac{1}{\rho} \cdot \frac{dC_{i}}{dx} + \overline{K}_{i} \cdot \sum_{j=1}^{m} \frac{\partial X_{j}}{\partial \rho} \quad i = 1, ..., NSP$$

$$\beta(C_i, T)$$
 PHI(I, 3) = $\overline{K}_i \sum_{j=1}^{I} \frac{\partial X_j}{\partial T}$ i = 1, ..., NSP

The equilibrium constants and their temperature derivatives are computed only for dissociation-recombination reactions; those quantities for the binary exchange reactions are computed by products and ratios of the dissociation-recombination reaction equilibrium constants and derivatives.

(2) Subroutine EF

This subroutine computes the dissociation-recombination reaction equilibrium constants and their temperature derivatives from the following formulae:

$$K_{j} = EK(J) = \frac{DATEF(J)}{T} \cdot exp \left[\frac{-\Delta H_{j}}{T} - \sum_{i=1}^{n} Ft_{i} \cdot v_{ij} + \sum_{i=1}^{n} Ft_{i} \cdot v_{ij} \right]$$

$$\frac{dK_{j}}{dT} \qquad DKT(J) = \begin{bmatrix} -\sum_{i=1}^{n} \left(\frac{Ht_{i}}{R_{i}}\right) \cdot \nu_{ij} + \sum_{i=1}^{n} \left(\frac{Ht_{i}}{R_{i}}\right) \cdot \nu_{ij} \\ T \end{bmatrix} \cdot \frac{K_{j}}{T}$$

where:

Ft. = species free energy at the current temperature

Ht; = species enthalpy at the current temperature

 Δ Hj = heat of reaction for the Jth reaction

DATEF(j) = is discussed in Subsection C. 5. e(2)

(3) Subroutine FLU

This subroutine computes the adiabatic component of the total derivatives f_i and the partial derivatives a_i and $\beta_{(i,j)}$ for the fluid dynamic equations. Pressure defined fluid dynamic equations are used. The summation terms, energy exchange term B, the diabatic heat addition term A, the Mach number, and all the partial derivatives of these terms are computed. The pressure and its derivatives are obtained from the pressure table.

For a stagnation streamline calculation, the pressure derivatives are obtained from the relationship:

$$\frac{dP}{dx} = -\rho \cdot V \frac{dV}{dx}$$

where V and dV/dx are defined by input tables.

Notes: $\Phi(i, \ell)$, $\ell = 1, 2, 3$ are defined under Subroutine DERIV $\Phi(i, 1)$, = $\beta(C_i, V)$; $\Phi(i, 2)$ = $\beta(C_i, \rho)$; $\Phi(i, 3)$ = $\beta(C_i, T)$

The following relationships may be helpful:

$$f_i = \frac{dC_i}{dx}$$
; $f_i = \frac{r^* \cdot \omega_i \cdot R \cdot T}{P \cdot V}$; $\frac{dC_i}{dx} = \frac{\omega_i r^*}{\rho \cdot V}$

Computation of the Summation Terms and their derivatives:

First Summation

S1 =
$$\frac{1}{R} \cdot \sum_{i=1}^{n} \frac{dC_i}{dx} \cdot R_i$$

$$\frac{\partial S1}{\partial V} \qquad DS1V \qquad = \frac{1}{R} \cdot \sum_{i=1}^{n} \Phi(i, 1) \cdot R_{i}$$

$$\frac{\partial S1}{\partial \rho}$$
 $DS1R\emptyset$ $= \frac{1}{R} \cdot \sum_{i=1}^{n} \Phi(i, 2) \cdot R_{i}$

$$\frac{\partial S1}{\partial T}$$
 DS1T = $\frac{1}{R} \cdot \sum_{i=1}^{n} \Phi(i, 3) \cdot R_i$

$$\frac{\partial S1}{\partial C_i} \qquad DS1C(I) \qquad = \frac{1}{R} \cdot \left[\sum_{i=1}^{n} \beta(C_j, C_i) \cdot R_j - S1 \cdot R_i \right]$$

$$i = 1, ..., NSP$$

Second Summation

S2
$$= \frac{1}{R \cdot T} \cdot \sum_{i=1}^{n} \frac{dC_{i}}{dx} \cdot h_{i}$$

$$\frac{\partial S2}{\partial V}$$
 DS2V = $\frac{1}{R \cdot T} \cdot \sum_{i=1}^{n} \Phi(i, 1) \cdot h_{i}$

$$\frac{\partial S2}{\partial \rho} \qquad DS2R\emptyset \qquad = \frac{1}{R \cdot T} \cdot \sum_{i=1}^{n} \Phi(i, 2) \cdot h_{i}$$

$$\frac{\partial S2}{\partial T} \qquad DS2T \qquad = \frac{1}{R \cdot T} \cdot \sum_{i=1}^{n} \left[\Phi(i, 3) \cdot h_i + \frac{dC_i}{dx} \cdot Cp_i \right] - \frac{S2}{T}$$

$$\frac{\partial S2}{\partial C_i} \qquad DS2C(I) \qquad = \frac{1}{R} \cdot \sum_{j=1}^{n} \frac{\beta(C_j, C_i) \cdot h_i}{T} - S2 \cdot R_i$$

Computation of the Energy Exchange Term B and its Derivatives:

BB =
$$\frac{Y-1}{V}$$
 · S2

$$\frac{\partial B}{\partial V}$$
 DBBV = $\frac{\gamma - 1}{\gamma} \cdot \frac{\partial S2}{\partial V}$

$$\frac{\partial B}{\partial \rho}$$
 DBBRO = $\frac{Y-1}{Y} \cdot \frac{\partial S2}{\partial \rho}$

$$\frac{\partial B}{\partial T} \qquad \qquad DBBT \qquad \qquad = \frac{\gamma - 1}{\gamma} \cdot \frac{\partial S2}{\partial T} + \frac{S2}{\sqrt{2}} \cdot \frac{\partial \gamma}{\partial T}$$

$$\frac{\partial B}{\partial C_i}$$
 DBBC(I) = $\frac{Y-1}{Y} \cdot \frac{\partial S2}{\partial C_i} + \frac{S2}{Y^2} \cdot \frac{\partial Y}{\partial C_i}$ i = 1, ..., NSP

Computation of the Diabatic Heat Addition Term A and its Derivatives:

$$A \qquad AA \qquad = S1-B$$

$$\frac{\partial A}{\partial V}$$
 DAAV = $\frac{\partial S1}{\partial V} - \frac{\partial B}{\partial V}$

$$\frac{\partial A}{\partial \rho}$$
 DAARO = $\frac{\partial S1}{\partial \rho} - \frac{\partial B}{\partial \rho}$

$$\frac{\partial A}{\partial T}$$
 DAAT = $\frac{\partial S1}{\partial T} - \frac{\partial B}{\partial T}$

$$\frac{\partial A}{\partial C_i}$$
 DAAC(I) = $\frac{\partial S1}{\partial C_i} - \frac{\partial B}{\partial C_i}$ i = 1, ..., NSP

Computation of the Mach Number and its Derivatives:

$$M^2$$
 $XM2$ $= \frac{V^2}{Y \cdot R \cdot T}$

$$\frac{\partial M^2}{\partial V}$$
 DM2V = $\frac{2 \cdot M^2}{V}$

$$\frac{\partial M^2}{\partial T} \qquad DM2T \qquad = -\frac{M^2}{T} - \frac{M^2}{Y} \cdot \frac{\partial Y}{\partial T}$$

$$\frac{\partial M^2}{\partial C_1} \qquad \text{DM2C(I)} \qquad = -M^2 \cdot \left[\frac{\partial Y}{C_i} \cdot \frac{1}{Y} + \frac{R_i}{R} \right] \quad i = 1, \dots, \text{NSP}$$

$$\frac{dV}{dx} \qquad FNX(1) \qquad = -\frac{1}{\rho \cdot V} \cdot \frac{dP}{dx}$$

$$\frac{\partial [FNX(1)]}{\partial x} \quad AL(1) \qquad = -\frac{1}{\rho \cdot V} \cdot \frac{d^2P}{dx^2}$$

$$\beta(V, V)$$
 BETA (1, 1) = $-\frac{1}{V} \cdot \frac{dV}{dx}$

$$\beta(V,\rho)$$
 BETA(1, 2) = $-\frac{1}{\rho} \cdot \frac{dV}{dx}$

The Gas Density derivatives are Computed:

$$\frac{d\rho}{dx} \qquad \qquad FNX(2) \qquad = \rho \cdot \left[\frac{dP}{dx} \cdot \frac{1}{\gamma \cdot P} - A \right]$$

$$\frac{\partial [FNX(2)]}{\partial x} \quad AL(2) \qquad = \frac{\rho}{\gamma \cdot P} \cdot \left[\frac{d^2P}{dx^2} - \left(\frac{dP}{dx} \right)^2 \cdot \frac{1}{P} \right]$$

$$\beta(\rho, V)$$
 BETA(2, 1) = $-\rho \cdot \frac{\partial A}{\partial V}$

$$\beta(\rho,\rho)$$
 BETA(2, 2) = $-\frac{1}{\rho} \cdot \frac{d\rho}{dx} - \rho \cdot \frac{\partial A}{\partial \rho}$

$$\beta(\rho, T)$$
 BETA(2, 3) = $-\rho \cdot \frac{\partial A}{\partial T} - \frac{\rho}{P \cdot v^2} \cdot \frac{\partial Y}{\partial T} \cdot \frac{dP}{dx}$

$$\beta(\rho, C_i) \qquad \text{BETA(2, i+3)} = -\frac{\rho}{\gamma^2 p} \cdot \frac{\partial \gamma}{\partial C_i} \cdot \frac{dP}{dx} - \rho \cdot \frac{\partial A}{\partial C_i} \quad i = 1, \dots, \text{ NSP}$$

The Gas Temperature derivatives are computed:

$$\frac{dT}{dx} \qquad \qquad FNX(3) \qquad = T \cdot \left[\frac{\gamma - 1}{\gamma} \cdot \frac{1}{P} \cdot \frac{dP}{dx} - B \right]$$

$$\frac{\partial [FNX(3)]}{\partial x} AL(3) = \frac{\sqrt{-1}}{\gamma} \cdot \frac{T}{P} \cdot \left[\frac{d^2P}{dx^2} - \left(\frac{dP}{dx} \right)^2 \cdot \frac{1}{P} \right]$$

$$\beta(T, V)$$
 BETA(3, 1) = - T · $\frac{\partial B}{\partial V}$

$$\beta(T,\rho)$$
 BETA(3, 2) = - T · $\frac{\partial B}{\partial \rho}$

$$\beta(T, T) \qquad \text{BETA(3, 3)} \qquad = \frac{1}{T} \cdot \frac{dT}{dx} + T \frac{1}{\sqrt{2} \cdot P} \frac{dP}{dx} \frac{\partial Y}{\partial T} - T \frac{\partial B}{\partial T}$$

$$\beta(T, C_i) \qquad \text{BETA(3, i+3)} = T \cdot \left[\frac{1}{\gamma^2 \cdot P} \cdot \frac{dP}{dx} \cdot \frac{\partial Y}{\partial C_i} - \frac{\partial B}{\partial C_i} \right]$$

$$i = 1, ..., \text{NSP}$$

For an adiabatic area defined calculation, the total derivatives a_i and β_{ij} for the fluid dynamic equations are computed using the following area defined equations:

The area ratio and its derivatives are computed from:

$$a = y^2$$

$$\frac{da}{dx} = 2 \cdot Y \cdot \frac{dY}{dx}$$

$$\frac{d^2a}{dx^2} = 2 \cdot \left[y \frac{d^2y}{dx^2} + \left(\frac{dy}{dx} \right)^2 \right]$$

where Y, dY/dx, d^2Y/dx^2 are computed via interpolation in the table of derivatives of the input wall table generated in Subroutine SLP.

The Gas Velocity derivatives are computed:

$$\frac{dV}{dx} \qquad \qquad FNX(1) \qquad \qquad = \frac{V}{M^2 - 1} \cdot \left[\frac{1}{a} \frac{da}{dx} - A \right]$$

$$\frac{\partial [FNX(1)]}{\partial x} \quad AL(1) \qquad = \frac{V}{M^2 - 1} \cdot \frac{1}{a} \cdot \left[\frac{d^2 a}{dx^2} - \frac{1}{a} \left(\frac{da}{dx} \right)^2 \right]$$

$$\beta(V, V) \qquad \text{BETA}(1, 1) \qquad = \frac{1}{V} \cdot \frac{dV}{dx} - \frac{1}{M^2 - 1} \cdot \frac{dV}{dx} \cdot \frac{\partial M^2}{\partial V} - \frac{V}{M^2 - 1} \cdot \frac{\partial A}{\partial V}$$

$$\beta(V, \rho)$$
 BETA(1, 2) = $-\frac{V}{M^2-1} \cdot \frac{\partial A}{\partial \rho}$

$$\beta(V, T)$$
 BETA(1,3) = $-\frac{1}{M^2-1} \cdot \frac{dV}{dx} \cdot \frac{\partial M^2}{\partial T} - \frac{V}{M^2-1} \cdot \frac{\partial A}{\partial T}$

$$\beta(V, C_i) \qquad \text{BETA}(1, i+3) = -\frac{1}{M^2 - 1} \cdot \frac{dV}{dx} \cdot \frac{\partial M^2}{\partial C_i} - \frac{V}{M^2 - 1} \cdot \frac{\partial A}{\partial C_i}$$

$$i = 1, \dots, \text{NSP}$$

The Gas Density derivatives are computed:

$$\frac{d\rho}{dx} \qquad \qquad \text{FNX(2)} \qquad = -\rho \cdot \left[\frac{M^2}{M^2 - 1} \cdot \left(\frac{1}{a} \cdot \frac{da}{dx} - A \right) + A \right]$$

$$\frac{\partial [FNX(2)]}{\partial x} \quad AL(2) \qquad = -\rho \cdot \frac{M^2}{M^2 - 1} \cdot \frac{1}{a} \cdot \left[\frac{d^2a}{dx^2} - \frac{1}{a} \left(\frac{da}{dx} \right)^2 \right]$$

$$\beta(\rho, V) \qquad \text{BETA(2, 1)} \qquad = \quad \rho \cdot \left[\frac{1}{(M^2 - 1)^2} \cdot \left(\frac{1 \text{ da}}{\text{a} \text{ dx}} - A \right) \cdot \frac{\partial M^2}{\partial V} \right]$$

$$+ \frac{1}{M^2 - 1} \cdot \frac{\partial A}{\partial V}$$

$$\beta(\rho, \rho)$$
 BETA(2, 2) = $\frac{1}{\rho} \cdot \frac{d\rho}{dx} + \frac{\rho}{M^2 - 1} \cdot \frac{\partial A}{\partial \rho}$

$$\beta(\rho, T) \qquad \text{BETA}(2, 3) \qquad = \quad \rho \cdot \left[\frac{1}{(M^2 - 1)^2} \cdot \left(\frac{1}{a} \frac{da}{dx} - A \right) \cdot \frac{\partial M^2}{\partial T} \right]$$

$$+ \frac{1}{M^2 - 1} \cdot \frac{\partial A}{\partial T} \right]$$

$$\beta(\rho, C_i) \qquad \text{BETA}(2, i+3) = \rho \cdot \left[\frac{1}{(M^2 - 1)^2} \cdot \left(\frac{1 \text{ da}}{\text{a dx}} - A \right) \frac{\partial M^2}{\partial C_i} + \frac{1}{M^2 - 1} \cdot \frac{\partial A}{\partial C_i} \right] i = 1, \dots, \text{NSP}$$

The Gas Temperature derivatives are computed:

$$\frac{dT}{dx} \qquad FNX(3) \qquad = -T \cdot \left[(\gamma - 1) \cdot \frac{M^2}{M^2 - 1} \cdot \left(\frac{1}{a} \frac{da}{dx} - A \right) + B \right]$$

$$\frac{\partial [FNX(3)]}{\partial x} \quad AL(3) \qquad = -T \cdot \frac{M^2}{M^2 - 1} \cdot \frac{\gamma - 1}{a} \cdot \left[\frac{d^2a}{dx^2} - \frac{1}{a} \cdot \left(\frac{da}{dx} \right)^2 \right]$$

$$\beta(T, V) \qquad \text{BETA}(3, 1) \qquad = \quad T \cdot \left[\frac{\gamma - 1}{(M^2 - 1)^2} \cdot \left(\frac{1}{a} \frac{da}{dx} - A \right) \cdot \frac{\partial M^2}{\partial V} \right]$$

$$+ \gamma - 1 \cdot \frac{M^2}{M^2 - 1} \cdot \frac{\partial A}{\partial V} - \frac{\partial B}{\partial V} \right]$$

$$\beta(T,\rho)$$
 BETA(3, 2) = $T \cdot \left[\gamma - 1 \cdot \frac{M^2}{M^2 - 1} \cdot \frac{\partial A}{\partial \rho} - \frac{\partial B}{\partial \rho} \right]$

$$\beta(T, T) \qquad \text{BETA}(3, 3) \qquad = \frac{1}{T} \cdot \frac{dT}{dx} + T \cdot \left[\frac{\gamma - 1}{(M^2 - 1)^2} \left(\frac{1}{a} \frac{da}{dx} - A \right) \frac{\partial M^2}{\partial T} + \gamma - 1 \cdot \frac{M^2}{M^2 - 1} \cdot \frac{\partial A}{\partial T} - \frac{\partial B}{\partial T} \right]$$

$$- \frac{M^2}{(M^2 - 1)} \cdot \left(\frac{1}{a} \frac{da}{dx} - A \right) \frac{\partial \gamma}{\partial T} \right]$$

$$\beta(T, C_i) \qquad BETA(3, i+3) = T \cdot \left[\frac{Y-1}{(M^2-1)^2} \cdot \left(\frac{1}{a} \frac{da}{dx} - A \right) \cdot \frac{\partial M^2}{\partial C_i} \right]$$

$$+ Y - 1 \cdot \frac{M^2}{M^2-1} \cdot \frac{\partial A}{\partial C_i} - \frac{\partial B}{\partial C_i} - \frac{M^2}{M^2-1}$$

$$\cdot \left(\frac{1}{a} \frac{da}{dx} - A \right) \frac{\partial Y}{\partial C_i} \right] \qquad i = 1, \dots, NSP$$

(4) Subroutine ADDXXX

This subroutine calculates the addition component of the total derivatives a_i and the partial derivatives a_i and β_{ij} and calculates the total and partial derivatives.

The addition components of the total derivatives are presented below:

$$\frac{dV}{dx}\Big|_{add} = \frac{\dot{M} - \dot{m}V}{1 + \overline{m}}$$

$$\frac{dC_i}{dx}\Big|_{add} = \frac{\dot{S}_i - \dot{m}C_i}{1 + \dot{m}}$$

$$\frac{dT}{dx}\Big|_{add} = \frac{1}{C_p} \left[\frac{E - \dot{m} H_T}{1 + \overline{m}} - \frac{V(\dot{M} - \dot{m}V)}{1 + \overline{m}} - \sum_{i=1}^{nsp} h_i \left(\frac{\dot{S}_i - \dot{m} C_i}{1 + \overline{m}} \right) \right]$$

$$\frac{d\rho}{dx}\Big|_{add} = \frac{-\rho}{T}\frac{dT}{dx}\Big|_{add} - \sum_{i=1}^{nsp} \frac{R_i}{R} \rho \frac{dC_i}{dx}\Big|_{add}$$

On option the adiabatic and addition components of the total derivatives are output from this subroutine.

(5) Subroutine IAUX1 (HL, H, QK, RK, JX)

This subroutine performs implicit integration according to the method discussed in Subsection C.3. The increments for the chemical species concentrations and the fluid dynamic variables at the forward point are calculated by solving the appropriate set of nonhomogeneous algebraic equations.

The calling sequence parameters are:

HL-last integration step size

H -current integration step size

QK-last increments for variables

RK-computed increments for variables

JX- l initial 3 steps

2 general step

3 special step

4 restart step

The total derivatives, $f_{i,n}$, and partial derivatives, $\beta_{i,j,n}$ at the back point are calculated in subroutines DERIV and FLU.

The special step calculation is used only in halving the step size if required.

After each integration step, subroutine IAUX obtains the derivatives at the then current axial position.

For implicit integration the equations used are:

Initial Step and Restart

$$k_{i, 1} = \left[f_{i, o} + \alpha_{i, o} h + \sum_{j=1}^{N} \beta_{i, j, o} k_{j, 1} \right] \cdot h$$

General Step

$$k_{i, n+1} = \frac{1}{3} \left[k_{i, n} + 2 \cdot \left(f_{i, n} + \alpha_{i, n} h + \sum_{j=1}^{N} \beta_{i, j, n} k_{j, n+1} \right) \cdot h \right]$$

Special Step

$$k_{i, n+1} = \frac{h_{n+1}^{2}}{\left(2h_{n+1} + h_{n}\right) \cdot h_{n}} \left[k_{i, n} + \left[f_{i, n} + \alpha_{i, n} h_{n+1}\right] + \sum_{j=1}^{N} \beta_{i, j, n} k_{j, n+1}\right] \cdot \frac{h_{n}}{h_{n+1}} \left(h_{n+1} + h_{n}\right)$$

(6) Subroutine IAUX (HL, H, QK, RK, JX)

This subroutine performs the iteration for the area defined, mass energy, momentum addition calculation. If the problem is pressure defined or an adiabatic-area-defined calculation, this subroutine merely calls Subroutine IAUX1 and then updates the derivatives at the forward point by calling Subroutine DERIV.

For the area defined, mass, energy, momentum addition calculation, the iteration proceeds as described below.

Prediction:

$$\frac{dP}{dx} = R \cdot T \cdot \frac{d\rho}{dx} + R \cdot \rho \cdot \frac{dT}{dx}$$

where

$$\frac{d\rho}{dx} = -\frac{1}{a}\frac{da}{dx} \cdot \rho \cdot \frac{M^2}{(M^2-1)}$$

$$\frac{dT}{dx} = -\frac{1}{a}\frac{da}{dx} \cdot T \cdot (\gamma-1) \cdot \frac{M^2}{(M^2-1)}$$

if

$$0.99 < M^2 < 1.01$$

then

$$\frac{dP}{dx} = 0.005 \cdot \frac{P}{H}$$

if

4.7

$$\frac{da}{dx} = 0.0,$$

then dP/dx from the previous step is used as the first estimate.

Iteration: Subroutine ITER is called successively to use the secant method to provide new estimates for dP/dx such that

$$f(A_{calc} - A_{input}) < \epsilon$$

Convergence: Convergence is obtained when

$$\left| \begin{array}{c} \frac{A_{calc} - A_{input}}{A_{input}} \right| < \epsilon$$

and the pressure at the forward point is computed from

$$P_{i+1} = P_i + \frac{dP}{dx} \Big|_i \cdot H$$

(7) ITER (F1, X1, XNEW, $N\phi\phi$)

The purpose of this subroutine is to find the root or zero of the algebraic equation

$$f(X) = 0$$

using the method of secant or false position. In particular this subroutine is designed to take advantage of the fact that the secant method will always find the root of the above equation if the root has been spanned.

Calling Sequence:

F1 is the value of the dependent variable, f, corresponding to the value of X1. (Input)

X1 is the value of the independent variable, X, which corresponds to F1. (Input)

XNEW is the predicted or new value of the independent variable. (Output)

NQQ is a flag such that

 $N\phi\phi = -1$ the first time ITER is called. (Input)

 $N\phi\phi$ = +1 upon subsequent calls. (Output)

Restrictions:

The user is expected to check for convergence as there are no internal checks made in ITER. A literal must not be input to this subroutine.

Method:

Subroutine ITER utilizes the secant method predictor formula

$$X_{i+1} = X_i - f_i \cdot (X_i - X_{i-1}) / (f_i - f_{i-1})$$

where the subscript i refers to the current value of X and f except for the first iteration in which the value of X is perturbed only slightly. When the root has been spanned, the subroutine saves 2 back values of f and X in order that the root may always be straddled and thus found. The linkage to the subroutine is set up so that if bounds on the root are known, then the value of XNEW may be disregarded and bounded values may be used for the first two guesses. This type of linkage necessitates that the value of X1 must be set equal to XNEW or the bounded value of X. In order to accelerate convergence, if the error within the bounded domain of the dependent variable exceeds a ratio of 10, then the new value of X is set equal to one half of the range.

(8) Subroutine INT

Provides control for the implicit integration procedure, determines the proper set of nonhomogeneous equations to solve, and, after

each integration step, computes the next integration step size according to the following relations:

$$h_{n+2} = 2h_{n+1}, \qquad \left| \frac{k_{i, n+1} - 2k_{i, n} + k_{i, n-1}}{3k_{i, n+1} - k_{i, n}} \right| < \frac{\delta}{10}$$

$$h_{n+2} = \frac{1}{2}h_{n+1}, \qquad \left| \frac{k_{i, n+1} - 2k_{i, n} + k_{i, n-1}}{3k_{i, n+1} - k_{i, n}} \right| > \delta$$

$$MAX$$

$$h_{n+2} = h_{n+1}, \qquad \frac{\delta}{10} \le \left| \frac{k_{i, n+1} - 2k_{i, n} + k_{i, n-1}}{3k_{i, n+1} - k_{i, n}} \right| \le \delta$$

$$MAX$$

On option, (JF=1) only the fluid dynamic variables are used in determining the next integration step size.

If the step size is halved for the fourth step, the integration is restarting using one-half the original step size.

The correspondence between equation number and physical property is:

Equation Number	Property
1	Velocity of gas
2	Density of gas
3	Temperature of gas
4 → NSP+3	Gaseous species mass fraction (1 → NSP) corresponds to (4 → NSP + 3)

(9) Subroutine LESK(Y)

This subroutine is a single precision linear equation solver which is used to perform the matrix inversions required by subroutine IAUX. Gaussian elimination is used with row interchange taking place to position maximum pivot elements after the rows are initially scaled.

(10) Subroutine MAIN

Provides overall logic control for the kinetic calculations, controls the shock calculation for both generalized oblique shock and the normal shock for the stagnation streamline, and prints the summary for the maximum and minimum reaction net production rates.

The normal shock calculation equations are presented below:

$$P_2 = \rho_1 V_1^2 + P_1 - (\rho_1 V_1) V_2$$

$$h_2 = \frac{{v_1}^2}{2} - \frac{{v_2}^2}{2} + h_1$$

$$T_2 = f(h_2)$$

$$\rho_2 = \frac{P_2}{R \cdot T_2}$$

(11) Subroutine OUTPUT

This subroutine provides conversion from internal computational units to output engineering units. The following output parameters are computed by this subroutine:

The pressure (in PSIA) is computed from:

$$P_{(PSIA)} = P/4633.056$$

The gaseous species mole fractions are computed from:

$$C_{i, m} = \frac{R_i}{R} \cdot C_i$$

The gas molecular weight is computed from:

$$MW = 49721.011/R$$

The percentage mass fraction change is computed from:

$$\triangle$$
(Mass Fraction) = 100.0 $\cdot \left(1.0 - \sum_{i=1}^{n} C_{i}\right)$

The gas heat capacity is computed from:

$$Cp_g (BTU/LB-^{\circ}R) = 3.9969 \cdot 10^{-5} \cdot Cp_g$$

The gas static enthalpy is computed from:

$$H_g (BTU/LB) = 3.9969.10^{-5} \sum_{i=1}^{n} C_i$$

$$\cdot (h_i - 905.770 \cdot R_i \cdot \Delta H_{F_i}^{\circ})$$

The percentage enthalpy change is computed from:

$$\Delta H_{T} = \frac{100 \cdot \left(H_{c} - \sum_{i=1}^{NSP} C_{i} \cdot h_{i} - V^{2} / 2 \right)}{HREF}$$

where

HREF =
$$\sum_{i=1}^{NSP} C_i \cdot \left(h_i - 905.770 \cdot R_i \cdot \Delta H_{F_i} \right)$$

evaluated at the initial conditions in Subroutine CONVRT.

The real gas constant and frozen gamma [Subsection C. 5f(2) (EF)] the enthalpy in internal units, the kinetic coupling terms A and B [Subsection C. 5f(3) (FLU)], the maximum relative error [Subsection C. 5t(8) (INT)], and the species concentrations in molecules per cc are also output.

(12) Subroutine OUTXXX

This subroutine is identical except in name to subroutine UTIL. It is used during the iteration for θ , the shock angle, during a generalized oblique shock calculation.

(13) Subroutine PLTSUB

This subroutine saves the current values of the variables requested for plotting and generates the proper labels and formatting for processing by subroutine DSCPLT. Plot information is saved on logical unit IPTAPE.

(14) Subroutine PRNTCK

For the option to print starting at step ND1, printing every ND3rd step up to step ND2, this subroutine checks whether or not the current step should be printed. If it is to be printed this subroutine calls OUTPUT.

(15) Subroutine UTIL (F1, X1, XNEW, NOO)

The purpose of this subroutine is to find the root or zero of the algebraic equation

$$f(X) = 0$$

using the method of secant or false position. In particular this subroutine is designed to take advantage of the fact that the secant method will always find the root of the above equation if the root has been spanned.

Calling Sequence:

- F1 is the value of the independent variable, f, corresponding to the value of X1. (Input)
- X1 is the value of the independent variable, X, which corresponds to F1. (Input)
- XNEW is the predicted or new value of the independent variable. (Output)

NOO is a flag such that

NOO = -1 the first time ITER is called. (Input)

NOO = +1 upon subsequence calls. (Output)

Restrictions:

The user is expected to check for convergence as there are no internal checks made in UTIL. A literal must not be input to this subroutine.

Method:

Subroutine UTIL utilizes the secant method predictor formula

$$X_{i+1} = X_{i} - f_{i} \cdot X_{i} - X_{i-1} / f_{i} - f_{i-1}$$

where the subscript i refers to the current value of X and f except for the first iteration in which the value of X is perturbed only slightly. When the root has been spanned the subroutine saves 2 back values of f and X in order that the root may always be straddled and thus found. The linkage to the subroutine is set up so that if bounds on the root are known, then the value of XNEW may be disregarded and bounded values may be used for the first two guesses. This type of linkage necessitates that the value of X1 must be set equal to XNEW or the bounded value of X. In order to accelerate convergence, if the error within the bounded domain of the dependent variable exceeds a ratio of 10, then the new value of X is set equal to one half of the range.

(16) Subroutine SHOCK

This subroutine calculates the downstream oblique shock conditions for an arbitrary pressure rise. The chemistry is assumed frozen across the shock. The method used is an iteration on the shock angle, θ , using the following:

1.
$$\theta^{(0)} = \arcsin \left[\sqrt{\frac{\frac{P_2}{P_1} \left(\gamma_1 + 1 \right) + \left(\gamma_1 - 1 \right)}{2 \gamma_1 M_1^2}} \right]$$

2.
$$u_1^{(i+1)} = V_1 \sin \theta^{(i)}$$

. 3.
$$v_2^{(i+1)} = V_1 \cos \theta^{(i)}$$

$$u_2^{(i+1)} = \frac{P_1 - P_2 + \rho_1 \left[u_1^{(i+1)}\right]^2}{\rho_1 u_1^{(i+1)}}$$

$$H_2^{(i+1)} = H_1 + \frac{1}{2} \left[u_1^{(i+1)} \right]^2 - \frac{1}{2} \left[u_2^{(i+1)} \right]^2$$

4. $T_2^{(i+1)} = f(H_2^{(i+1)})$, obtained by iteration using subroutine UTIL.

5.
$$\rho_{2a}^{(i+1)} = \frac{P_2}{R T_2^{(i+1)}}$$

$$\rho_{2b}^{(i+1)} = \frac{\rho_1 u_1}{u_2^{(i+1)}}$$

6. if
$$\left| \frac{\rho_{2a}^{(i+1)} - \rho_{2b}^{(i+1)}}{\rho_{2a}^{(i+1)}} \right| < \epsilon$$
 go to 9.

- 7. Call OUTXXX to obtain new value for $\theta^{(i)}$ using secant method
- 8. Go to 2.
- 9. Compute and output downstream conditions:

$$Y_2 = \frac{C_P(T_2)}{C_P(T_2) - R}$$

$$V_2 = \sqrt{u_2^2 + v_2^2}$$

$$M_2 = \frac{V_2}{\sqrt{Y_2 R T_2}}$$

If the upstream Mach number is less than one, no shock calculation is possible and the calculation is terminated at this point.

If the specifed conditions are inconsistent, i.e., if

$$\sqrt{\frac{\frac{P_2}{P_1}}{2 + 1} + (\gamma_1 - 1) + (\gamma_1 - 1)} > 1.0$$

a normal shock calculation is attempted using $\theta = \pi/2$ and no iteration. An error message is provided for both of the above error conditions.

(17) Subroutine SCRX

This subroutine screens the reaction set at each axial station for those reactions which are necessary to assure a relative accuracy for a specified chemical species. Summaries of an ordered set of those reactions which produce and those reactions which destroy the specified species, along with total production and destruction rates are provided.

The subroutine screens those reactions which produce or destroy the specified species and retains those which produce or destroy the species at a rate greater than the input criterion (a percentage of the total production or destruction per unit normalized length). A logical vector is generated with the entry TRUE if the reaction is required or FALSE if the reaction is not required.

(18) Subroutine TREE (L, LIND, N)

The purpose of this subroutine is to reorder an input vector L containing N components so that

$$L(1) \leq L(2) \leq \ldots \leq L(N).$$

L may be either real or integer.

The vector LIND must be input as a vector of integers 1, 2, . . N. The vector LIND will be output as a vector containing the integer numbers which were the original position numbers of the L components.

The method used by subroutine TREE is described by ALGORITHM 245, TREE SORT 3 by Robert W. Floyd, "Communications of the ACM." December, 1964.

(19) Subroutine CONAD

This subroutine calculates the mass, momentum, energy, and species addition functions to the gas phase streamtube resulting from the condensation of a single gaseous species.

Subroutine VAPOR is called at each integration step to determine location of current thermodynamic state relative to liquid-vapor or solid-vapor coexistence line on the pressure, temperature (P, T) surface.

Once condensing vapor becomes saturated, the critical droplet radius and nucleation rate are computed. The addition function components are not calculated until the nucleation rate surpasses a threshold value EJMIN (input under \$PROPEL). The addition functions are computed from the following:

$$\dot{M} = \dot{m}V + g \frac{(1+m)}{\rho V} dp/dx$$

$$\dot{H} = (1-2g) \dot{m}h' - (1+\overline{m}) \dot{g} \frac{dh'}{dT} \frac{dT}{dx}$$

 $\dot{m} = dg/dx$

$$\dot{S}_{i} = \dot{m}$$
, for $i = condensing species$

= o , for remaining species

The rate of change of the condensed mass fraction is obtained from

$$\frac{\mathrm{d}g}{\mathrm{d}x} = \frac{4\pi\rho_{\mathrm{L}}}{\rho\mathrm{V}} \left[\frac{1}{3} r^{**}(\mathbf{x}) J(\mathbf{x}) \frac{A(\mathbf{x})}{A_{\mathrm{o}}} + \frac{\mathrm{d}r}{\mathrm{d}x} \int_{\mathbf{x}_{\mathrm{o}}}^{\mathbf{x}} r(\mathbf{x}, \xi)^{2} J(\xi) \frac{A(\xi)}{A_{\mathrm{o}}} \mathrm{d}\xi \right]$$

The liquid droplet properties are obtained from Subroutine DROPS.

(20) Subroutine VAPOR (PV, PVS, DPVSDT, TS, IP, JJ, ISAT)

This routine computes for a specified species JJ the vapor pressure (PV), saturated vapor properties, and the location of the current state with respect to both the triple point and the liquid-vapor or solid-vapor coexistence line. Vapor pressure is computed from

$$P_v = C_i \frac{Ri}{R} P$$

For pressures and temperatures above the triple point, the saturated vapor pressure (PVS) corresponding to temperature T, derivative of PVS with respect to T, and saturated vapor temperature (TS) corresponding to pressure P are obtained from expressions of the form

PVS =
$$C_1 + C_2/T + C_3/T^2$$

TS = $1./[C_4 + C_5 \ln{(PV)} + C_6 \ln^2{(PV)}]$

DPVSDT = -PVS (
$$C_2$$
 - 2.0 C_3 /T) / T^2

At or below the triple point, the following expressions are used

PVS =
$$\exp(C_8 - C_7/T)$$

TS =
$$C_7/[C_8 - \ln{(PV)}]$$

DPVSDT = PVS
$$C_7/T^2$$

where the constants C_1 through C_8 are determined from data and are input under PROPEL. Preliminary values of these constants for water vapor are stored internally and need not be entered.

The additional parameters in the calling statement are

IP = 0 Current state corresponds to triple point

= 1 Above triple point

= 2 Below triple point

ISAT = 0 Vapor is unsaturated

= 1 Vapor is saturated

(21) Subroutine DROPS

This subroutine computes liquid droplet properties and derivatives as a function of temperature. Above the triple point, latent heat and density are computed from

$$L = A_1 - A_2T - A_3T^2$$

$$\rho = A_4 - A_5 T_D + A_6 T_D^2$$

$$\sigma = A_{11} - A_{12}T_{D}$$

At or below the triple point, the following are used

$$L = A_7 - A_8T$$

$$\rho - A_9 - A_{10}T_D$$

$$\sigma = A_{11} - A_{12}T_D$$

First and second derivatives with respect to temperature are obtained by differentiating the above expressions. The coefficients A_1 through A_{12} are input under \$PRQPEL. Preliminary values of these constants for water vapor are stored internally.

c. Plotting Subroutines Descriptions

(1) Subroutine LINK 50

This subroutine reads the quantities saved for plotting from logical unit IPTAPE into the proper buffer areas and calls subroutine DSCPLT for the actual plotting.

(2) Subroutine DSCPLT

This subroutine is a generalized plotting routine utilizing Calcomp software to produce plots. It requires subroutines SCAL and MAXMIN.

(3) Subroutine MAXMIN

This subroutine finds the maximum and minimum entries in an array.

(4) Subroutine SCAL(XMAX, XMIN, XI, DX, XO, XE)

Given the maximum and minimum of a variable and the number of units (inches, cm., etc.) available for plotting, this routine:

- 1. Determines the most efficient scale per plotting unit of the form $1.0 \cdot 10^a$, $2.0 \cdot 10^a$, $5.0 \cdot 10^a$ where a is an integer.
- 2. Adjusts the minimum scale value so that the plot begins at a multiple of the scale value.

Calling Sequence:

XMAX = the maximum value of the variable (input)

XMIN = the minimum value of the variable (input)

XI = the number of units (in., cm, etc.) available for

plotting (input)

DX = the scale selected for the plot grid (output)

XO = the first plot grid value (output)

XE = the last plot grid value (output)

The algorithm used is given below:

$$W = \frac{XMAN - XMIN}{XI}$$

Then select s = 1, 2, or 5 such that

W = DX is a minimum

where

*

$$A - 1 + \log\left(\frac{w}{s}\right)$$

or

$$B = [A] - 1 \text{ if } A = [A]' \text{ or } A < 0$$

$$C = s 10^B$$

$$D = \left| \frac{XMIN}{C} \right| C$$

or

$$D = \left(\left[\frac{XMIN}{C} \right] - 1 \right) C \text{ if } C \cdot D > XMIN$$

$$E = W/C$$

and

$$DX = C$$

$$XO = D$$

$$XE = E$$

also

if W=O then DX=XO=XE=O

C. 6 PROGRAM USER'S MANUAL

This program was developed on the CDC 6500 computer using the FORTRAN IV language. Conversion to another computer system should be straight forward provided sufficient core storage (120,000g words) is available. Program overlay extends three levels deep including the executive level, when used as a subprogram to CONTAM.

The description of the operation mode and input to the computer program is divided into the following six Subsections:

- C. 6. 1 OPERATION MODE-Automatic or manual.
- C. 6. 2 \$THERM \(\textstyle \text{Namelist input which controls the thermodynamic data input.} \)
- C. 6. 3 THERMODYNAMIC DATA-Optional.

- C. 6. 4 TITLE CARD-Also serves for plot labels.
- C. 6. 5 SPECIES CARDS-Species to be considered and their initial concentrations.
- C. 6. 6 REACTION CARDS-Input of reactions and rate data.
- C. 6. 7 \$PROPEL-Namelist input for a specific case.

Card listings for the complete input for several sample cases are given in Section C. 7.

a. Operation Mode

4 3

Two modes of operation (automatic or manual) are available and specified through input variable $KM \not ODE$ in the executive program, CONTAM.

(1) Manual Mode (KM \emptyset DE = 1)

In the manual mode of operation the <u>KINCON</u> subprogram is independent of the other subprograms. The user is free to specify, via input data (cards), the streamtube initial conditions, boundary conditions, and program options. A complete set of data (inputs described in Subsections C. 6b through C. 6g) is required for each streamtube calculation. However, if a specific nozzle/plume streamtube, generated by either the nozzle or plume portion of <u>MULTRAN</u> is desired; subprogram <u>SLINES</u> may be utilized in conjunction with <u>MULTRAN</u> to specify initial conditions and pressure distribution while operating in the manual mode. The variable NSL, which specifies the number of streamtubes processed by subprogram <u>SLINES</u>, is input through either the executive program or <u>SLINES</u> and is tested to determine if input data from <u>SLINES</u> via TAPE 8 is to be used in the manual mode of operation.

If NSL = 0, TAPE 8 will not be read and all data must be input.

If NSL \$\neq\$ 0, the initial values (Z, PI, T, V, and EXIT), as well as the streamtube pressure distribution (NTB, ZTB (I), and PTB (I)) are read from TAPE 8. If successive cases are run, TAPE 8 will be read successively until either the final case is completed or the number of streamtubes on TAPE 8 is exhausted. It should be noted that any or all data read from Tape 8 may be overridden by card input.

(2) Sequential Mode (KMØDE = 0)

The automatic mode of operation is designed to calculate both chemical-kinetic and single-species condensation effects (a two-pass calculation performed in that order) for a number of streamtubes calculated by MULTRAN and SLINE subprograms. This mode requires only one set of input data and is utilized in conjunction with subprogram SLINES (or a TAPE 8 previously generated by SLINES).

In the automatic mode, KINCON will accept input data describing initial gas-phase composition, chemical reaction and rate coefficients, and pertinent integration control parameters (described in Subsections C. 6. b through C. 6. g). Initial streamtube conditions and pressure distribution are read from TAPE 8, and kinetics calculation is then performed along the pressure-defined streamline. During the kinetics calculation pass the vapor pressure of the condensible species is compared with the saturated vapor surface to determine the vapor state. Once the saturated vapor state is reached, the streamtube conditions at the saturation point (location, temperature, pressure, velocity, and species composition) are written onto TAPE 1. The streamtube area ratio, referenced to the saturation point, is then computed and written on TAPE 1 during the subsequent integration.

TAPE 1 thus contains, at the completion of the kinetics pass, the initial conditions and area ratio distribution for the streamtube beginning at the saturation point.

If the saturation point is countered on the kinetics pass, a second pass is made to compute condensation. This pass, utilizing TAPE I generated during the kinetics pass, invokes frozen-chemistry and single-species condensation options to compute condensation effects along the area defined streamtube. Following the completion of the condensation calculation, initial conditions for the next streamtube are read from TAPE 8 and the calculation procedure is repeated until all streamtubes have been completed. If the saturation point is not reached on a given kinetics pass, the condensation calculation pass is not performed and the program proceeds to the next streamtube.

Inputs in the automatic mode consist of all inputs described in Subsections C.6.b through C.6.f and a limited number of inputs in \$PROPEL. Inputs required in \$PROPEL are the following:

RSTAR

Axial distance normalizing factor (nozzle

throat radius)

PRINT VARIABLES

(ND1, ND2, ND3)

INTEGRATION VARIABLES

(HI, HMIN, HMAX, DEL, JF)

Inputs that are available but not required include MISCELLANE-OUS VARIABLES (except IFLAST), INTERMEDIATE OUTPUT VARIABLES, and certain CONDENSATION OPTION VARIABLES.

b. \$THERMØ

Permits the generation of a master thermodynamic file or the use of a tape file previously generated master thermodynamic file.

٠	P	E	d	
	۲	ľ		
h	31	P	od	
	3		뼥	

Variable		Value	Description
NUCHEM	=	1	A master thermodynamic file will be generated for this case on tape unit 4. Species and thermodynamic data will be read from unit INTAPE (nominal=5, the input file) and a new thermodynamic file will be generated on unit 4. The new thermodynamic file will be end-filed and rewound after generation.
	Ξ	0	A previously generated master thermodynamic file will be used for this case. The master thermodynamic file must be file TAPE 4. No other input variables are required for this option.
MAXSP	=	Input	If a master thermodynamic file is to be generated this variable specifies the number of species for the master file.
LIST	=	1	A list of species named for those species master file will be output.
	=	0	This output will be deleted.
LISTX	=	1	Thermodynamic functions (CP, H, F) will be output for each species (one page per species, 52 lines per page). Species names, molecular weights, and heats of formation will also be printed.
	=	-1	Only a table of species names, molecular weights, and heats of formation will be printed.
	=	0	The above output will be deleted.
INTAPE	=	Input	Tape file from which thermodynamic data is to be read (nominal = 5).
ADM			

\$END

c. Thermodynamic Data

For a NUCHEM = 1 option in \$THERM ϕ the program will read MAXSP Master Species cards containing: species symbolic identifier, molecular weight, and $\Delta H^{\circ}F_{298}$, and then read MAXSP sets of thermodynamic data (CP, H, F) checking names and card sequences. The Master Species cards must be sequenced sequentially in columns 41-50 (I10 format) and must correspond directly to the order in which the thermodynamic data is to be read. Thermodynamic functions consist of 10 cards per function, 5 values per card corresponding to temperature values of $100 \rightarrow 5,000^{\circ}K$ at $100^{\circ}K$

intervals. Table C-I lists the species for which thermodynamic data is currently available. For species which do not appear in Table C-I, this input may be obtained directly from the JANAF tables. Table C-II is a sample listing of the thermodynamic function input for the species N₂.

Master Species Cards

Column	Information
1 - 10	Not used.
11 - 16	Species Symbolic Identifier, 6 alphanumeric characters (left justified).
17 - 20	Not used.
21 - 30	Species molecular weight (F) format.
31 - 40	Species AH°F209 (F) format, K cals/mole.
41 - 50	Species ΔH°_{F298} (F) format, K cals/mole. Right justified sequence number used for sequence checking on input (IIO format).
51 - 80	Not used.

It should be noted that a species is identified by the name assigned by the user. In general this name is the chemical symbol, e.g., O, O₂, H, H₂. However, it may be useful to define a dummy species with all the properties of another species but which may be treated in a special manner, e.g., the percentage of the total amount of a species which is designated as an inert (possibly to simulate incomplete mixing or combustion). This may be done by defining species O and OX where OX is identical to O except in name, but does not appear in any reaction.

Master Thermodynamic Function Cards

Column	Information
. 2 - 10	Not used.
2 - 10	Function value at (100 + 500 (n-1)) °K, n = card number.
11	Not used.
12 - 20	Function value at (200 + 500 (n-1)) °K.
21	Not used.
22 - 30	Function value at $(300 + 500 (n-1))$ °K.
31	Not used.
32 - 40	Function value at (400 + 500 (n-1)) °K.
41	Not used.
42 - 50	Function value at (500 + 500 (n-1)) °K.
51 - 60	Not used.
61 - 66	Species symbolic identifier, left justified.
67 - 68	Not used.
69 - 70	Function Definition CP, H, or F, left justified.
73 - 76	The word CARD.
77 - 78	Card number 1-10 right justified.
79 - 80	Not used.

Species symbolic identifier, function definition, and card numbers are checked for consistency on input.

Table C-L. SPECIES RESIDING ON MASTER THERMODYNAMIC TAPE

1 1150	19 SF4	37 WCL3	55 C2F2	73 0052	90 ALF
2 H,	Per SF6	3% PF	56 03	74 104	91 ALF?
3 1	71 0	30 PF2	57 HO2	75 MH	92 ALFT
4 72	12 6 2	40 IF3	58 405-	74 1H2	93 ALUCL
r . ,;	23 C'	41 '-OCL	59 NA	77 H.	94 ALUF
ķ u	24 00	47 HOF	60 NA+	7# 8H	95 ALCLE
7 6 191	28 Qu	43 PCLF	61 HAQ	73 BHS	96 ALCL2F
e Fo	76 G.	44 PCL2F	62 CF2	ви на	S7 ALCLES
9 44	27 6 3	45 POLF?	e3 CF3	81 4246	98 CH=
10.1	4 C+4	45 1	64 LF4	95 (555	99 CH20
41.	75 C282	47 CL	65 C2F4	F7 "2"3	100 0.400
17		44 CL2	66 NO2	ee al	101 WAR
11 .	Si Hon:	49 t	67 120	BE ALC	192 HZ2
14 12-	32 BUT	50 HCL	68 HC0	er also	103 HZ
15 10	53 HA	51 OLF	A9 HCO+	87 ALCL	104 GHX
16 1 -	sa any	52 CNCL	70 C2H	Bu stots	105 PHOTON
17 6-	is bal	53 CNF	71 430+	SO ALCES	106 67
10 2	16 8042	54 (F	72 H202		

Table C-II. SAMPLE LISTING FOR THERMODYNAMIC FUNCTION CARDS FOR N_2

15.00							
6,9560,	5,9570,	6,7610,	6,9900,	7,0690,	N2	60	
7,1960.	7,3500,	7,5120,	7,6700,	7,8150,	N2	CP	CAPD 1
7,9450,	8,0610,	8,1620,	8,2520,	8,3300,	_	CP	CARD 2
8,3980	8,4580.	8,5120.	8,5590,		N2	CP	CAPD 3
8,6380,	8,6720.	8,7030.		A,6010.	NS	CP	CARD 4
8,7790	5.8000.		8,7310,	8,7560,	N2	CP	CARD 5
		8,5200,	8,8380,	8,8550,	N2	CP	CARD 6
8.8710.	8,8850,	8,9000.	8,9140,	8,9270,	N2	CP	CARD 7
8,9390	9,9500,	8,9620,	8.9720,	8,9830,	N2	CP	CARD 8
8,9930	9,0020,	9,0120,	9,0210,	9,0300,	N2	CP	CARD 9
9,0390	9,0480,	9,0570,	9,0660,	9,0740,	N2	CP	CARD10
-1.3790	6830,	,0130,	.7100.	1,4130,	_		
2,1250,	2,8530,	3,5960,	4.3550,	5,1290,	N2	H	CARD 1
5,9170,	6.7180.	7,5290,	8,3500.		N2	• н	CARD 2
10.0150	10.8530.	11,7070,	12.5600,	9,1790,	N2	н	CARD 3
14,2800.	15,1460.	16.0150.		13,4180.	N2	H	CARD 4
18,6380	19.5170.		16,8860,	17,7610,	N2	H	CARD 5
	• -	20,3980,	21,2800.	22,1650,	N2	H	CARD 6
23.0510	. 23,9390.	24,5290,	25,7190.	26,6110,	N2	Н	CARD 7
27,5050.	25,3990.	29,2950,	30,1910,	31,0890,	N2	н	CARD 8
31,9880.	32,8880,	33,7880.	34.6900,	35,5930,	N2	H	CARD 9
36,4960,	37,4000,	38,3060,	39,2120,	40,1190,	N2	H	CAPD10
51,9570.	45.4070,	45,7700.	46,0430,	46,5610,	NZ	_	
47,1430,	47,7310,	48,3030.	48,8530,	49.3780.	-	F	
49,8790	50.3570.	50,8130,	51,2480,	51,6650,	N2	F	CARD 2
52,0650	52.4480.	52,8160.	53,1710,		NS	E	CARD 3
53,8420,	54.1600.	54,4680.		53,5130,	N2	F	CAPD 4
55,3350,	55.6060.		54.7660,	55,0550,	NS	F	CARD 5
56,6190,		55,8700.	56,1270,	56.3760,	N2	F	CARD 6
	55,8560,	57,0870,	57.3120.	57,5320.	N2	F	CAPD 7
57,7470	57,9570,	38,1620,	58,3620,	58,5590,	N2	F	CARD 8
58.7510.	58,9400.	59,1240,	59,3050,	59,4820,	N2	F	CARD 9
59,6570,	59.8270.	59,9950,	60.1600.	69.3220.	N2	F	CARD10
				- , ,	14.	•	OWERTA

d. Title Card

The title card contains free field information which will be written as a header label for the program output. The first 40 characters will be written as a label on each plot if plotting is requested.

e. Species Cards

This input is prefixed by a single card with SPECIES in columns 1 to 7 and with the words MASS FRACTIONS or MØLE FRACTIONS in columns 9-22. If the identifier for mass or mole fractions is omitted, mass fractions are assumed. Up to 40 species cards may be input. Only those species specified by input species cards will be considered. The order of the input species cards is independent of the order in which the species appear on the master thermodynamic data file. However, the order of the input species cards does define the species order for the specific calculation, and other input referencing individual species must refer to the order of the input species cards. Species Cards are described below:

Column	<u>Function</u>
1 - 10 No	t used.
11 - 16 Syr	mbol (left justified).
17 - 20 No	t used.
	lue of initial species concentration (if zero must input as 0.0) free field F or E format.
	er identification if desired.

Symbols for Species Identification

A chemical species is identified symbolically by six alphanumeric characters as follows:

The species symbol must agree with that given on its master species card (columns 11-16) used in generating the master thermodynamic file. If the species is ionized, the degree of ionization is indicated by a + (or -) sign followed by an integer describing the degree of ionization (if no integer is given the species will be assumed singly ionized). The species symbol may not contain the characters * or =. The special species symbol PH Φ T Φ N is reserved for specifying radiative reactions.

Examples:

Symbol	Interpretation
CL	CI
NA+	NA ⁺ K ⁺⁺
K+2	K ⁺⁺
CL2-2	C/2-
H2O2	H ₂ O ₂

f. Reaction Cards

This input is prefixed by a single card with REACTIONS in columns 1 to 9. Up to 15 dissociation reactions and a total 150 reactions may be input following this card. Only one card per reaction is permitted. Cards specifying dissociation-recombination reactions must precede cards specifying exchange reactions. The content and format of the reaction cards are defined as follows:

(1) Each card is divided into five fields, separated by commas. Each field contains:

Field 1 the reaction.

Field 2 A = followed by the value of A.

Field 3 N =followed by the value of N.

Field 4 B = followed by the value of B, the activation energy

(Kcal/mole).

Field 5 available for comments.

Rules for specifying the reaction are given in C. 6. e(2) below. The values A, N, and B define the <u>reverse</u> reaction rate, k, as

$$k = A \cdot T^{-N} \cdot e^{-(B/RT)}$$

All three reaction rate parameters must be input. The numeric value of each parameter may be specified in either I, F, or E format. If E format is used, the E must appear before the exponent.

There may be no blanks between the characters A and equal sign, the N and equal sign, and the B and equal sign.

Input rate parameters are in units of cc, °K, mole, sec.

(2) The general form of a reaction is:

$$N_1*Symbol_1 + N_2*Symbol_2 + \dots = N_a*Symbol_a + N_b*Symbol_b + \dots$$

where the left-hand side represents reactants and the right-hand side represents products.

Each symbol must be as defined on an input species card (see the description of SPECIES CARDS).

The multipliers, N, must be integers and represent stoichiometric coefficients. If no stoichiometric coefficient is given, the value 1 is assumed.

It is required that

$$N_1 + N_2 + \dots \leq 10$$

and

$$N_a + N_b + \dots \leq 10$$

Examples:

Reaction

Interpretation

NA++CL- = NACL
B+2+M-2 = BM
$BE+2+2*\phi H = BE\phi H\phi H$

 $Na^{+} + C\overline{l} = NaCl$ $B^{++} + M^{--} = BM$ $B^{++} + 2\emptyset H^{-} = Be (\emptyset H)_{2}$

(3) The dissociation—recombination reactions specifying third-body terms must precede other types of reactions, and must be followed by the directive:

Column 1

END TBR REAX

All reactions prior to the above directive will have a third-body term added to each side of the reaction. For example:

$$H_2 = H + H_1 \dots$$

END TBR REAX

is the same as

$$H_2 + M = H + H + M$$
, . . .

where M is a generalized third body. Specific third-body effects may be included by inputting specific third-body reaction rate ratios XMM (J, I).

- (4) Radiative reactions may be considered using the special species $PH \phi T \phi N$. The $PH \phi T \phi N$ may only appear on the left-hand side of the equal sign for a reaction.
- (5) The reaction set is terminated by a card containing LAST CARD in columns 1 to 9.

g. \$PRØPEL

Case Vari	ables	Units
*Z	Initial normalized axial position	None
*PI	Initial pressure	PSIA
*T	Initial temperature	°R
*V	Initial velocity	ft/sec
RSTAR	Axial distance normalizing factor (normally throat radius for nozzle calculations)	Inches

*EXIT Normalized axial distance for run termination None NØCHEM = 1 if a frozen chemistry case is desired None

Plot Variables (plotting not available in automatic mode of operation)

IFP = 1 Plotting requested ITP ~ 1 Plot temperature IRØP Plot density Species numbers for desired species concen-ICHEMP(1) =tration plots, up to 30 species Plot functions vs normalized distance IXP = 1 MLP = 1 Plot species MØLE fractions

MSP = 1 Plot species MASS fractions

Print Variables

HI

ND1	Print every ND3 rd step beginning with the ND1 st step
ND2	until the ND2 nd step. (The initial conditions and the
ND3	EXIT point are always printed.)

Integration Variables

HMIN	Minimum normalized step size
HMAX	Maximum normalized step size
DEL	Relative error criterion
JF	 = 0 all variables considered for step size control = 1 only fluid dynamic variables considered for step
	size control.

Initial normalized step size

The variable which controls the step size; i.e., has the maximum relative error; is printed in the normal output under Integration Parameters, labeled Governing Equation. The number: variable correspondence is as follows: 1=T; 2=RO; 3=V, 4=Species 1; 5=Species 2; ... NSP + 3 = Species NSP.

Table Input Variables

*NTB Number of input table entries for pressure table (101).

JPFLAG

Determines the type of differentiation used to obtain derivatives for all input tables. Reference should be made to Mass, Momentum, Energy and Species Addition Functions for other input tables controlled by JPFLAG.

^{*}Values are read from TAPE 8 and need not be input when operated in automatic mode (KM ϕ DE = 0) or in a manual mode (KM ϕ DE = 1) and using streamtube data from TAPE 8 (NSL \neq 0). If input, they will override TAPE 8 values for first streamtube only.

= 1 derivatives of input table obtained by simple difference formulae with second derivatives defined as 0. Normally used in automatic mode.

= 2 derivatives of input tables obtained by SPLINE fit (NTB ≤ 20).

= 3 derivatives input along with tables.

= 4 derivatives of input tables obtained by parabolic differentiation (NTB ≤ 20).

*PSCALE = Multiplicative constant for the input pressure table.

*ZTB (1) = Normalized axial positions for input tabular values for pressure table (always input).

*PTB (1) = Pressure table (PSIA) (always input).

*DPTB (1) = Pressure table derivative (for JPFLAG = 3 option).

Miscellaneous Variables

XMM (J,I) Reaction rate ratio effect on reaction J of species I.

TU (1) Temperature above which approximate extension of JANAF tables will occur for each species (nominal = 9,000°R).

IFLAST For overlay reasons must be set = 1 on the last case of a run.

TSTOP Time at which a run will arbitrarily be terminated (CP time). Ignore in automatic mode.

Intermediate Output

IDQDX = 1 Print total derivatives.

IDXJDX = 1 Print individual reaction net production rates.

IEQØUT = 1 Print equilibrium constant and its temperature dependence in internal units.

IRATE = 1 Print reaction forward and reverse rates in internal units.

^{*}Values are read from TAPE 8 and need not be input when operated in automatic mode (KM \emptyset DE = 0) or in a manual mode (KM \emptyset DE = 1) and using streamtube data from TAPE 8 (NSL \neq 0). If input, they will override TAPE 8 values for first streamtube only.

IMASDX	= 1	For an IADDF = 1 option print chemical and addition function components of total derivatives.
IØP XF	= 1	Print influence coefficient vectors.
IØPVAR (1)		= Set consecutive entries equal to K for variables for which influence coefficient vectors are to be output, where

<u>K</u>	Variable	
1 2 3	V R Ø T	
4	Species number	1
•	•	
•	•	
•	•	
NSP+3	Species number	NSP

For example: IQPVAR(1) = 4, 3, will output influence coefficient vectors for species number 1 and for temperature.

Mass, Momentum, Energy, and Species Addition Functions

<u>Variable</u>		
IADDF	= 1	Addition functions will be input.
IXTB	the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the s	Number of entries in addition function tables ≤ 40.
IADD ØP	= 0	Addition functions input via tables.
	= 1	Addition functions defined via multiplicative factors. See EFACT (I), XMFACT (I), SPFACT (I, J) below. Note that IXTB factors must be input.
	= 2	A subroutine for addition calculation will be supplied by the user.
XADSCL	=	Multiplicative scale factor for all addition functions. Note that input of XADSCL as 1.0/(p. V. A) will provide automatic normalization (per unit initial streamtube mass flux) for input addition functions.
ADDX (I)	=	Table of normalized axial stations for all input addition functions.

ADDMAS (I) Mass addition rate, per unit normalized length, per unit initial streamtube mass flux (unitless). ADDE (I) Energy addition rate, per unit normalized length, per unit initial streamtube mass flux (BTU/1b). ADDEX (I) Auxiliary energy addition rate, per unit nor-= malized length, per unit initial streamtube mass flux (BTU/lb) which will be added to the energy addition rate ADDE (I). Note that ADDEX (I) is independent of EFACT (I) for IADD OP = 1 option and will be added to ADDE (I) before modification by XADSCL. ADDMØM (I) Momentum flux, per unit normalized length, per unit initial streamtube mass flux (ft/sec). NPØINT (I) Species number (for current case) to relate species addition functions to specific species. See example under ADDSP (I, J). NSPADD Number of entries in NPØINT (I) table. ADDSP (I, J) Species mass addition rate per unit normalized length, per unit initial streamtube mass flux (unitless). I=1, . . . IXTB corresponding to number of entries J=1, ... NSPADDe. g., NSPADD = 2NPOINT = 3.5ADDSP (I, 1) corresponds to Species 3 ADDSP (I, 2) corresponds to Species 5 For IADDOP = 1, ADDE (I) computed as EFACT (I) ADDMAS(O) *EFACT(I). For IADD ϕ P = 1, ADDM ϕ M (I) computed as XMFACT (I) ADDMAS (I) *XMFACT (I). For IADD Φ P = 1, ADDSP (I, J) computed as SPFACT(I, J) =ADDMAS (I) *SPFACT (I, J). Restart flag indicating that a case is being IHTØTF = 1 restarted and directing the initial enthalpy to be HTOTX. HTØTX Initial enthalpy if a case has been restarted (ft^2/sec^2) .

DMASDX (I) = For JPFLAG = 3 option, derivative of the mass addition rate d(ADDMAS)/dx.

DEDXT (I) = For JPFLAG = 3 option, derivative of the total energy addition rate d(ADDE+ADDEX)/dx.

DMØMDX (I) = For JPFLAG = 3 option, derivative of the momentum addition rate d(ADDMØM)/dx.

DSPDX (I, J) = For JPFLAG = 3 option, derivative of the species addition rate d(ADDSP)/dx.

Generalized Oblique Shock Calculation

A generalized oblique shock calculation is specified by input of a pressure table containing a pressure discontinuity, and a pointer designating the shock location.

UNITS

NSHØCK = Pointer designating the last entry in the pressure table prior to the shock. For example: $\frac{P_2}{P_1} = \frac{PTB \text{ (NSHØCK+1)}}{PTB \text{ (NSHØCK)}}$ Entries NSHØCK and NSHØCK+1 in the pressure table must have the same axial position.

SHKBUG = 1.0 provides intermediate output. None

0.0 provides no intermediate output,

SMAXIT = Maximum number of iterations None during a generalized oblique shock

calculation.

SKEPS(1) = Relative convergence criterion for None

temperature iteration during a generalized oblique shock calculation.

SKEPS(2) = Relative convergence criterion for None

overall iteration during a generalized oblique shock calculation.

Normal Shock Stagnation Streamline Calculation

The normal shock stagnation streamline calculation option performs a normal shock calculation from specified upstream and downstream velocities and continues the calculation as a velocity defined streamtube.

			<u>UNITS</u>
NSTAGV	=	1 specifies a normal shock stagnation streamline calculation.	None
VEL1	=	Upstream velocity for the normal shock calculation.	ft/sec
VTB(1)	=	Array defining the velocity as a function of normalized axial distance. VTB(1) is defined as the downstream velocity for the normal shock calculation.	ft/sec
NVTB	=	Number of entries in the velocity profile NVTB ≤101.	None

Reaction Screening Input Variables

1

If a reaction screening calculation is requested, the program performs a two-pass calculation. The first pass utilizes the complete reaction set and determines those reactions which must be retained to satisfy the input criteria for each species screened. The second pass redoes the first calculation with an edited reaction set and provides a summary page comparing both calculations.

			<u>UNITS</u>
ISCRF	=	1, specifies a reaction screening case for ISCSP (I) species.	None
ISCSP(1)	= .	Species number for those species to be screened ≤40.	None
EPSCR(1)	=	Relative retention criterion for each species to be screened. Defined as the maximum change in mass fraction relative to production or destruction of the species per unit normalized length for all reactions involving the corresponding species ≤ 40 .	None
ISCBUG	z	1, provides intermediate output during the reaction screening procedure.	None

Area Defined Option Input

Area Defined Calculation With Mass, Energy or Momentum Addition:

ITAREA = Input maximum number of iterations for area ratio calculation. A nonzero value for ITAREA triggers the area ratio iteration logic. After maximum iterations, the program outputs an error message, accepts the most recent values, and continues.

ABAR(1) = Area ratio table (A/A_0) .

XARTAB(1) = Normalized axial coordinates for the input

area ratio table.

DABARX(1) = Derivative with respect to normalized axial

distance of the input area ratio table (if input

derivative option used).

NPATAB = Number of entries in area ratio table ≤40.

ICALDA = 0 derivative of area ratio table controlled by

JPFLAG.

= 1, 2, 3, 4 replaces JPFLAG control for area

ratio table ONLY.

AREPS = Relative convergence criterion for area ratio

iteration.

Single Species Condensation Option

Condensation calculations require the use of both the mass, momentum, and energy addition option and the area-defined streamtube option. The pertinent parameters for these options are included below:

*IADDF = 1 Addition function option will be utilized.

*IADDØP = 3 Single species condensation routines will

provide addition functions.

*ITAREA = Maximum number of iterations for area ratio

calculation.

*XARTAB(1) = Normalized axial coordinate for input area

ratio table.

*ABAR(1) = Area ratio table (A/A_0) .

^{*}Values for these variables are calculated internally when operated in automatic mode (KMØDE = 0) and need not be input.

*NPATAB = Number of entries in area ratio table ≤ 40.

*DABARX (1) = Derivative with respect to normalized axial distance of input area ratio table (if input

derivative option used).

AREPS = Relative convergence criterion for area

ratio iteration.

*NOCHEM = 1 Frozen chemistry option is utilized.

ICØND = Number signifying the location of condensing

species with respect to the species input

order.

The following inputs describe the vapor/liquid properties. A preliminary set of constants for water is stored internally and will be used if the following inputs are ignored. If improved values for water are available or if a different condensing species is being considered, some or all of the following may be input:

TLIM = Limiting temperature for saturated vapor

test. State of condensible vapor will not be

tested above TLIM (units = *R).

EJMIN = Threshold value for nucleation rate below

which condensation effects will not be

computed (units = $1/ft^3$ -sec).

ETA = Condensation (or sticking) coefficient for

water vapor/liquid.

PCØNST (I) = Array of constants utilized in describing

vapor state (see subroutine VAPØR).

 $PCONST(1) = C_1$ (atm)

 $PCONST(2) = C_2 \quad (atm - {}^{\circ}R)$

 $PCONST(3) = C_3 \qquad (atm - (^{\circ}R)^2)$

 $PCONST(4) = C_A (1/^{\circ}R)$

 $PCONST(5) = C_5 \qquad (1/{}^{\circ}R)$

 $PCØNST (6) = C_6 \qquad (1/{}^{\circ}R)$

 $PCONST(7) = C_7$ (°R)

 $PCØNST(8) = C_8$ (-)

^{*}Values for these variables are calculated internally when operated in automatic mode (KMØDE = 0) and need not be input.

DCØNST (1) =
$$A_1$$
 (BTU/lb)
DCØNST (2) = A_2 (BTU/lb)- ${}^{\circ}$ R)
DCØNST (3) = A_3 (BTU/lb)- ${}^{\circ}$ R²)
DCØNST (4) = A_4 (lb/ft³)
DCØNST (5) = A_5 (lb/ft³) - ${}^{\circ}$ R)
DCØNST (6) = A_6 (lb/ft³ - ${}^{\circ}$ R²)
DCØNST (7) = A_7 (BTU/lb)
DCØNST (8) = A_8 (BTU/lb - ${}^{\circ}$ R)
DCØNST (9) = A_9 (lb/ft³)
DCØNST (10) = A_{10} (lb/ft³ - ${}^{\circ}$ R)
DCØNST (11) = A_{11} (Poundal/ft)
DCØNST (12) = A_{12} (Poundal/ft - ${}^{\circ}$ R)

DCØNST (13) = Condensed phase heat

capacity (BTU/lb - *R)

C. 7 SAMPLE CASES

A sample case illustrating the abilities of the KINCON program as a subprogram to CONTAM has been presented in the program description of CONTAM, Section 7.0. In this mode, the KINCON program analyzes the nonequilibrium chemical kinetics and condensation of plume species for prediction of contamination effects on bodies submerged in bipropellant plumes. Several additional KINCON program options are available when run in the manual mode as an independent program. Sample cases are included in this appendix to illustrate these options.

a. Reaction-Rate Screening Case

Computer input card listing and selected output for the rate screening of pure-air chemistry system are presented in Tables C-III through C-V.

The original system consists of a set of 14 species and 37 reactions as presented in the card-image listing in Table C-III. The complete reaction set is utilized on the first calculation pass. Those reactions which must be retained to satisfy the input criteria for each species screened are determined. A second pass recomputes the case using the edited reaction set and provides a summary page comparing both results.

Output for a rate-screening case differs from a standard kinetics run only by the addition of intermediate reaction-rate printout for each screened species at each output station and a summary page comparing the results of the original and edited reaction sets. Table C-IV contains the reaction rate output for screened species E-(electron) at axial position 20.0. A similar page is output for each remaining screened species. The summary page is included in Table C-V.

b. Oblique Shock Case

Card image listing for an oblique shock calculation in air is included in Table C-VI. A discontinuity in the input pressure table identifies the shock location as shown in pressure table output, Table C-VII. Results of the shock calculation are presented in Table C-VIII. The output for the integration to and from the shock is the same as the standard kinetics output and has been omitted.

c. Normal Shock-Stagnation Streamline Case

Card-image listing for a normal shock stagnation streamline calculation in air is included in Table C-IX. The streamline downstream of the normal shock is defined through an input velocity distribution table. Velocity table is output as shown in Table C-X. The normal shock calculation output is included in Table C-XI. The remaining output for this case is of the same format as the standard kinetics run and has been omitted.

d. Automated Kinetics and Condensation Case

The following sample case was completed to illustrate the operation of the KINCON subprogram in the automatic mode including both kinetics and water vapor condensation passes in a single case. The test case corresponds to a streamline bounding 90% of the total mass flow for a typical MMH/NTO engine. The initial streamtube temperature was purposely reduced from the original combustion chamber value to ensure condensation within the region of interest.

A card-image listing of the case is shown in Table C-XII. The streamline conditions (initial pressure, temperature, and velocity) including the pressure distribution were obtained from the output of the MULTRAN subprogram and are presented in Table C-XIII. Printout of the input data and the pressure distribution table were omitted since that information is available in Tables C-XII and C-XIII. The initial streamline conditions for the kinetics pass are presented in Table C-XIV.



Table C-III. CARD LISTING FOR RATE-SCREENING CASE

PURE AIR SCHEENING		FHOURAN SCHEEN		
PECLES				
• • • • • • • • • • • • • • • • • • • •	1,760000	****		
14	0,2259369			
;)	4,737153743			
· · · · · · · · · · · · · · · · · · ·	1, 001/37-7			
	7,921345 =7			
<u> </u>	2.0654731-16			
	1,96494256			
	7.7.54212			
٠.٠.	2.22374: 07			
''& '	7.536630007			
102-	1.076755-10			
າડ	4,47:535:04			
; ∪ ₹	7,9915190-31			
ACTIONS				
12 = 0 + 1	1.1541-4 8 1.	2. Hm 0.0.		
45 m % + , " " = ,	1,595417) (# 5	1,12, 9= 0,0,		
VC = N + F ,	4,06+14, # 9,	i, d≡ ^.î,		
102 8 100 + C A A	F 1 . 1 E + 15 . 48	He =1.79.		
VC = NO+ + 5- , A	is 5,675+25,	= 2.5, b= 0.9,		
03 = 0 + 12 , /=	1.006+19, 1=	2,, 0m 0,0, 1 m ALL		12,49
02- = 02 + E- , ns	13.51 16. 185.i.	380.0.		14.24
J- # 0 + [- , A=				15,23
		(NOS XIN)		16.77
10 + L-	AR SACEPICA	B G G BB Tara		17,15
THE REAL				
		F =1,5, AE 5,94,	1	4
3 4 03 = 10 4	AB 1 SEPTIME	WE 0.0. DE D.M.	2	<u> </u>
	A 4 3544	L= 9,7; 4m 79,5; .m 0,7; 4m 27,7;	-	
ic a kg a myo t g	1 14 TIS A14	** *** *** *** **** ****	57 51	51 52
C + 02 = U + 11.52		v= 0, 10 c= 1, 15.	23	56
12 4 02 = 1 4 3		0	24	
C + 02 B A + 13	4 3 4 6 5 6 6 6 1 1 .	18 U.U. 18 C.T.	2:	
		1.8 0.0. RE 2.57.	26	57
		P. NE 7.C. CE U.C.	27	
2 + 02 = 1.2 - 3	4 3.06417	NB 0.6. 98 24.4.	241	
		, MB 0,0, BB 0,4,	3.	
2 + 10 = 123 +	A# 1.25+1.	NE Date PE U.D.	3	. •
		S D D BE D	31	
3 + E- * u- + 'z	A 4.7E+5.	.z -0,63, kz 9,34,	37	79
3 + E= = U2- + 3	AR 1, VE+14	្នាត់ ប្រើប្រាក្សា មិន ក្រុង	33	
				7,111
120 + E- = C- + -2	48 6. 'E.	WE DADA DE UADA	3.4	81
120 + E- # C- +	A= 6.76+C.	, Ta C, Ua pa O, T,	34	

Table C-III-Concluded

```
AB 9.6E+13.
                                                                    162
                                                                 3
                                                                    12
 02 + 0= # U7= + 7 , A= 4,3-+15, .= 0,7,
                                                                 31
02 + NO2- = U2- + .... A 4.54+14, RE 5.0, RE 6.0,
                                                                    131
NC2- + 0 = 102 + 0- , AB 7,25+14, NE 6,6.
                                                                    154
 48 1.5, BE 4,5,
                                                                 4:
                                                                44
                                                                    12
                                            E8 '.',
 NC + C2 = NA+ + U2- , A= 3,66+17,
                                    Nº 1.0.
                                                                     129
                       AS:
                          3.4±+17.
                                                                 44
 NC + MD2 = "U+ + "C2=. A= 3.66+17.
                                                                    142
                                    N= 1.0.
LAST CARD
PSPRCPEL
 NTd=5, PTF(1;=5-1,0, ZTR(1)=0,0,10,6,20,0,30,6,1,754, Z=0,0, PI=1,0,
PI B 1.0. HSLALEED. 020H. 10381. EYITED. 45. 1847LD. 7. VESROU.D. RSTARE12.0.
 HIBD.1, HM1780.1, HTAX89.0, Daleu.005, JF81,
 ISCRF#1, 15USF(1)#9,0,7,6,12,17,44, GPSCP(1)#4+1,UF#2,3+1,06#2,
 NE38Z
 ICGUX#1.
 SEND
```

The water vapor saturation point is reached at station 24.988 as shown in the station output, Table C-XV. Conditions at the saturation point are saved as initial conditions for subsequent condensation calculation pass. An area ratio table is constructed as the integration continues downstream as can be seen by the area ratio printout. Table C-XVI shows the termination of the kinetics pass when the static temperature dropped below 180°R. The condensation calculation pass begins at the saturation point and follows the area ratio table computed during the kinetics pass. Table C-XVII presents this area ratio table plus the first derivative as output at the beginning of the condensation pass. Initial conditions for the condensation pass are included in Table C-XVIII; intermediate printout of pertinent water vapor properties and nucleation rate are also shown. Variables include

PV = vapor pressure (atm)

PVS = saturated vapor pressure (atm)

PRATIO = PV/PVS

RS = critical droplet radius (cm)

EJ = nucleation rate $(1/cm^3 sec)$

TS = saturated vapor temperature (*R)

As the flow continues to expand to higher degrees of supersaturation, the nucleation rate increases triggering droplet growth (condensation). Tables C-XIX and C-XX present the output at typical locations in the condensing region. These tables illustrate the additional printout in the condensing region (where nucleation rate is greater than threshold value). The

Table C-IV. SAMPLE REACTION-SCREENING OUTPUT

********	NO	REACTION	24									
ANIAL POSITION	DESTRUCTION	D(C(1))/DX 7.53288253E-20	4,95092844E-14 6,41667151E-14									
TING SPECIES EN		REACTION 25	28	29	26	, 6 0	30	,	5,69021127E-14	1,136760756-13		
	PRODUCTION	5(1))/D	.197307596-	513964E-1	745279E+1	3107916-1	2291-32E-	151ejņ	WET PRODUCTION	NET DESTRUCTION		

Table C-V. FINAL SUMMARY OUTPUT AFTER SCREENING - RESULTS OF ORIGINAL AND SCREENED RUNS

VARIABLE	ORIG FINAL COND	SCREENED FINAL COND
VEL	3,8070000E+03	3,8000000000000000
DENS	1,179935046-05	1,17993504E-05
TEMP	4.700009016+03	4,700005016+03
.12	7.600002536=01	7.60000253E-01
Ũ2	2,25934373E=01	2,25934373E-01
NO.	4.04039502E=03	4,04039501E-03
N .	5,62376673E+07	5,62376794E-07
U	1,002291986-02	1,00229198E-02
NO+	3,96949443Em07	3,90950464E-07
U2-	2.90312842E010	2.80494462E-10
0-	2,73467463E=09	2,03513961E-09
t •	7,05313769E+12	7,05313731E-12
M05	1.08509134E=06	1.08509778E-06
KSO	1,051615326-04	1.08141731E-08
NCK-	1.02571348F-10	1,02570492E-1U
03	5.70692828E+09	5.70186569E-09
NO3	9,98479043E-11	9,99081970E-11
REACTION .	9	
		•
REACTION :	10	
REACTION	18	
REACTION	19	
REACTION :	20	
REACTION	25	
REACTION	27	
	28	
REACTION :		
	29	
REACTION	2 9 3 5	
REACTION :		

CARD LISTING FOR OBLIQUE SHOCK CALCULATION IN AIR Table C-VI.

##\$55 FRACTION FIELD STREATURE NO. 20 ##\$55 FRACTIONS OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476 OZ .209476	STHERMO NECHEMBO, LISTEL SEND		·	0	
.00934 .00.00 .00.00 .00.00 .00.00 .00.00 .00.00	11				
.0 .0 .0 .0 .0 .0 .0 .0 .0 .0 .0 .0 .0		134			
AR1.0E18. WRI.U. BRG.O. AR1.0E18. WRI.U. BRG.O. AR1.0E15. WRI.S. BRG.O. AR1.0E15. WRI.S. BRG.O. AR1.0E13. WRI.S. BRG.O. AR1.0E13. WRI.C. BRG.O. AR1.0E13. WRI.C. BRG.O. AR1.0E13. WRIAR.S. EXITE.U56928. EFF. BWAXE1.O. DELE.O. WRI.C. DELEGET. OF AVIVEL.S.54682.11.607338.21.724.44.11.158.45	, n				
AE1, DE18, WE1, U. BEG, U. AE1, OE15, NEU, S. BEU, U. AE1, OE15, NEU, S. BEU, U. AE1, SE13, NEU, S. BED, U. AE1, SE13, NEU, L. BED, U. AE1, SE13, NEU, L. BED, U. AE2, SE20, NEI, S. BEC, U. BETS, 483, AE2, SE20, NEI, S. BEC, U. BETS, 483, AE2, SE20, NEI, S. BEC, U. BETS, 483, AE2, SE20, NEI, S. BEC, U. BEL, C. U. BEL, C. U. SE41, S. S. S. S. S. S. S. S. S. S. S. S. S.	0				
AE1.0E16. WE1.0. BEG.O. AE1.0E15. NEU.S. BEG.O. AE1.0E15. NEU.S. BEG.O. AE1.8E 8. NEO.S. BEG.O. AE1.5E13. NEU.C. BEC.O. AE2.9E20. NEI.S. EKITE.U56928. E-4. BEAXELS. DELE.D. KURRES. JPELAGEL. 491951.5.54852.11.607338.21.7244.71.156.45	•				
AE1,0E18, WE1,0: BEG.O. AE1,0E18, NEU,5: BEG.O. AE1,8E 8, NEU,5: BEG.O.1, AE1,5E13, NEU,0: BEC.O.1, AE2,5E13, NEU,0: BEC.O. AE2,9E23, NEL,2: BEC.O. E-4, BEAE1,0: NSTARE,5: EXITE,UE6928, E-4, BEAE1,0: DELE,O.1, NUTE16,1. AS1,41, VE2250.; NSTARE,5: EXITE,UE6928, E-4, BEAE1,0: DELE,O.1, NUTE16,1. AS1,41, VE2250.; AS1ARE,5: EXITE,UE6928, E-4, BEAE1,0: DELE,O.1, NUTE16,1. AS104, U.96:147,18:167,68:100,42:117,28:44,11:158,45					
AE1.0E18. WE1.0. BED.D. AE1.0E18. NED.S. BED.D. AE1.8E 8. NET.E. BED.D. AE1.5E13. NEU.C. BETO.483. AE2.8E20. NEL.2. BEC.D. E-4. BEAXELS. BELE.D. NEL.2. BEC.D. AE2.9E20. NEL.2. BEC.D. AE2.9E20. NEL.2. BEC.D. AE2.9E20. NEL.2. BEC.D. U.96.147.18.17.68.100.42.117.28.44.11.158.45					
AE1,0E16, WE1,0, BE0,0, AE1,0E16, NE0,5, BE0,0, AE1,8E 8, NE0,1, BE0,0, AE1,5E13, NE0,1, BE79,488, AE2,8E20, NE1,2, BE79,488, EE4, BEAXELO, BE79,488, AE2,8E20, NE1,2, BE6,0, AE1,5,54892,11,607338,21,7242,0,31,837449		**			
ARIOE15 NRU.5 BRU.0. ARIOE15 NRU.5 BRU.0. ARIOE15 NRU.5 BRU.0. ARIOE13 NRU.0. BRO.0. ARIOE13 NRU.0. BRO.0. ARIOE13 NRU.0. BRO.0. ARIOE13 NRU.0. BRO.0. ARIOE13 NRU.0. BRO.0. ARIOE13 NRU.0. ARIOE13 NRU.0. ARIOE13 NRU.0. ARIOE13 NRU.0. ARIOE146. ARIOE146. ARIOE146. ARIOE146. ARIOE15. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16. ARIOE16		1,			
##1.9615, N#0.5, B#0.0, ##1.9613, N#0.5, B#0.0, ##1.9613, N#0.0, B#79.483, ##2.9620, N#1.2, B#0.0, ##2.9620, N#1.2, B#0.0, ##4.961.5,3484.0, HSTAR#,5, EKIT#,U56928, ##4.961.5,34892.11,607338.21,724202,31,637449 ##91951.5,54892.11,607338.21,724202,31,637449		. 0#1	WEL. U.	BRO.0.	TION
##1.8E 8.		*1,9E1	NEC. 5.	Sec. D.	200
## ## ## ## ## ## ## ## ## ## ## ## ##		26.0E1	NEU.D.		200
## ## ## ## ## ## ## ## ## ## ## ## ##				77777	LION
### PET PET PET PET PET PET PET PET PET PET	.:	9E	N. 1 . 5.	BRA. Dot.	
** A*2,9620, N*1,2, B*79,483, ** T*431,41, V*22500, HSTAR*,5, Extr*,U56928, ** E** HMAX*1,0, DELE,C1, KUT*16, JPFLAGE1, 491951,5,548952,11,607338,21,724202,31,837449 2004, U,960147,18,167,68,100,42,117,28,144,11,158,45	• 7.	SE1	N. D. L.		
** AFRICAL NELSON NELS. BEG.O. ** TE431.41. VE2500. RSTARE, 5. EXITE, UE6928. E-4. BEAXELO. DELE, CI. KUTELG. JPELAGELO. 491951.5.54692.11.607338.21.724200.31.637449	0	3	1,01		_
. T=431,41, V=2500, RSTARE,5, EXITE,U56928, E=4, bMAXE1,0, DELE,CI, KURE1G, JPE, AGE1, 491951,5,54695,11,607338,21,724302,31,637449	•) (3-042	B870,483	
. T=431,41, V=22503, RSTARE,5, ExITE,U56928, E-4, bEAXEL, DELE,CI, RUNELS, JPELAGEL, 491951,5,54695011,607338,21,724242,31,637449		D D	N . T . N	10.0mp	Z
1951,5,548852,11,607338,21,7243U2,31,837449 6,147,18,157,68,100,42,117,28,144,11,156,45	- 9	.41, VE22503,, RS		T*, U56928,	PSGALER, 00694444,
-,5, ,5,1,491951,5,548852,11,607338,21,7242J2,31,837449,52,056573, 4,2,2014, 4,1HU,96,147,18,157,69,100,42,117,22,144,11,158,45,	ه د د		1		
.5.1.491951.5.548822.11.607338.21.7243U2.31.837449.52.056573. 4.18.2304. 4.180.96.147.18.157.68.100.42.117.28.144.71.158.45.	. f.				
4,1HU,96,147,18,157,69,100,42,117,28,144,11,158,45,	5,1	5,546952,11,60733	38,21,7243,	9,31,637449	1,52,056573,
	4.1HU.96.147	118,157,68,100,42	2,117,28,14	4, 11,158,45	

Table C-VII. PRESSURE TABLE

*****	SHOCK	OCCURS AT TABLE ENTRY	3	*******
1	x	P		DP/DX
\$	-1.0000000E+00	9.20040000E+00	0.	
8	-5.07000000E-01	9.20840000E+00	0.	
3	0.	9.20040000E+00	0.	
4	0.	1.00960000E+02	-2.26	413425E+01
•	1.49196100E+00	1.47180000E+02	-1,11	897475E+01
6	4.54885200E+00	1.07680000E+02	-4,62	266508E+00 ·
7	1.160733806+01	1.00420000E+02	6,32	60068iE-01
•	2.178430205+01	1:17280000E+02	2,16	953827E+00
•	3.18374490E+01	1.44310000E+02	1,35	712132E+00
10	5.20605730E+01	1,58450000E+02	6.99	199590E-01

FORTRAN symbols correspond to the symbols used in the analysis in the following manner for primary variables of interest;

$$XMBAR = \overline{m}$$

$$HINT = \int_{0}^{x} \dot{H} dx$$

$$G = g$$

$$DHPDX = dh'/dx$$

$$DGDX = dg/dx$$

$$DRDX = dr/dx$$

$$RSTAR = r**$$

$$JRATE = J$$

Table C-VIII. SHOCK CALCULATION

		ore false the remarkable as a	
BEGIN SHOCK CALCULAT	ION, AXIA	L POSITIO	N = 0.00000
*****************	••••••	*******	••••••
		•	
(4)			
The second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second secon	per in 1990 d annual is special		
			1
CALCUL	ATED SHOO	CK CONDITI	ONS.
SHOCK ANGLE	(DEG)	1.07112	5386+01
	(DEG)		538E+01 096E+00
SHOCK ANGLE Deflect ang	(DEG)	1.07112	538E+01 096E+00
SHOCK ANGLE Deflect ang	(DEG)	1.07112 8.47695 1.96687	538E+01 096E+00
SHOCK ANGLE DEFLECT ANG P2/P1	(DEG) (DEG)	1.07112 8.47695 1.76687	538E+01 096E+00 101E+01
SHOCK ANGLE DEFLECT ANG P2/P1 PROPERTY PRESSURE (PSIA) TEMPERATURE (DEG-R)	(DEG) (DEG) UPSTREAM	1.07112 8.47695 1.96687	538E+01 096E+00 101E+01 DOWNSTREAM
SHOCK ANGLE Deflect and P2/P1	(DEG) (DEG) UPSTREAM	1.07112 8.47695 1.76687 4 2988-02 218-02 3758-04	538E+01 096E+00 101E+01 DOWNSTREAM

 $HPRIME = h^{\dagger}$

DHPDT = dh'/dT

XMDOT = m

XMOM = M

 $\mathbf{E} = \mathbf{\dot{H}}$

All values of variables printed out under the above symbols are in the internal computing units of poundal, pound mass, BTU, foot, and second.

CARD LISTING FOR NORMAL SHOCK STAGNATION STREAMLINE CALCULATION IN AIR Table C-IX.

	PASS AND 1 ART DETECTOR ART DE BRE DE DE DE DE DE DE DE DE DE DE DE DE DE			NEILUS BEG.D. PERCTION 1	SECTION 2 OEC.O. PEACTION 3	NEG.5. BE6.001. REACTION 6 NEG.0. GE0.0. REACTION 7 NEG.0. GE77.484. REACTION 8 HEU.0. HEACTION 9	AXETON DELEGATO FIXERS DELAGETO	
--	----------------------------------------------------------------------------	--	--	--------------------------	--------------------------------	---------------------------------------------------------------------------------------------------	---------------------------------	--

Table C-X. INPUT VELOCITY DISTRIBUTION TABLE

	<u> </u>	VELOCIT	YTAPLE
1		1,70160000E+03	-3,35797665E+04
-	2,57:10111111-13	1,5158000000+03	-3,15487113E+**
3	2.140000005=13	1,329100006+03	-7,35603113E+04
1	7.7100000000-03	1,442800000E+03	-1,354085606+04
-5	1,02000000000	1,35A70UDDE+A3	
6	1,265000004=02	1,270700006+03	-1,34435798E+04
7	1,5420000000-02	1,1848000000+13	-1,342412456+54
 -	1,799000000±=02	1,09590000E+n3	
7	2,056000008-02	1,01310000F+03	-3,33463035F+14
1 0	2.3130(0006-02	9,275000006+02	-8,33073930E+04
11	2,5700000000	#,41900010E+72	-7,32484975E+14
2	2.427000000-02	7,5650000000+12	-3.31796615E+n4
3	1.004000npF=n2	6,71300000E+02	-7,314229575+64
4	\$,3416H5H0H=H2	5,86230050F+02	-5,30739300E+14
ķ	3, 593090305-02	5.0130000000+02	-3,29766537E+64
6	3.0550CCCUE-02	4,167000006+02	-3,291828796+84
. 7	1,11200000000	3,32100000E+02	-3,28 ^155 45+^4
Ħ	4.369000008-02	2,48100000E+02	-3,262645915+84
ς	4.5260000000-02	1,6440000000002	-3,16881323E+04
,	4,4 48000001-02	4,419\$0000E+41	-3,19944356E+** 4
1	5.140006066-72	\mathfrak{g}_{ullet}	-3,27407U04E+04

Table C-XI. SAMPLE OUTPUT FOR NORMAL SHOCK CALCULATION

BURNAL SECTION STANSATION STRAWNERS IN MALCHEATTON HOW STEEL

TALESTATES S OF COCCUPIENS

t til ,▼v	ur Alamar	J A STOTAH
ロ 5 05 07 (STA)	ა, გაკეცეგაე — ცე	3.77035/2556614
11 17 2 T) ((20.Gen)	4,შეგა ნეციეგაცე	3.71145/2566476
2015 学校マイチーデアデオカ)	აქ,უთქიუმუთნოუბ	3.71145/256



Table C-XII. KINCON SAMPLE CASE-DATA LISTING

008 -	,0452			
HRO	2738		•	
60	18017			
HZ	020			
NS	,42509761			
NO	1005			
OM	00967			
02	,000764			
H	1,69E011 ,00115			
N	1,396-6			
0 .	.000817			
EACTIONS	14469			
02 = 2+0.	A=3,3E17,	N=1.0.	8=0.0.	REACTION 1
NZ B ZUN,	A49,6217,	N=1,0,	B=0.0.	REACTION
NO = N + O.	A=7,2E15,	N=0.5.	B=0.0.	REACTION 3
H2 = 2+H,	A=5.0E18.	N=1.0.	8=0.0.	REACTION
HZO B H + OH,	A-1,17617,	N=0.0.	B80.0.	REACTION S
OH = H + O;	A#2,3E16,	N=0.0.	B=0.0.	REACTION 6
CO2 = CO + O,	A=5,1E15,	iv=0.0.	B=3,58,	REACTION 7
CO * C + O.	YR9'059'.	NEC. U.	8=50,,	REACTION &
ND TBR REAX				•
NO + 0 = 02 + N,	A#3,0E11,	N=+0.5.	B=7,13,	REACTION 9
NU + N + NZ + U;	VESSI CETS!	NEU.U.	8=75,5,	MENUTION IN
N2 + 02, = 2+NO,	A=1,0E13,	N=0.0.	8=79,489,	REACTION 11
co + o2 = co2 + o,		N=0.0.	B=54,15,	REACTION 12
CC + N = C + NO,	A=1,3E10,	N=0.3,	8=0.5,	REACTION 13
CO + O = C + O2,	A=2,4E13,	N=0.0.	B=1,99,	REACTION 14
CC + H = C + OH,	A=1,2E14,	N=0.0.	B=25,83,	REACTION 19
COS + H = CO + OH'		Neg. 0.	821,08,	REACTION 18
0H + 0 = H + 02; 0H + H2 = H2O + H;	A82,24E14,	N=0.0.	8=16,5,	REACTION 17
2 40H B H2O 4 O,	4×5,75613,	N=0.0.	8=20,1,	REACTION 18
H2 + 0 = 0H + H,	4=7,33E12,	N=0.0.	Balk'1	REACTION 20
H2 + 02 = 2+0H,	A=4.98E23.	N=2.5.	B≖U,0, ⊌=85,7,	REACTION 21
NO + CU = COZ + NI		N=0.U.	HB9,93,	MEACTION 22
NO + H = OH + N.	A=3,4E13,	N=0.0.	B=1,38,	REACTION 23
CC + CO : CO2 + C,		N=0.0.	8=9,93,	REACTION 24
AST CARD	-1	4-4141	3-71707	INTERNATION BY
PROPEL				
RSTARBO, 0972, JPFLA	G=1, DEL=.01.	ND3=10. H	l=.00C1. HM1N=	,0001, HMAX=1.
COND(1)=12*0. T=25			a : 4 a a a B b	* ****

Table C-XIII. STREAMLINE GENERATION PROGRAM

CP TIME = 33.799

PP TIME = 192,713

-STREAMLINE	1)= 90.00000	PERCENT OF THE TOTAL MASS PLOW
ZINIT =	0.000000	(NONDIMENSIONAL - S/RC)
PINIT #		(PSIA)
TINET		(DEG ₁ R)
VINIT #	2587,734814	(FT/SEC)
ZFINAL	44,569804	(NONDIMENSIONAL - Z/RC)
HOKSK .	78	(NO, OF PRESSURE TABLE POINTS)

AXIAL	RADIAL	STREAMLINE	PRESSURE
DISTANCE	DISTANCE	DISTANCE	(PSIA)
(Z/RC)	(R/RC)	(S/RC)	
-6,237351E-01	1,010896E+00	0.	6,752437F+01
=5,613616E-01	9,992094E-01	6,345886E-02	6,550104E+01
	9,8842655=01		-6,4662425+0 6
	1111	•	6,246165E+01
			6,019785E+01
			8,430127E+01
			5,614706E+01
	the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the s	the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the s	5,386846E+01
			B,1558615A01
-			4,951774E+01
			4,715078F+01
			4,4783595401
	·		4,274839E+01
			4,038052E+01
그 사람이 가득하게 되었습니다. 아이들이 나를 다면 되었습니다. 그렇게 되었습니다.			-3,8040445+01
			3,573715E+01
			3,374438E+01
			3,152130E+01
			2,935697F+01
			2,726202F+01
그리다 그리는 그리는 그리는 그리는 그리는 그리는 그리는 그리는 그리는 그리는			2,4485325 <u>40</u> 1
			2,406621E+01
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	and the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contract of the second contrac		£,5726585401
			1,4791706+01
			1,386588E+01
			1,295506E+01
			1,206744E+01
2.007078E+00	1,3303035-00	Z,330072E+00	1,120840E+01
	DISTANCE (Z/RC) -6,237351E-01	DISTANCE (Z/RC) -6.237351E-01 1.010896E+00 -5.613616E-01 9.992094E-01 -4.366146E-01 9.792893E-01 -3.742411E-01 9.713276E-01 -2.494940E-01 9.589293E-01 -1.871205E-01 9.545219E-01 -2.7470E-01 9.545219E-01 -2.7470E-01 9.502609E-01 -2.7470E-01 9.502609E-01 -2.7470E-01 9.502609E-01 -2.74205E-01 9.617624E-01 -2.74205E-01 9.680336E-01 -2.742411E-01 9.680336E-01 -2.742411E-01 9.680336E-01 -2.742411E-01 9.680336E-01 -2.742411E-01 9.680336E-01 -2.742411E-01 9.680336E-01 -2.742411E-01 9.680336E-01 -2.742411E-01 9.680336E-01 -2.742411E-01 9.680336E-01 -2.742411E-01 9.680336E-01 -2.742411E-01 9.680336E-01 -2.742411E-01 9.680336E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01 -2.742411E-01 9.838889E-01	DISTANCE (Z/RC) (R/RC) (S/RC) -6.237351E-01 1.010896E+00 05.613616E-01 9.992094E-01 6.345886E-02 -4.366146E-01 9.792893E-01 1.897968E-01 -3.74241E-01 9.713276E-01 2.526764E-01 -3.74241E-01 9.743276E-01 3.780439E-01 -2.494940E-01 9.589293E-01 3.780439E-01 -1.871205E-01 9.545219E-01 4.405729E-01 -1.871205E-01 9.492547E-01 5.654367E-01 -3.552714E-15 9.484006E-01 6.278160E-01 -2.37351E-02 9.492547E-01 5.654367E-01 -3.74241E-01 9.502609E-01 7.525830E-01 -1.871205E-01 9.529274F-01 8.150134E-01 -2.4940E-01 9.529274F-01 8.150134E-01 -2.4940E-01 9.529274F-01 8.150134E-01 -2.4940E-01 9.617624E-01 9.400784E-01 -2.4940E-01 9.6336E-01 1.002766E+00 -2.30535E-01 1.04635E+00 1.299366E+00 -2.30535E-01 1.047388F+00 1.299366E+00 -2.30535E-01 1.047388F+00 1.459507E+00 -2.30535E-01 1.047388F+00 1.459507E+00 -2.30535E-01 1.047388F+00 1.459507E+00 -2.30535E-01 1.047388F+00 1.756937E+00 -2.30535E+00 1.226632E+00 1.756937E+00 -2.26244E+00 1.226632E+00 2.260882E+00 -2.345742E+00 1.226672F+00 2.129195E+00 -2.345742E+00 1.226672F+00 2.129195E+00 -2.345742E+00 1.226672F+00 2.129195E+00 -2.345742E+00 1.226672F+00 2.260882E+00 -2.345742E+00 1.226672F+00 2.260882E+00 -2.260882E+00

Table C-XIII-Concluded

7.4	0 4544755.00	4 4070005400	0.0405505.00	0 5040445404
34	2,156435E+00	1,407000E+00	2,840550E+00	9,581216F+00
35	2,313262E+10	1,447797F+00	3.002596E+00	M.816376F+00
34	2,478760E+00	1,490560E+00	3,173530E+00	A, 084436E+00.
37	2,655024E+00	1,535833E+00	3,355528E+00	7.3680695+00
38	2,845226E+00	1,584638E+00	3,551879E+00	6,652436E+0U
	3,0483445480	1.636619E+00	3,7418435400	5,0747795400
40	3,269320E+00	1,693105E+00	3,989624E+00	5,314822F+00
41	3,508983E+00	1.754318E+00	4,236981E+00	4,702522F+00
42	3,7731426+00	1 821724E+00	4,509604E+00	-4.121188F+00
43	4.065000E+00	1.896143E+00	4.810801E+00	3.581002E+00
44	4,392306E+00	1.979536F+00	5,148564E+00	3,0755816+00
45	4,761856E+00	2,073639E+00	5,529907E+00	2,610257F+00
46	5,1852R8E+00	2,181412E+00	5,966838E+00	2,183015E+00
47	5,675233E+00	2,3060876+00	6,472398E+00	1,797046E+00
48	6,251182E+00	2,452654E+00	7.066704E+00	1.451938E±00
49	6,625002E+n0	2,547389E+NU	7,452340E+00	1,272373E+00
50	7,074635E+0U	5,660688E+00	7,916029E+00	1,086394E+00
-51	7,8153136+00	2,849148F+00	8,680306E+00	8,6672015-01
52	8,571396E+00	3,041428F+00	9,460456E+00	7,013092F=01
53	9.265366E+00	3,218157£ +00	1.017658E+11	5.882468F=01
54	9,975806E+00	3,399777E+00	1.090986E+01	4,941469E=01
55	1.0685n3E+01	3,581162E+00	1,164192E+01	4,249876F=01
56	1,067373E+01	3,574772E+00	1,165490E+01	4,264300F-01
57	1.076411E+01	3.404541E400	1.175387E+01	1.163831E-01
58	1,113066E+n1	3,702849E+00	1,212951E+01	
				4,638904E=01
59	1,125574E+01	3,738687E+QU	1,225963E+01	4,641027E=01
	1,142516E+01	3,787626E+00	1,243597E+01	4,589467F=01
61	1,156782E+N1	3,429747E+00	1,258472E+01	4,641766E-01
65	1,180306E+01	3,899483E+00	1,283008E+01	4,481104E-01
63	1,2072285401	3,977769E400		4,8025945-01
64	1,242947E+01	4,074703E+00	1,348056E+n1	5,042020F-01
65	1,285652E+01	4,183837F+00	1,392133E+01	4,871803E=01
66	1,314293E+01	4.257693F+00	1,421712E+01	4,860031E=01
67	1.3574146+01	4.370228E+0U	1,466277E+01	4.534370F=01
68	1,411218E+01	4,512423E+00	1,521928E+01	4,268618E=01
60	1,4781545+01	4.484822E400	1,5910985401	3,0800185-01
70	1,663745E+01	5,1209416+00	1,781699E+n1	2,243533F=01
71	1.816543E+01	5.496204E+00	1,939037E+01	1.536901E-01
72	1.906691E+01	5.647968E+00	2.030454E+01	-1.482508E=01
73	2,135328E+01	6,248347E+00	2,266842E+91	1.130785E=01
7.4	2.379229E+01	6,939570E+00	-	
75		8.144826E+00	2,520350E+01	4,0966695-02
	2,6844595401		2,848513E+01	1,5236525=02
76	2,942882E+01	9,186441E+00	3,127138E+01	9,9732098-03
77	3,145359E+01	9,973589E+00	3,344378E+01	7,958811E=03
78	4,158523E+01	1,454961E+01	-4.456980E+01	2,352045E=03

Table C-XIV. INITIAL CONDITIONS KINETIC STREAMTUBE CALCULATION AXIAL POSITION = 0. INPUT NORMALIZING AXIAL SCALE FACTOR (FT) 8. 10000E-03

•	PLOH PROPERT						KINET	10 0	KINETIC COUPLING TERMS	TERM	-			
26.50			=	••••	1111	###	14000	22	COUPLING TORN B		12		2,007004706.00 2,000400016.00	
-554			(1) (1) (1) (1) (1) (1) (1) (1) (1) (1)			111	20	S LNEW	218		3,	0000	1.000000000.1	
354	100		7U/L0-0	ê :		77	PAK I	3,	144 104 C	SE SE		19487	-1.689407376-11 0.	
•					70440407	CHENICAL COMPOSITION	8	*	3					
ě	HO, SPECIES MASS PRACTION HOLE		FRACTION	10	FRACTION MOLBG/CC	HOLBC/C		9	89E0188	MA 58	FRACTION	10 HOLE	FRACTION	NO. SPECIES HASS PRACTION HOLE PRACTION HOLEC/CC
•	300	980	6, F200005-02	8.91	41-814401 T.07918-17	7,07918	17	~	2 H20	2,736	0006-01	2.9	10-225-01	2,7380006=01 2,9909286-01 7,2421E-1
+	8	3	30050001304	4.4	71336-01 31A4098418	3,44086	3	+		3,000	0005-02	-	184036-01	3,0000006-02 1,9824036-04 4,74056-44
•		4,89	4,2509766-01	2.0	61978-01 7,25066-18	7,29066.	•	•	2	2,000	0006-03	1.31	16656-03	2,0000006-63 1,3116698-03 3,1848E-1
•	*	9,67	P.6700068-03 1.11	1,11	.86218-02 2,71688-17	2,71606	17	•	30	7,64	000E=04	•	10694E-04	7,640000E=04 4,698654E=04 1,1499E+16
-		4	6,6800005m11 8,74	8,20	#0018-11 4,72358-00 10 H	4,723964		=	_	1.180	0000-03	2,24	32646-02	1,1500006-03 2,2482646-62 5,45166-17
=	=	1.3	1,3000006-04	1.	1,9986496-06 4,74166-13 18 0	4,74166	27	=		6.170	000E-04	7.5	191 B4E - 84	6.170000E=04 7.509:84E=04 1.8427E+14

Table C-XV. KINETIC STREAMTUBE CONDITIONS AXIAL POSITION 2,41693E+01

	PAGESCAE VELOCITY	(P81A)	14) /850)	5,74884214E+09 6,78042174E+02 7,44408137E+03		NON THE	7684 B			6199	-7.619937926-01	**	
	DENSITY (LB/ MINALPY SHO MINALPY SHO MINALPY SHOP MINALPY SAPER MINALPY SAPER MINALPY SAPER	ALPYCHO (BT) ALPYCHO (BT) CAPACITY(BT) EN GAMMA	TV (LB/FT3)	24 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4		1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	CURRENT STEP SIZE PERCENT ENTHALTY CHANG 1.0 - SUHHATION C(1) HAXIMUM RELATIVE ERROR POVERNING FOUNTION	C.1.)		0440 0470 14040	1.024000006-01 -8:100478816-00 -1.436006976-11 1.077177266-03	2645	
0	SPECIES		HASS FRACTION HOL	HOLE FRACTION	-CMEMICAL COMPOSITION MOLEC/CC NO. SPE	NO.	SPECIES	HASS	FRACTION	T CL		FRACTION	#OLEC/CG
	202	7,11	7,116331E-02	3,216600E+02	4,0189E+15	~	H20	2,623	2,827214E=01		3,121782E-01	E-01	3,9004E-16
	3	4413	1,979046E=01 1,4	1,4054966-01	1,7561E+16	•	# CH	1,994	1,9996145-02		1.9931406-01	E-01	£,4653E+16
	N 2	4,29	4,2509846-01	3.018460E-01	3,77136+16	•	0	2,001	2,001244E=03		1,326677E-03	E-03	1,6576E-14
	#0	1,74	1,748476E-06	2,0450716-06	2,55516+11		20	3,637	3,837964E-04		2,385905E-04	E-04	2,9810E+13
	U	4,02	4,0299505-12	6,67486 <u>15</u> e12	4,33936+05	97	*	7,289	7,2854746-84		1,437A03E-02	E-05	1,7964E-15
	z	1,07	1,876187E-08	2,664416E-08	3,32906+09	12	0	7,34	7,349727E=07		9.138047E-07	E-07	1,1417E+11
	4	SATURATION Ze 2.	110N POINT H20 2,4988506+ 2,5983806+		214988500E+01 5441789E+02 AREA RATION AREA RATION		-00	487±320E+03 00E+00 67E+00	P 0		i		
1		.7	2.5602900E	00E+01	۱.	F	1.12364706-00						



Table C-XVI. KINETIC STREAMTUBE CONDITIONS AXIAL POSITION 3, 52285E+01

•	PER PROPERTIES	EN7155		Z X	KINETIC	COUPL ING	TERMS				
T .	RESSURE	CA100	6.97243082		COUPLING	7684		40		- 10703302E-07	
•	ELOCITY FEBRUARIA	(FY/SEC)	7,86861495E+03	r en a		MARGAL MASSAGE	962634				
	BENEITY (LB.	LO/FT3) HO (BTU/LB)		: S N	ENT	STEP SIZE		•	0960	00006-01	
4 2	-	ACITY(BTU/LB-DEGR	-	ĺ		SUMMATION	CHANGE	1	4114	-5.0433950-6-00	
1	NI	- 5	1		₹ ‡	RELATIVE	ERROR	~	0	4003912E-04	
7	CIDENCIDO EOS	H(1) (F12/SEG2)	69,72961720Ee07	E-07 CHEMICAL COMPOSITION	904	**					ı
9	SPECIES	SPECIES MASS FRACTION MO	HOLE FRACTION		2	SPECIES	MASS	FRACTI	NO	MASS FRACTION MOLE FPACTION	HOLEC/CC
	203	7,116333E-02	3,216602E-02	9,3960E+14	~	H20	2,827	2,827214E=01		3,121783E-01	9,1190E+15
*	1 60	1,0790465-01	1.4054965-01	4,10565415	+	4	1,996	30-3679666 ¹ 7		1,973145E-81	8,7630E+15
•	8	4,2509846-01	3,018461E-01	8,81726-19	•	9	2,001	2,001243E-03		1.326677E-03	3,8754E+13
~	10	1,63669E=06	1,914335E-06	5,59206+10	-	05	3,837	3,837965E-04		2,389904E-04	6,9695E+12
•	9	4,6207645012	6,6242545-12 1,9496Engs	8486848	2	=	7,2A5	7,2A50865-04		1,437727E-02	4,19976-14
#	2	1,0695016-08	2,6549226-08	7,75536+06	12	0	6,333	8,33311E=07		1.0360716-06	3,0265E+16
	ı		77006-01 12006-01 12006-01 13006-01	44444	• • • • • •	7,51038006 7,51038006 1,16166246 1,67310706					
		Z4 4,0962900E Z8 4,1782100E	06-01 06-01	AREA RATIO		. 0366796E-0 . 0732400E-0	7946-00 4006-01				
		4									

Table C-XVII. AREA RATIO TABLE

-1			
1	2,4985000E+01	1,00000000000000	0 •
2	2,51933000E+01	1,05773674E+00	3,02360939E-01
-	2,549810005+01	1,12784784E+A8	3,402682915-01
4	2,56029000F+01	1,197110636+00	3,75357475E-01
5	2,58077000E÷01	1,27759346E+0U	4,12841205E-01
-	2,49125000E+A1	1,364210396+00	4,52080543E=01
7	2,62173000E+01	1,46309663E+00	4,93586234E=01
8	2,64221000E+01	1,56438331E+00	5,34413881E=01
-	2,6626900E+81	1,681992566+90	1,74269361E+01
10	2,68317000E+01	1,80360412E+00	6,11919442E-01
11	2,70365000E+01	1,93263476E+00	6,46n7n464E=n1
13	2,724130005+01	€,04^23459€+∩0	6 ₁ 75455076E=01
13	2,74461000E+01	2,20930117E+00	6,98907936E=01
14	2,76509000E+01	2,35450728E+00	7,15425464E+01
15	2,78557000E+01	P,50233944E+00	7,24108045E=01
16	2,90605000E+01	2,65113916E+00	7,24603912E-01
17	2,32653000E+01	2,79913720E+00	7,16138891E-01
-18	2,347010005+01	2,9414494 5 6+00	7,125834095-01
19	2,87261U00E+01	3,12754171E+00	7,43507550E-01
20	2,89921000E+01	3,32514552E+00	7,96910279E-01
21	2,928930005+01	7,574344585+00	*,32199993E-01
22	2,95965000E+01	3,83644914E+00	8.480978576-01

Table C-XVII-Concluded

23	2,99037000E+01	4,09743291E+00	8,37479712E-01
24	3,02400005+04	4,350994475+00	*,00741477E=01
25	3,05181000E+01	4,58940859E+00	7,27663481E+01
26	3,09277000E+01	4,87258586E+00	6,17132395E-01
27	3,443970005+01	5,15415740E+40	5,32042607E-01
28	3,19517000E+01	5,41739749E+00	4,85100428E-01
29	3,25661000E+01	5,70457492E+00	4,17098491E-01
30	3,33053000E+01	6,01534988E+00	3,444976n9E=01
31	3,44093000E+01	6,33955291E+00	3,51928117E=01
32	3,60477000E+01	6,95232330E+00	3,95030554E-01
73	3,48440005+01	7,31038000E+00	4,62962691E-91
34	3,76861000E+01	7.71084136E+00	5,19581533E-01
35	3,85053000E+01	8,16166239E+00	5,87320313E-01
36	3,932450005401	A,473106965+00	
37	4,01437000E+01	9,25869520E+00	7,71221106E=01
38	4,09629000E+01	9,93667562E+00	8,99477990E-01
30	4,478310005+01	1,078239995+01	9,713481696-01

Table C-XVIII. INITIAL CONDITIONS KINETIC STREAMTUBE CALCULATION AXIAL POSITION = 2, 49885E+01 INPUT NORMALIZING AXIAL SCALE FACTOR (FT) 8, 10000E-03

				MOLEC/CC	3,06176-16	07-904-0	1,30076-14	2,35936-13	\$103861015	9.2212E+10		F-00 75- 456,667	E-00 TS- 454.047	E-00 75- 494.067	1-00 T3- 496, 064	E-00 TS- 494,040	100
		1.000000000.1 -0.000000000.1 -0.0000000000		SPECIES HASS PRACTION HOLE PRACTION	3,1217026-01	1940417 - 10-917124617 - 20-961726617	1.3266776-03	2,3090096-04	1,4377016-00	9,340096-87	00-90	PRATIO: 1.14619026-00	PRATIO: 1.14619686+60	PRATIO 1.1462198E-00	PAATION 1.14624826+88	PAATIO: 1.1463302E-06	PRATION 1,14647426-09
WINETIC COUPLING TERM	COUPLING TER- B	•	MF0611100	NO. SPECIES MASS PRAC	2 H20 2,8872146-81	\$679000 ¹ 7 - 7 -	6 NO 2,8812436-83	8 02 3,837944E=64	-t- + -1-3083605-6+	12 0 7,5120976-67	AREA RATIO . 1.00006-00	FXPL 0. 24004746-04	PVS = 0,2498617E-04 EXPJ= 0,	PVS & 9.2496071E-04	PVS = 9.24936336-84 EXPL= 0.	EXPL. 0. 2404020E-04	PVS - 9.2471930E-04
	4,9001,426-12 COU		CHONICAL CO	PACTION HOLE PRACTION HOLEC/CC	3,2166016-02 3,1793E-15	\$+4004006-01 - 1130756-14	3.010461E-01 2,9797E-16	2.0214076-06 1,99968-11	4,4744316-12 4,98485-05	2.6416796-08 2,62796-89	MBAR+1 = 0.	PV= 1.04021996-03	PV0 1,060217fe05	PV- 1,0602133E-03	PVP 1,0402046E-03	PV- 1,06016096-03 R8- 0,07319646-07	Pve 1.06016835-03
	UNE (PEIA) 17 (PT/SEC)			NO. SPECIES HASS PRACTION HO	7,1163315-02	4,0700466-04	4.2509646-01	1,720320E-06 2.	4,000006-12 4,	1,0742596-06 2.	.0	Z & 2,49895986-01	Z 0 2,49999986+01	2 # 2,4001508E-01 17ARCTO 0	2 0 2,49925006.01 TARCTO 0	Z 0 2,40049006+01	70-9006900+ 7 - Z
	100 N	20 17 10 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	5	40. SPE	300 T	8	5 w2	7 04	9	7 77	. 270						

Table C-XIX. KINETIC STREAMTUBE CONDITIONS AXIAL POSITION 2, 70052E+01

		# # # # # # # # # # # # # # # # # # #		HOLEC/CC	1,29226-16	0.10706-19	9,49106-13	9.87696-12	8.9517E+14	3.86676-10		TERM # 9,92415586+21 6 # 1,529642E-FS	발	46-02	7E-02 TS- 436,717	9E+02		4E+02 TS= 436,497	4E-f2 TS- 435,687	76-02 75- 436,677
	0.	2,000000000000000000000000000000000000		TION HOLE FRACTION	3,121667E-01	2 1.0731746-01	1,326599E-03	2,3850496-04	1,437409E-02	7 9,341151E-D7	146-00	E = 1,465A714Ee12		PRATICS 4,006444E+02	PRATION 4.0410787E-02	PRATION 4,0764409E+02		PRATIO: 4.03637046+02	PRATION 4.05403846-02	PAATION 4,07186376.02
OUPL IN	COUPLING TERM B	P SIZE HALPY CALNOS TION C(I) TION C(I)	CHEMICAL COMPOSITION	NO. SPECIES MASS PRACTION	.5 2 H20 2,027104E=01	.5 4 H2 5,00045E-6	.6 6 NO 2,001274E-03	.0 0 02 3,8%022E-04	15 10 H 7,285470E=04	10 12 0 7,513071E-0	AREA RATIO . 1,00094E+0	2.0000000E=n3 RPRIME 1.5298692E=n5 HINT		6252116E 8886666	PVS = 8.5417458E=07 EXPJ= 1.2276672E=05	PVS = 0,497772E=07		PVS = 0.5417450E=07 EXPJ= 1.2211901E=05	FXP1 1.2442071E-07	FYS - 8,4574112E-07 EXPJ- 1,7475764E-05
H 0-356496949		1,096749960 1,096731061600 1,096319660 1,09674119600 1,38696960 1,4869860 1,4869860 1,4869860 1,4869860	TEDINENS	MOLE FRACTION MOLEC/CC	3,216694E-02 1,3315E-15	11-1321016-01 5,01016-12	J. n105116-01 1,24956-16	2,n21531E-06 8,3661E-10	6,4745326-12 2,76296+05	2,661723E-08 1,1018E-09	HRAT-1 = 9,99985E=01	290E+01 DELZ = 170E+13 XIBAR = 100E+13	200	PVe 3,49	1 PV= 3,4517867E=04	Pve 3,4477629Eeg4	2,7011250CE+01	TOTAL 3,4477690Es04	Pve 3,4457531Eest ASe 2,999750Eest	Re 2,99420026-04
PLOW PROPERTIES	PRESSURE (PS1A) VELOCITY (FT/SEC) TEMPERATURE (DEC.E)			NO, SPECIES MASS FRACTION MOLE	C02 7,116439E-02	3 CO 1.0706765-01	9 N2 4,251048E+01	7 OH 1.728346E-06	C 4,6299266-12	11 v 1,874287E=08	*8141.52987E-05	Z = 2.7609290E+ TEH2 = 1.0210476E+	04000	2	2 e 2,70042506+n1	Z • 2,7011250E•01	STEP SIZE HALVES AT Z .	Z = 2,7100250E+01	Z = 2,7010259E+C1	Z = 2.73112506+01 TARCT= 0

Table C-XX. KINETIC STREAMTUBE CONDITIONS AXIAL POSITION 2, 90277E+01

	e special de special de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servicion de servici												80	E1,7472219E+08	
	Ť				MOLEC/CC	9,9604E-19	3,79426-15	2,5525€+13	4,5903E-12	2,74426-14	1,7972E-16		46.8		
		9.00000000000 -8.74568898-60 -4.234834715-12 1.318145978-03	•	•	NO. SPECIES HASS FRACTION FOLE FRACTION	3,105306E-01	1,0776446-61	1,329n39E-03	2,391592E-04	1,4417046-12	9,3632626-07		4,1369606E-12	7,8795803E+23	9.4144203EABB
7£8+\$	• •	w w			HASS FRACTION	2,8117356-01	2,0030316-62	2,005943E-03	3,8462486-04	7,3010056-04	7,5201736-07	3,362296.00	# I I		
KINETIC FOUPLING TERMS	COUPLING TERM &	CURRENT STEP SIZE PERCENT ENTWALFY CHANGE 1.0 - SUMMATION C(1) MAXINUM RELATIVE ERROR		MOI 1 1 90	NO, SPECIES	2 H20	4 15	02	• 02	10 H	12 0	AREA RATIO .	5.0000000E-34	15005776-10	-1.0448248E-01
M I M I	E-03 COUP.		E-07	CHEMICAL COMPOSITION	HOLEC/CC	611086E+14	21-31-04-15	5,00736+15	3,8392E+10	1,26415+05	5,1209E+08	78446-01	DELZ = 5.00		
	7,305200036- 6,506102406- 7,652263996-	91076192 27515332 979638623 37663888	-5,62218244E•	Ŭ		3,224260E-02 6	1,4000466-01 3	3.029696E-01 9	2.026316E-06 3	4,4003316-12 1	2,668024E-08 5	MBAR-1 - 9.97	F-01 DE	E+21 RS	E-03
	(PSIA) (FT/SEC)	TV (LE/FT3) LEVENO (8TU/LS) OLECHIA HELGHT CAPACITV(8TU/LS-DEGR) N GAMMA	SUM C(1)+4(1) (F72/SEC2)		NO. SPECIES MASS FRACTION HOLE FRACTION	7,1316916-02 3,	- 1-0033186-01 1.	4,2601596+01 3.	1,732051E-06 2,	4,6385635-62 6,	1.8783046-08 2.	*2,15566E*03 HBA	ERM2 = 2.6380530	407 - 1,3242619	MD07 1.3573853
FLOW PROPERTIES	PRESSURE	DEMBITY (LE ENTHALTY (LE HEAT CAPACE FROZEN BARE	SUM C(1)		NO. SPECIE	7 005	85	5 N2	7 OH	3 6	11 N	MBAR	mF e	22	N .

C. 8 REFERENCES

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Appendix D

SURFACE

DEPOSITION AND SURFACE EFFECTS COMPUTER PROGRAM

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Appendix D

SURFACE

DEPOSITION AND SURFACE EFFECTS COMPUTER PROGRAM

D. 1 INTRODUCTION

The computer program SURFACE described in this appendix is a subprogram to the Plume Contamination Effects Prediction Computer Program, CONTAM. It computes the effect of direct plume impingement on sensitive satellite surfaces in terms of changes in the thermal and optical properties of the surfaces.

a. Objectives

- 1. Analyze and model the direct impingement interaction of the plume from a chemical thrustor with typical satellite surface configurations and materials to predict changes which occur on the satellite surface by chemical deposition and mechanical abrasion.
- 2. Analyze and model the surface changes which occur during subsequent periods when the thrustor is not operating.
- 3. Analyze and model the behavior of satellite optical ports and thermal control surfaces subsequent to their being affected by plume impingement and space exposure sequences to predict changes in transmissivity and thermal balance factors.

b. Data Base

Required input to the SURFACE Subprogram includes:

- 1. The gasdynamic, thermodynamic, and chemical constitution description of a plume as computed by the <u>MULTRAN</u> (Appendix B) and <u>KINCØN</u> (Appendix C) subprograms.
- 2. A configurational and material description of the external sensitive surfaces of the spacecraft.

c. Output

The <u>SURFACE</u> Subprogram of the <u>CØNTAM</u> program includes determination of plume impingement effects in two related area:

- 1. Surface chemical/mechanical effects on a satellite caused by direct plume impingement or exposure to the space environment.
- 2. Optical/thermal effects on satellite functioning caused by the surface chemical/mechanical changes.

d. Program Outline/Summary

The program is a generalized system model consisting of:

- 1. A general configurational description of a satellite with multiple thrustors and functional projections such as solar cell arrays—up to 175 separate structural elements can be treated.
- 2. A variety of structural materials commonly used on the exteriors of satellites, such as alloys, thermal control paints, platings, viewport materials, and solar cell materials. These may be combined to model multilayer surfaces. Up to 25 different materials may be used.
- 3. A variety of typical plume constituents which may be gases, vapors, or condensed droplets/particles. Up to 25 different constituents may be treated.
- 4. The configuration of the plume and its local properties are a necessary input. This information may be obtained by examination of the <u>KINCON</u> and <u>MULTRAN</u> subprograms.
- 5. Program output consists of a description of the current condition at up to 175 satellite locations. The description includes, for each location, structure composition and surface state, deposit chemical composition, deposit quantity, deposit state and surface condition, surface and substrate temperatures, thermal radiation coefficients for heat balance calculations, and optical transmissivity coefficients for sensor efficiency calculations. Average values for thermal and optical coefficients are also listed.

e. Program Physical Configuration

The <u>SURFACE</u> program consists of about 7 boxes of punched cards. (The size of an object deck in punched form has not been estimated.) The standard data supplied with the program consists of about 1/2 box of cards. If a complete set of plume characteristics data is prepared from <u>KINCON</u> and <u>MULTRAN</u> and used as input to <u>SURFACE</u> in punched card form, this will add one to four more boxes.

The physical size of the program when in card form makes it preferable to store the program and data in a magnetic tape or disc file when it is in use, and just to keep a backup copy on punched cards.

The tape file of <u>SURFACE</u> prepared at MDAC is a multi-file master file which has the structure illustrated in Figure D-1. This figure, a Table of Contents for the tape, is the first file on the tape, and it is normal practice at MDAC to copy this contents file to output every time the tape is used.

It is advisable to keep this table up to date by correcting it whenever modifications are made to the program or data and a new file created with the modified contents. This is readily done because the second file on the master tape is the UPDATE \emptyset PL for the Table of Contents. (The Deck Name is STR.)

D.2 ANALYSIS

a. Satellite Structure

SURFACE accepts data input to describe the external surface of the satellite. The surface is divided into a preselected, arbitrary number of locations given in a cylindrical coordinate system. Additional inputs include the locations of all projections from the surface, the height of such projections, and the angle they present to the main axis of the satellite. The initial surface temperature at each location is noted. The program prints a set of tables showing all configuration data.

In the satellite-based cylindrical coordinate system, the origin is placed at the forward vertex of the satellite, with the satellite axis lying on the X-axis. The cylindrical coordinates of any point in this system are given in terms of (X, R, 0), with 0 measured in radians in the plane of the Y-Z axis from an arbitrary zero (see Figure D-2).

For purposes of defining the various conditions and effects on the surface of the satellite, the exterior is divided into segments, each with a specified area. Each segment is given an identification number, and its location is recorded in the cylindrical coordinates of its centroid. The shapes of the segments are appropriate to the surfaces on which they lie—curved squares on the cylindrical surface, annular segments and circles for the ends, etc. The assignments are completely flexible and can be changed by simply changing the data cards. In general, there is no need to describe the satellite exterior completely. If the areas which can interact with the plume are already known, then the input can be limited to these areas.

b. Projection Configuration

Data are input to describe the configuration and structure of projections above the satellite surface, including sensors, solar cells, and thrustors. These are also broken down into conveniently sized segments. Tables are printed showing the data for each projection.

c. Structural Materials

Data are input to describe the structural-mechanical properties of typical materials used for exterior structures of satellites. Materials include aluminum, gold plating, solar-cell cover glasses, infrared ports, windows, ultraviolet ports, and white and black thermal-control coatings, plastics, etc. Typical or handbook data are currently input. Definitive data for specific alloys or compositions can be added easily when information on their behavior is required. The input data are listed.

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Figure D-1. Table of Contents of SURFACE Master Tape

Figure D-2. Satellite Coordinate System

d. Optical and Thermal Coefficients for Structural Materials

Data are input to define properties such as a solar absorptivity (α_s) , thermal emissivity (ϵ_{TH}) , transmittivity (τ) , and reflectivity (ρ) at selected wavelengths, etc., as functions of surface finish for the structural materials. Typical or handbook data are currently supplied. Specific data inputs for selected compositions of materials can be added as required. The input data are listed.

e. Physicochemical Properties of Propellants and Deposits

Data are input to define the physical and chemical properties of plume species and deposits at selected reference temperatures. Properties of propellants at other temperatures are then calculated. However, there are insufficient data for some of the condensed reaction products to allow accurate extrapolation of these properties, and data for these materials are treated as constant with temperature.

f. Optical and Thermal Coefficients for Deposits

Data are input to define thermal and optical coefficients— a_s , ϵ_{TH} , a_s , and ρ —as functions of temperature. Input data are listed.

g. Other Data

When certain data normally input in c. to f. are not supplied, appropriate values are estimated from other data that are input. Thus, if the critical temperature is not supplied, it can be internally estimated from the 3/2's rule and the boiling point: $T_c = 3/2$ Tp. As another example, missing optical transmissivities at various wavelengths are estimated from the refractive index. A listing of missing property data for each material is made.

h. Initial Assignment of Materials

Data are input to assign specific structural materials to each satellite segment. As many as three layers of materials may be assigned to any segment. Usually only one or two layers of structural materials are input, leaving at least one layer for deposits formed by the plume. The data are entered into two arrays; one is the surface description array SURDES, and the other is a transfer matrix used to search for properties corresponding to the materials and their conditions.

A table is printed showing the materials assigned to each segment of the satellite and projections, and the initial surface condition (finish or roughness, and temperature).

i. Suppression of Listing

The tables listing initial input data can be suppressed by a flag set during initialization. This reduces redundant output when the same initial conditions are used in a sequence of runs. Error, warning, and timing messages are not suppressed.

j. Optical/Thermal Surface Parameters

This routine takes the transfer matrix of materials assignments to segments and searches the stored optical and thermal coefficients for the appropriate data. Interpolations/extrapolations are conducted if the specific data needed have not been entered during input. (The optical and thermal properties of structural materials are tabulated as functions of surface finish; the properties of the deposits are given as functions of surface temperature.) The data appropriate to specific segments are entered in the array SURDES.

The interpolation/extrapolation routines include the following features:

- 1. No extrapolation of optical/thermal properties is done below the value given for the lowest temperature or surface finish in the tables: the minimum value is used.
- 2. Extrapolation above the highest temperature/surface finish is a simple linear ratio from the values for the two greatest temperatures/surface finishes.
- 3. Interpolation is by ratio from the nearest adjacent values, using essentially the method given by Wiberg (Reference D-1).

The new description of the surface is printed out.

k. Effective Values of Heat Transfer Coefficients

This routine calculates the values for solar absorptivity and thermal emissivity and their ratio, averaged over the whole satellite exterior (excluding projections). The calculated values are printed out.

1. Transmission Through Transparent Segments (Windows, Optical Ports)

The percentage transmission of an incident normal ray through the actual thickness of transparent material is calculated at eight wavelengths (infrared to ultraviolet) and the results are listed. Averaged values are calculated for devices which extend over more than one segment.

m. Test/Precalculated Surface State

Card data are input describing the physical state of selected structural segments. Such data include presence of deposits, thickness of deposit and/or thickness of original surface, surface finish, and surface temperature. The data are listed.

A flag is set during initialization to activate or bypass this module.

n. Optical/Thermal Surface Parameters and Heat Transfer Coefficients

The program then returns to the calculation of surface parameters and coefficients above, this tests for the precalculated surface state, and lists the values obtained.

o. Reinitialization

If the precalculated results module (m and n above) is active, the results from this module may be discarded after listing, or they may be used as the basic condition prior to plume and space exposure. A flag set during initialization selects the mode to be followed.

p. Exposure Sequence

A set of input data define the exposure sequences to be undergone by the satellite. The definitions include types of exposure (plume or space), their sequence and durations. Flags and counters are set by these inputs to select the sequence of computation routines appropriate to the sequence of exposures.

q. Plume Impingement Locations

Those satellite segments upon which the plume impinges directly are identified by considering the configurations of the plume and satellite. Only these identified segments are used in calculation of plume impingement effects.

The intercept of the plume with the satellite is calculated. The variables used are satellite configuration, thrustor location, and plume geometry. A paraboloid shape for the plume is assumed. The latus-rectum of the plume paraboloid, which defines the shape of the plume, is equal to four times P. P is a function of thrustor size, configuration, etc., and of the time in the cycle (pulse transient).

The locations impinged by the plume are functions of the geometry of the plume and the configuration of the satellite. The border of the impinged area is founded by simultaneous solution of the equations for the plume and the surface. The array SURDES is searched to find all segments whose coordinates are on or inside the border. These segment identification numbers are put into the array called AFFSEG.

The computation is done in two separate modules. The first is the intersection of the plume with the satellite surface. Currently this module is restricted to satellites of near cylindrical cross section. The second module determines the intersection with projections. The computation assumes an approximately plane surface on the projection which can lie at any solid angle to the axis of the plume. The configuration of the system used for these calculations is the data originally entered in the program. There is no provision for recalculating the intersection of movable projections after they have changed position.

Impingement on one location by the plumes from two separate thrustors fired simultaneously is not treated; however, sequential firing of two or more thrustors is treated.

r. Space Effects Locations

The satellite segments which have a surface deposit formed by previous plume exposure cycles are identified. Only these segments are used in calculation of space exposure effects.

The identification is done simply by searching the array SURDES, which lists current surface state, for the presence of deposited materials as the topmost layer of a segment.

s. Transformation of Coordinates (Plume Exposure Only)

The <u>MULTRAN</u> and <u>KINC ϕ N</u> subprograms treat the plume in thrustor-based cylindrical coordinates. In order to identify the locations of impinged satellite surfaces in the plume, a transformation to satellite-based coordinates is performed on the plume flow field. These calculations require inclusion of a radial angle in the plume coordinates because the cylindrical symmetry of the isolated plume is lost in the presence of the satellite structure. The origin for measuring this angle is defined by the plane in which the central axis of the thrustor and the central axis of the satellite both lie.

t. Surface Properties

The physical and mechanical properties of the surfaces of the affected segments are found and entered in the array SEARCH. These properties determine the type of interaction with the plume.

u. Surface Effects

KINCON and MULTRAN outputs are used to determine data on the characteristics of the plume at the location of each impinged segment at each time slice. If complete data on the plume characteristics are not available, the program assumes that conditions prevailing st the previous time increment or at the adjacent segment (or both) apply to the segment of interest. This allows the program to be run with as few as a single set of plume characteristics—in such a case there is automatic provision for startup and trail-off transients.

As a first step, the abrasion wear is calculated for a single time increment of approximately a microsecond (the exact width is controlled by the user). The amount of abrasion in this interval is then compared with a standard wear depth value, input by the user (and defaulting to 10^{-10} inch). If wear is less than the specified amount, the program considers wear as insignificant and branches to calculation of condensation from the plume vapor. (The two processes are treated as mutually exclusive.) After completing the condensation routines, direct deposition of plume droplets is calculated.

The results of the three processes are stored temporarily. The effects are then calculated for the next time interval, using the results from the previous interval as the base.

v. Abrasion

All abrasion is assumed to be the result of impacting particles (droplets). There are no provisions for abrasive wear resulting from gas impact or for thermal ablation. If abrasion occurs, the impacting drops are assumed to depart along with the abraded surface material. The wear relation used is the fatigue wear term of Neilson and Gilchrist (Reference D-2). The rate of material removed per weight of impinging condensed phase is calculated by means of tabulated values for the fatigue wear parameter for the several materials, plus constants for the equation used to calculate the value at specific velocities.

The wear rate constant (lb_m material removed per lb_m material impinged) for a particular material is a function of the impingement velocity up to a lower critical velocity; it then becomes constant. The critical velocity is a property of the material undergoing abrasion. Some data available suggest that there are upper critical velocities at which the form of the wear equation changes, but the data are insufficient for definitive application, therefore the program does not model the changes.

In the calculation of abrasion effects, the droplets are grouped into two size categories: the uncombusted material (large) and the condensed combustion products (small). An average diameter is used for each size category.

Two effects are calculated—the wear depth and the resulting surface finish. If the abrasion is sufficient to completely remove the surface layer, subsequent calculations are conducted with the characteristics of the newly exposed layer.

w. Condensation

The model used for the condensation process is one which uses the analogy between heat and mass transfer, based on Trebal's model (Reference D-3). The model uses heat transfers coefficients, without requiring input data on mass transfer or deposition rates.

The determination of deposition requires the diffusivities of the condensing species in the gas stream. These are calculated from the boiling point, molecular radius, and density. The assumption is made that each condensable species diffuses independently through a medium consisting solely of nitrogen gas. Next, using the specific heat, viscosity, and molecular weight of the plume, the Prandtl number and Schmidt number are calculated.

For each condensable species, separate film heat transfer coefficients are calculated based on plume properties and on substrate properties. The one based on plume properties is used to define a mass transfer coefficient. Then using the bulk heat transfer behavior of the substrate, the heat of condensation of the species of interest, and the heat and mass transfer coefficients, the rates of deposition of both heat and mass are obtained by simultaneous solution of the equations, with vapor pressure of the condensing species used as the pivot in the calculations. The process is carried out for each species, then the results are summed. The final substrate and surface temperatures are also calculated.

Chemical reactions between respective species as they condense are also computed to define the final composition of the deposit. The calculations are repeated for each time increment in the pulse, using the appropriate plume characteristics.

x. Deposition

The direct contact of plume particulates is then treated. Droplets which are not moving fast enough to cause abrasion wear are assumed to have a probability of adhering to the surface. The probability is calculated as a function of droplet mass, droplet velocity, and impingement angle; then the total quantity of deposited material is determined from the probability and the droplet mass flux.

The calculations are repeated for each time increment.

v. Net Effects

After exposure to the plume is complete, reactions between species deposited at various times and by various processes are calculated to define the final deposit condition.

The surface finish of the deposit is calculated, based primarily on temperature and deposit thickness.

The total quantity (depth) of surface deposit at each segment is compared with a user-input standard, typically about 10-6 in. If the quantity deposited is smaller than the standard, then the deposit is ruled negligible and the data are deleted.

z. Space Exposure Effects

The model used for space exposure effects is primarily a calculation of evaporation rate of volatile deposits as a function of surface temperature, heat transfer from the substrate, and vapor pressure and heat of evaporation of the deposits. A zero back pressure is assumed. The method of calculation is similar to that used for evaporation of surface films in the thrustor chamber (see Appendix A).

aa. Recycling/Termination

The program prints a summary of the calculations carried out for the completed exposure. It then checks for instructions about further exposure cycles, and proceeds to follow the instructions. If all cycles are complete, a closing summary is printed.

ab. Program Operations Information

A clock is kept to measure the length of time each overlay requires, and this information is listed after return from the overlay. To keep core and disc space reservations to the minimum required for program execution, internal field length redefinition (SRFL) and disc file release (DRØPIT) are used extensively. Appropriate messages are printed in the Day File when these routines are called. The field length redefinition routine SRFL, when

used to increase the field length, sets all locations in the newly acquired core region to zero. SFRL is written in COMPASS and included in <u>SURFACE</u>.

A counter is used to keep track of all calls to external routines which are part of the program (library and system calls are not monitored). At normal termination, this count is listed by a program subroutine. For abnormal termination, going to EXIT, a small auxiliary program prints the same information.

The standard CDC library function EXP has a serious drawback: If the value supplied for exponentiation is less than or equal to -675.82, the result is set to zero, but no warning is given of this action. (A warning notice is printed if the value is too large.) A modified version of EXP which does give a warning message has been coded (in COMPASS) and this function is included in the <u>SURFACE</u> program.

Overlay calls to levels which are 8_{10} or greater are coded in octal numbers, using a terminal B (e.g., CALL OVERLAY (fid, 12B, 1)) to avoid the confusion inherent when the calls are decimal while the Overlay identification cards are octal. *

The Program Card for <u>SURFACE</u> includes as parameters the names of all files on which Overlays are stored, even those which are not released by DRØPIT (see Figure D-3). This listing on the Program Card was found to be necessary to prevent random losses of unlisted Overlay files when DRØPIT was used on an Overlay file that was listed.

D. 3 PROGRAM FUNCTIONS

a. Outline

This section contains descriptions of the various modules—overlays and subroutines—in <u>SURFACE</u>. These descriptions are given in the order of their presentation in Table D-1, which shows the Overlays which function for each major subtask.

The calling sequences for overlays and subroutines is outlined in Figure D-4.

b. First Principal Overlay (0,0)-Program TM42

There is an additional preliminary 0, 0 level Overlay, not shown in Table D-1, which can be used to set initial timing before even starting to copy the main program 0, 0 Overlay into core. Its use is optional. If it is used, it calls the main program automatically.

This module prints the data and a summary of the CP and PP time used since the job started, prior to starting on the <u>SURFACE</u> program proper.

Program TM42 calls Program P1942.

^{*}Note: In this Appendix, all overlays are referred to by their octal level numbers used in definine the overlay, e.g., QVERLAY (ABC, 11, 2) is referred to as (11, 2).

4 ,

Figure D-3. Program Card for SURFACE (P1942)

Table D-I
SURFACE PROGRAM OVERLAY STRUCTURE

Initiation and Program Ex	ecutive 0,0
Structure and Properties	Input 2, 1 1, 0 14, 0 2, 0
Materials Assignment and Surface Conditions	Initial 3,0 11,0-11,2
Precalculated Post Expos and Surface Conditions	4,0 7,0 11,0-11,2 10,2
Plume Exposure Sequence Surface Conditions (This sequence loops back on itself several times)	
Space Exposure Sequence Surface Conditions	and 6,0 7,0 11,0-11,2 10,1
Termination	16,0 10,0

c. Principal Overlay (0,0)

(1) Program P1942

This routine initializes most storage and control variables and flags. It reads in the data defining the exposure sequence to be followed and sets up the appropriate sequence of calls to primary overlays. These data are on file ADATA (TAPE16).

It is possible to turn off listing of some of the normal outputs which describe materials properties and satellite structure, and also the computations with precalculated exposure data. It is done by adding a single card to the 0,0 Overlay. This can be useful for reducing the amount of paper generated when several computer runs are to be made with the same initial data. The card assigns the value 7 to I6F. The usual location of use it is noted in the (0,0) listing. Reversion to normal listing at any point can be caused by inserting a card equating I6F to 6; in any case reversion occurs automatically upon entry to operating sequences which calculate plume or space exposure effects.

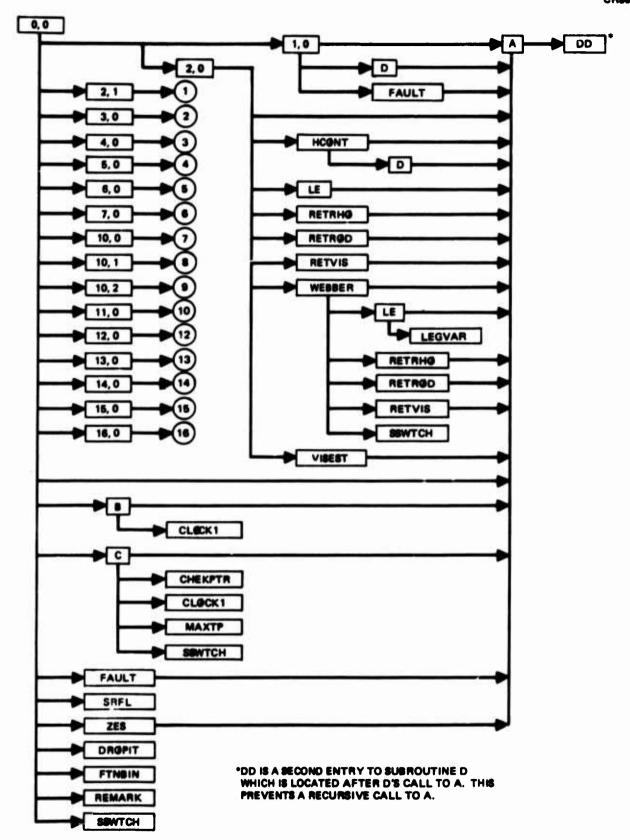


Figure D-4. Part A. Program Segment Linkage Pattern

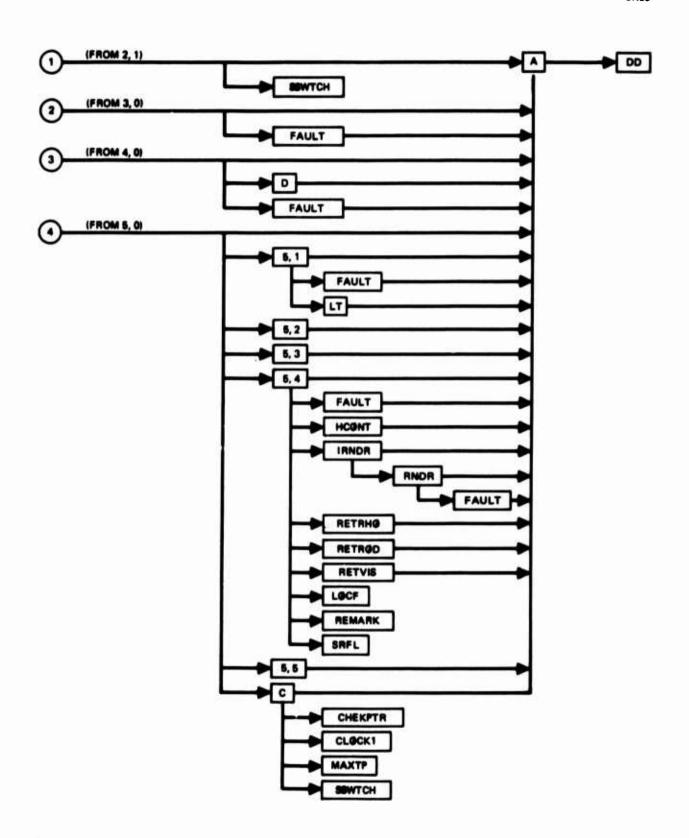


Figure D-4. Part B. Program Segment Linkage Pattern

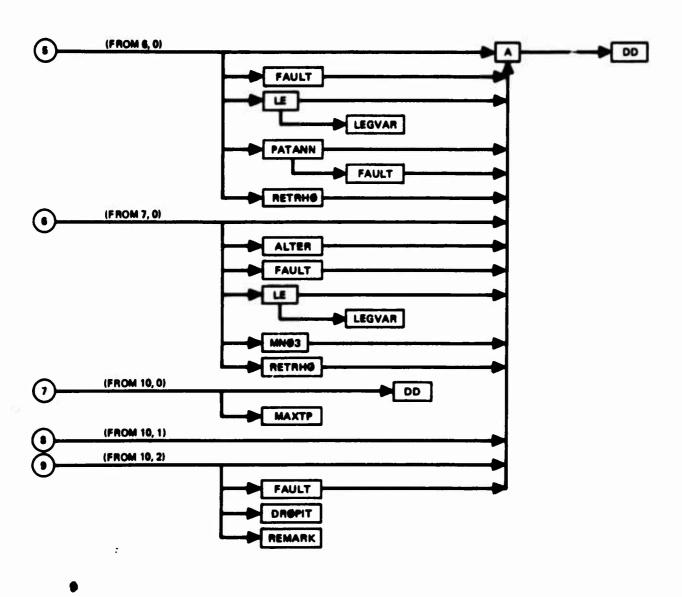


Figure D-4. Part C. Program Segment Linkage Pattern



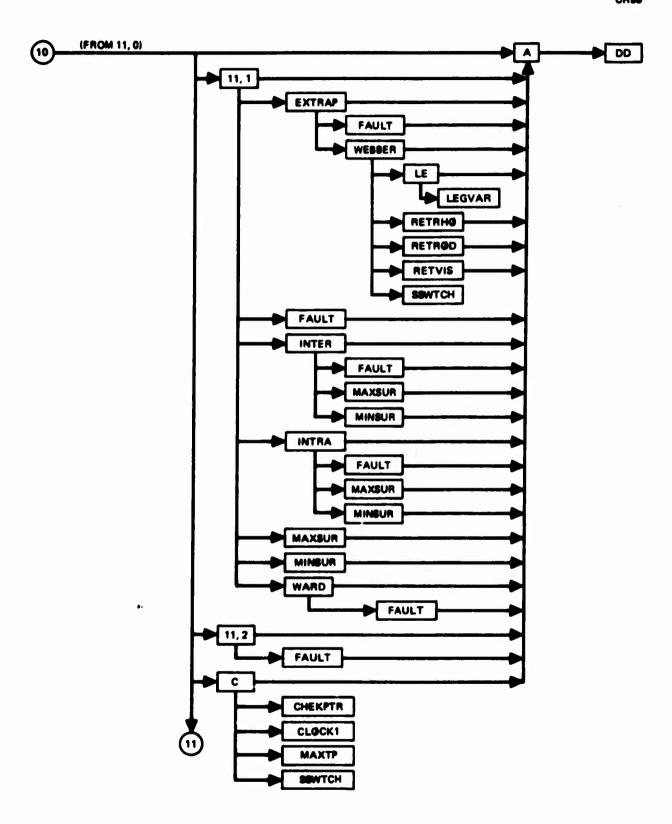


Figure D-4. Part D. Program Segment Linkage Pattern

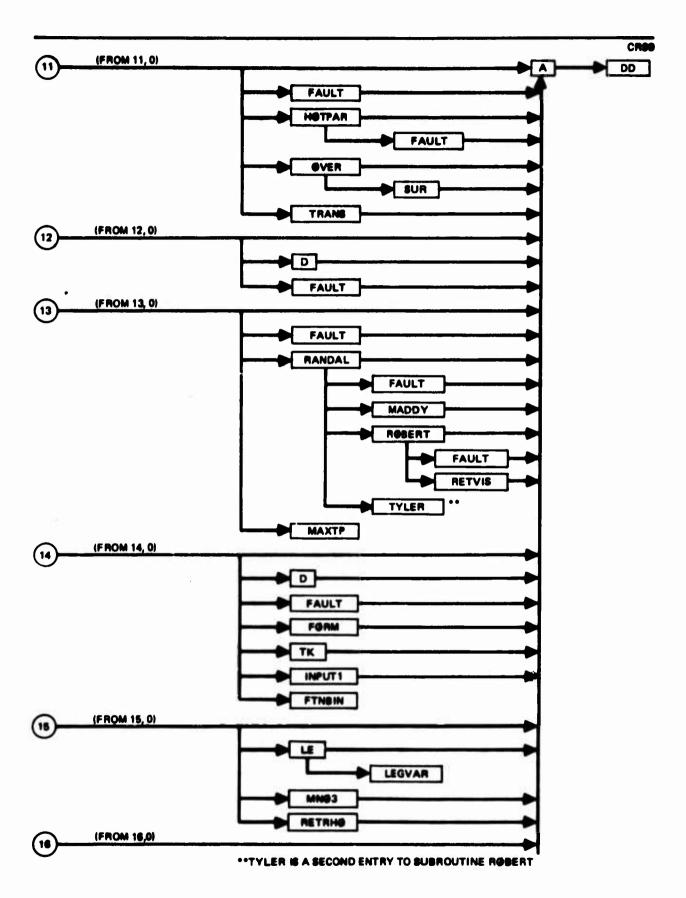


Figure D-4. Part E. Program Segment Linkage Pattern

Overlay (0,0) calls all primary overlays, the secondary overlays (2,1), (10,1), and (10,2), and the subroutines A, B, C, D, and ZES.

(2) Subroutine A

This subroutine writes the file (CQUNT = TAPE15) which records the number of calls to Overlays and Subroutines. It is called by all program units. It calls Subroutine D in case of a fatal error.

(3) Subroutine B

This subroutine initializes the clock for timing program operation. It calls Subroutine A.

(4) Subroutine C

This subroutine has two functions: (1) It keeps track of and prints the time needed to process each primary and secondary overlay and the total PP and CP time used on the job after finishing each overlay. (2) It will write a CHECKPØINT file of the <u>SURFACE</u> program if switched on to do so. The file is taken after return from the nthOverlay call, where n lies between 3 and 6, inclusive, and is repeated at every tenth subsequent return from an Overlay call. Each new CHECKPØINT file is written over the previous one. The CHECKPØINT file operation is enabled by turning on any Switch with a number from 3 to 6.

Subroutine C calls Subroutine A.

(5) Subroutine D

This subroutine prints a Dayfile listing of the numbered $ST \emptyset P's$ in the program and their corresponding sources. It calls Subroutine A.

(6) Subroutine ZES

This subroutine interprets the data which define the next exposure sequence and sets the flag which calls the appropriate computing sequence. It calls Subroutine A.

(7) Function HCØNT (TEMP)

This subroutine calculates the enthalpy HCØNT (= H) of the condensed phase of volatile plume constituents as a function of the temperature:

- (a) If T (= TEMP) is below the melting point of the material, then H = T * Cpsolid
- (b) If T is equal to or greater than the melting point, then H = Melting Point * C_{psolid} + H_{fusion} + (T-M. P.) * C_{pliquid}

The calculation is done using SI units.

Function HCONT calls Subroutines A and D.

(8) Function RETRHØ (RTEMP)

This subroutine approximates the SI reduced density of a liquid as a function of reduced temperature, RTEMP, along the saturation line, based on the curve of Hougen and Watson. The calculation is identical to REDRH \emptyset in \underline{TCC} .

Function RETRHO calls Subroutine A.

(9) Function RETRØD (RTEMP)

This subroutine calculates an approximation of the reduced density difference (liquid-vapor) as a function of reduced temperature along the saturation line. The calculation is identical to REDRØD used in TCC.

Function RETRØD calls Subroutine A.

(10) Function RETVIS (RTEMP)

This subroutine calculates the reduced viscosity (SI units) of a fluid as a function of reduced temperature. The value resulting is that for the liquid along the saturation line up to the critical temperature, and for the vapor along $P_{\rm O}$ above the critical temperature. The computation is identical to REDVIS in \underline{TCC} .

Function RETVIS calls Subroutine A.

(11) Function RNDR (X. N)

This subroutine will round off a real number, X, of any size to a real number containing N significant digits, without changing the location of the decimal point. The statement functions IRNDR, RNDI, IRNDI are used in conjunction with RNDR for input or output of integer variables.

Function RNDP calls Subroutine A.

(12) Subroutine FAULT (PØSN)

This subroutine is called when certain types of program-defined errors are detected. These errors may be fatal or nonfatal. For both types, an error message, giving the location of the error, is printed. The program is transferred to a termination step if the error is fatal. If it is nonfatal, the count of nonfatal errors is increased and the program returns to computation. However, if the error count reaches an input-data defined limit (defaulting to 10), the error is treated as fatal and computation is terminated. PØSN is the statement number (coded in Hollerith) at the location of the error. It is assigned in the calling routine immediately prior to the call to FAULT.

Subroutine FAULT calls Subroutine A.

(13) Subroutine LE (X, N)

This subroutine is used to check the validity of a real variable X (whether it is in range, indefinite, or infinite) before the variable is used in calculations, setting N to 0, -1 or +1 respectively. It calls the library routine LEGVAR for the check.

Subroutine LE calls Subroutine A.

(14) Subroutine VISEST (LQGV, N, X, Y, I)

This subroutine is used to calculate the viscosities of mixtures of liquids, given the viscosities X and Y of the component liquids. The number of concentrations calculated is given by I, which defaults to 10 if no value for I is supplied. The logical variable LQGV switches printing of the results off/on; if LQGV is .TRUE., the results are printed. When N is set to 1, a demonstration calculation using N_2Q_4 and MMH is carried out.

Subroutine VISEST calls Subroutine A.

(15) Subroutine WEBBER (RTEMP, X, Y, LQGV)

This subroutine is used to calculate the reduced values of a number of physical properties of liquid or gaseous plume/deposit constituents at the reduced temperature RTEMP. Either one or two substances (X and Y) can be handled at once. All the results are stored in the array REDPRP.

The routine checks to determine whether the particular temperature/constituent pair has already been computed before proceeding.

The logical variable LQGV switches on listing of the results when it is .TRUE..

The properties for which reduced property values are calculated are vapor pressure, density, density ratio if two substances are being treated, viscosity, and surface tension.

Vapor pressures is calculated by use of the Calingaert-Davis equation:

$$\ln P = A - B/(T - 43.)$$

This is applied by evaluating A and B at the boiling point of the fluid, then the A and B are used at the temperature of interest.

Density and viscosity are obtained by calls to the functions RETRHØ and RETVIS.

Subroutine WEBBER also calls Subroutines A, LE, and RETROD

d. Primary Øverlay (1,0) - Program P4210

This routine reads and stores input data which have the following names and functions:

FACTR - controls width of time slice on space exposure

DECPTH - significant abrasion depth lower limit

DEDPTH - significant deposition depth lower limit

NAME - symbolic names of structural features, structural materials, and plume/deposit constituents

(7) The parachor, PCHØR, is calculated by means of the relation

PCHØR = (Molecular Weight * Surface Tension)^{1/4}
Critical Density * RETRØD (Reduced Room Temperature)

= (AMW * TENS) ** 0.25/(RCRIT * RETRØD (TRED))

- (8) The liquid enthalpy, vapor pressure, specific gravity, viscosity, and surface tension are calculated at a series of temperatures for $N_2 \not O A_1$, MMH, and water, using the equations and subroutines describer earlier, and the results are printed.
- (9) The latent heat content of the vapor, DELH, at room temperature is calculated by means of the relationship.

DELH = specific heat (vapor) * room temperature * enthalpy at absolute zero - (specific heat (solid)) * freezing point + latent heat of freezing + specific heat (liquid) * (room temperature-freezing point)

Subroutines are used so the equation is

DELH = HVAP (TEMP (ATM)) - HCØNT (TEM).

- (10) A call to Subroutine WEBBER is made, and the density, viscosity, and surface tension are calculated at assigned temperatures, read from file ADATA (TAPE16).
- (11) A call to Subroutine VISEST is used for calculating the viscosity of mixtures of liquids from the viscosities of the component liquids.

Program P4220 calls Subroutines A, HCØNT, RETRHØ, RETRØD, RETVIS, LE, WEBBER, and VISEST.

g. Secondary Overlay (2, 1) - Program P4221

This overlay is used to read in NAMELIST values of changed data from the INPUT file (TAPE17) and stores them on TAPE18 for access by user routines. It is activated by turning on SWITCH 1 by means of the SCØPE control card.

SWITCH, 1.

If desired, it will also list the values supplied. This is controlled by the value of the flag RDWRT—if RDWRT is greater than zero, the input is printed.

There are eight NAMELIST sets, which include the following variables.

NAST1 — Used by ϕ VERLAY (0, 0) for changing exposure cycle assignments:

- 1. NPI number of sequences to be run
- 2. NPSEQ sequence of exposure cycles
- 3. ISXTST number of space exposing
- 4. IPXTST number of plume exposures
- 5. IGØTST total number of exposure cycles

NAST2 — Used by ØVERLAY (4,0) for changing assigned effect inputs:

- 1. IIS total number of segments affected
- 2. EFFECT conditions on segments

NAST3 — Used by \emptyset VERLAY (5, 0) for changing values for the variables:

- 1. TØTIME plume operating time
- 2. DELTIME time slice width for plume exposure calculations

NAST4—Used by ØVERLAY (5, 0) for changes in assignments to the variables:

- 1. P quarter latus-rectum of the plume
- 2. DELP change of P from cycle to cycle
- 3. NTR I.D. number of active thrustor (source of plume)

NAST5 — Used by \emptyset VERLAY (10, 2) for checking whether the assigned data used in \emptyset VERLAY (4,0) are to be the basis for further calculations instead of using the initial conditions:

ISTST - flag to select which data for the base

NAST6 and NAST7 – Used by \emptyset VERLAY (12, 0) for input of new or additional plume characteristics data in addition to card input and data supplied by $\underline{KINC}\emptyset N$:

- 1. IPIS identifies cycle number
- 2. IP2S identifies segment to which initially input plume characteristic data apply
- 3. J1 new cycle number
- 4. K1 new segment
- 5. PLMCHR new plume characteristics

NAST8—Used by ØVERLAY (11, 0) to control evaluation of averaged thermal transfer coefficients.

- 1. PRJFLG flag to turn on processing of averaged thermal coefficients on a projection
- 2. MS11 SURDES location of segment with lowest ID number to be included
- 3. MS21 SURDES location of highest segment ID to be included

Program P4221 calls Subroutines A and D.

h. Primary Overlay (3,0) — Program P4230

This overlay reads in the assignment of structural materials and initial surface roughness and temperature for specified segments of the satellite surface and projections from file BDATA (TAPE8). The data are added to the array SURDES which already contains the coordinates of each segment. The data are also placed in the transfer array SEEK, which is used to search for the properties (optical and thermal) which the surface exhibits. SEEK is buffered out onto TAPE12.

No computations are conducted in P4230, other than to check the ID's and locations of individual segments.

Program P4230 calls Subroutines A and FAULT.

i. Primary Overlay (11,0)

The set of overlays and subroutines based on Primary Overlay (11,0) function to calculate the current state of each segment in terms of optical and thermal energy transmission coefficients, based on the current physical and chemical condition of the surface of the segment, resulting from initial preparation or plume or space exposure.

(1) Program P42110

The program first reads in from TAPE12 the latest values for SURDES (surface description), SEEK (the segments affected by the latest sequence of exposures), and several constants and auxiliary arrays. A check is made that the data in SURDES are consistent, and that a top layer still exists for each segment. If the top layer has been destroyed, (e.g., by abrasion), the second layer is moved into the top position, etc. The calculations of the state of each segment are made by table interpolation during multiple calls to the secondary Overlays (11, 1) and (11, 2) and associated Subroutines, and the numerical results are returned in the array VALUE. A final check is made to ensure that segment ID's match in the arrays SURDES, SEEK, and VALUE, then the data are entered into SURDES.

The temperature of the surface is checked, and the emissivity is corrected for temperature.

The data are then printed out in a large tabulation in which each segment is described completely—location, deposit thickness, materials, surface condition, optical and thermal constants. In cases of slight abrasion, where the thickness might require printing a string of 5 to 19 nine's or else rounding up to no apparent effect, a special routine prints the value in the form:

n· n9(6)9nn

where the bracketed 6 indicates 6 additional nines in between the two printed nines. Subroutines ØVER and SUR perform this task, storing the results in arrays SURH and SURI.

Subsequent to printing the data for each segment, the average values over the whole satellite for absorptivity and emissivity are calculated by summing and dividing by the number of segments, and listed. The calculation is done in subroutine HØTPAR. The results are EFALFA for the average solar absorptivity, EFEPSN for the average thermal emissivity, and EFRATØ for their ratio. The results are printed. The percent change in EFRATØ from the baseline value is calculated and printed.

The program next checks TAPE18 for the presence of NAME-LIST NAST8 directives which cause a calculation of the average heat transfer coefficients for projections. If the directive is present (PRJFLG > 0) the program recalls HØTPAR and calculates EFALFA, EFEPSN, and EFRATØ for specified projection locations. This call can also be used for calculating the constants for specified limited regions of the hull of the satellite.

Next, all transparent segments are isolated, and the percent transmission for normally incident rays of light through any deposits and the transparent structure elements are calculated at seven wavelengths, from far infrared to ultraviolet. When there are several segments of one transmission type (such as a solar cell array) the average transmission for the group of segments for each wavelength is calculated and printed. These calculations are done in the Subroutine TRANS.

All subroutines local to (11,0) and its secondaries are attached to (11,0). These include EXTRAP, H ϕ TPAR, INTER, INTRA, MAXSUR, MINSUR, ϕ VER, SUR, TRANS, and WARD. They are described following the sections on (11,1) and (11,2).

Overlays and Subroutines called from P42110: P42111 (ϕ VERLAY (11,1)), P42112 (ϕ VERLAY (11,2)); A, C, ϕ VER, FAULT, H ϕ TPAR, and TRANS.

(2) Secondary Overlay (11, 1) - Program P42111

This program initiates the search for property data which correspond to the current surface condition. The array VALUE is used to hold the results of the search. The array ALPHA is used to hold data originally read in as PRØPTY, CHEMIC or MATERAL, now stored on files, and read in as required for current searching needs. The data contained in ALPHA become the primary source of information on the surface state.

The data in array SEEK are first checked to be sure that the layers of materials correspond to those in SURDES, except of course, for new deposits or removal by abrasion/evaporation. Subroutine WARD makes this test, and performs simple corrections.

Then, working with one segment at a time, and on each segment from the top layer down, one layer at a time, the material present is identified by its code number. If it is a deposit that is present, the substance present in highest concentration is taken to represent the deposit. Deposit temperature is obtained by octal masking of the appropriate location in the array SEEK.

A check is made to be certain that the surface condition sought is legal (not a negative number). If illegal, a warning message is printed and the segment is bypassed.

If the condition is legal, the file TAPE2 (which contains optical/thermal properties) is searched until the material corresponding to that on the segment is found. The data for this material are copied to array ALPHA. A flag called $H\emptyset LD$ is set with the material's code number: this will be checked when the next segment is processed, and if the material and $H\emptyset LD$ are the same, the file search will be skipped.

The surface condition of the segment is compared with the entries in ALPHA. If a match is found, the properties corresponding to the condition are copied to array VALUE. Where a specific property value is not available for the surface condition of interest, interpolation/extrapolation from values for other surface conditions must be carried out—Subroutines MINSUR, MAXSUR, INTER, INTRA, and EXTRAP are used for such calculations—the specific routines called depend on the type of calculation needed.

In similar fashion, if the surface condition does not correspond exactly to an entry in ALPHA, extrapolation/interpolation is used to identify its exact relation to the entries in ALPHA, then the surface state data are calculated. After completion of the calculations, the results are entered into the array VALUE.

The calculation recycles until data for all layers of all segments have been entered into VALUE, then a return to ØVERLAY (11,0) is made.

PROGRAM P42111 calls Subroutines A, WARD, MINSUR, INTER, MAXSUR, EXTRAP, INTRA, and FAULT.

(3) Secondary Overlay (11, 2) - Program P42112

This program enters the data on surface state from VALUE into SURDES. It first checks the information currently in SURDES against the entries in VALUE, and if they are the same, no further action is taken. A check is then made of whether the sequence of materials in VALUE is possible (e.g., that no cases exist where the top layer is aluminum and the second is a deposit). Such errors will abort the program. Then the sequence of layers in VALUE is compared with the sequence in SURDES. After allowing for deposition and erosion, if the sequences do not correspond, the data in VALUE are bypassed and the next segment is taken up.

If deposits exist on the segment in both SURDES and VALUE, their identities are compared. They should be the same, both plume and space exposure routines call Overlays (15,0) and (7,0) to make sure of this. However, post exposure data supplied as input data in Overlay (4,0) does not call those overlays, so in that case the concordance may not exist. If such is the case, some simplified assumptions about the chemistry of deposit interactions are made, and the final products calculated. If this process has occurred, the data on surface state in VALUE are incorrect, and the routine returns to (11,0) with flag GETGET set to cause another call to (11,1).

Eventually the material identity test is satisfied.

Then (11,2) transfers the data in VALUE to the array SURDES and returns.

Program P42112 calls Subroutines A and FAULT.

(4) Subroutine EXTRAP

This subroutine, called from (11,1), is used to interpolate/extrapolate physical/chemical properties as a function of temperature required to determine the surface state. It reads TAPE3 until it finds the material ID number, then puts the data into the array CHEM, then calls Subroutine WEBBER to do the actual computations. The results are returned in the array REDPRP.

Subroutine EXTRAP calls Subroutines A, FAULT, and WEBBER.

(5) Subroutine HØTPAR

This subroutine is called by the Overlay (11,0). It calculates the overall effective values for α , ϵ , and the α/ϵ ratio corresponding to a particular set of surface conditions. Local values of α , ϵ , and the area of each segment are read from the file SURDES. The products of α times area and of ϵ times area for each segment are separately summed, then divided by the total area. Normally, the complete main body of the satellite is used for the calculations, and projections such as solar cell arrays are ignored. However, by using Namelist input NAST8, the routine can be commanded to calculate the coefficients for any subsection of the satellite, including and/or limited to projections.

Subroutine HØTPAR calls Subroutines A and FAULT.

(6) Subroutine INTER

This subroutine, called from Overlay (11, 1), is used to interpolate values for surface optical/thermal properties when no values corresponding to the surface finish of interest are available in the tabulation. It searches to find the closest lower and higher values, and the corresponding finishes. A factor is calculated from the finishes

$$Q = \frac{R_{\emptyset} - R_{L}}{R_{U} - R_{L}}$$

The surface property required is then calculated.

$$S_{\emptyset} = (S_{U} - S_{L})*Q + S_{L}$$

where S indicates the surface property, R the finish, L the lower value, and U the upper value.

Subroutine INTER calls Subroutines A, FAULT, MINSUR, and MAXSUR.

(7) Subroutine INTRA

This subroutine, called from overlay (11, 1), is used to interpolate values for surface optical/thermal properties when interpolation is necessary on the surface finish entry as well as the desired property entry. The interpolation routines are the same as in INTER.

Subroutine INTRA calls Subroutines A, FAULT, MINSUR, and MAXSUR.

(8) Subroutine MAXSUR

This subroutine, called from Overlay (11, 1) or by Subroutines INTER and INTRA in turn called from (11, 1), is used when there is no value of a selected surface thermal/optical property corresponding to the surface finish of interest. It searches through the tabulation to find the property entry which corresponds to a surface finish which is less than but as close as possible to the surface finish of interest.

Subroutine MAXSUR calls Subroutine A.

(9) Subroutine MINSUR

This subroutine, called from Overlay (11, 1), INTER and INTRA, is the mirror image of MAXSUR. It is used to find the value of a surface property corresponding to the least increase in surface finish, when a value corresponding exactly to the surface finish is not available.

Subroutine MINSUR calls Subroutine A.

(10) Subroutine ØVER

This subroutine, called from Overlay (11,0), is used to display a long series of the numeral 9 in condensed form, avoiding the requirement for a printing format of 20 positions to prevent rounding up. This is used for printing the thickness of materials after exposure to the plume, where a minor amount of abrasion can give a long series of 9's. The routine encodes the number in decimal form, then decodes it as a one-dimensional array of individual numbers. It then inspects each number in turn, starting at the most significant. It counts the number of 9's in any sequence of 9's only, and if the number of 9's is greater than 5, it sets up the special format. Any number other than 9 in the sequence counted resets the counter to zero.

The special format consists of modification, suitable for computer printout, of the number structure found in some mathematics publications,

$$x \cdot x \cdot 9_n \cdot 9 \cdot x \cdot x$$

where the subscript in n is a number which indicates how many nines lie between the two nines printed on the line (that is, there are n + 2 nines in the sequence). The x's indicate any other numeral. In SURFACE, the above structure is presented as

$$x \cdot x 9 (n) 9 x x$$

The procedure works only for real numbers — extension to exponential numbers (scientific notation) would not be difficult but does not seem to be needed.

Subroutine ØVER calls Subroutines A and SUR.

(11) Subroutine SUR

This subroutine, called from ØVER, stores the number describing the thickness of a layer in one of two arrays, SURH or SURI, for transmission to Overlay (11,0). The array is selected by means of a flag which is set by the use to which the number will be put.

Subroutine SUR calls Subroutine A.

(12) Subroutine TRANS

The subroutine, called from Overlay (11,0), calculates the percent transmission of light (UV, visible, and IR) through all layers of transparent locations on the satellite—sensors, solar cell arrays, and windows. The approach is based on the usual relation for intensity at any depth;

$$\frac{I}{I_o} = e^{-kt}$$

which is applied to each layer, including deposits, and on extra loss of intensity at all interfaces. The loss at each interface is calculated from the reflectivity appropriate to the wavelength—if reflectivity data are not supplied, it is taken as 4 percent for internal interfaces and 2 percent for external interfaces.

The results are returned to (11,0) for display printing, but are not stored.

Subroutine TRANS calls Subroutine A.

(13) Subroutine WARD

This subroutine, called from Overlay (11, 1), is used to transfer all lower layers upward one layer in the segment description if the data indicate a zero or negative thickness for a top layer in the array SEEK. It checks the results for consistency with the data in SURDES.

Subroutine WARD calls Subroutine A and FAULT.

j. Primary Øverlay (4,0)—Program P4240

This overlay has the function of accepting input data which define the postexposure condition of the exterior of the satellite when this data originates from sources other than program <u>SURFACE</u>. Such data may be input simply to test the overall program or may have been generated from other sources, such as tests, or even from previous runs of <u>SURFACE</u>. The data are input from file CDATA (TAPE9).

Prior to input of the data, the initial state of the satellite surface as contained in SURDES is written out and stored in TAPE4. This feature allows use of the defined postexposure state simply to test the program, and then later to calculate exposure effects on a pristine surface. If this is done, the initial state is restored downstream by a call in Overlay (10, 2).

The data read in by Overlay (4,0) are stored directly in the array EFFECT, which is same array for storing the results from plume or space exposure. The input data are listed.

Program P4240 calls Subroutines A, D, and FAULT, and calls in TAPE12 to obtain the current values in SURDES for writing on TAPE4.

As mentioned in the description of the main Overlay (0,0), the call to Overlay (4,0) can be bypassed if assigned condition calculations are not wanted.

k. Primary Øverlay (7,0)

(1) Program P4270

This program calculates the final chemical composition of deposits on satellite segments after a plume or space exposure cycle, and writes the results in appropriate locations of the arrays SURDES and EFFECT. The computations include calculation of chemical reactions between pre-existing and newly deposited layers.

There are a number of steps in the calculations in this program.

- (a) This array EFFECT is checked for the presence of data with which to calculate.
- (b) Local, temporary arrays are copied from SURDES and EFFECT.
 - (c) The presence of deposits is tested.

The following steps are carried out for each affected segment before returning to the main overlay.

- (d) If there are no deposits, appropriate changes in SURDES are made in response to data in EFFECT and program returns to Overlay (0,0).
- (e) If there are deposits, a check is made for their presence both prior to and after the exposure.
- (f) If deposits are present only prior to the exposure, the exposure has removed them. The appropriate changes in SURDES are entered and the program returns to (0,0).
- (g) If deposits are present only after exposure, exposure has caused their formation. The layer structure is checked for consistency, then SURDES is modified and the program returns to (0,0).
- (h) If deposits are present both before and after exposure, their constituents and percentage composition are equated to a set of local variables.
- (i) The segment temperature is noted, the melting point of all deposit constituent and density-temperature coefficients are read in from TAPE 3.
- (j) The surface temperature is compared with the melting points of the deposit constituents. If all constituents are solids, no reactions will occur, and the following calculations are made.
- 1. The mass of each constituent is determined from the percent concentrations in the new and old deposits.
- 2. The masses of species present in both old and new layers are summed.
- 3. The total mass and the percent concentrations of all species are calculated.
- 4. The two most abundant species are identified (plus water, if present).
- 5. The identities and concentrations of the two most abundant species, plus the concentration of water, are stored in SURDES and EFFECT, and the program returns to (0,0).
- (k) If at least one constituent is liquid, chemical reactions are possible. The species present are checked to see if there are any mutually reactive substances.

Allowed reactions are

$$N_2 \phi_4$$
 + any base - base nitrate
 $HN\phi_3$ + any base - base nitrate
 $HCL/HCL \cdot aq + NH_3/NH_4 \phi H$ - ammonium chloride
 $N_2 \phi_4$ + water - nitric acid

Displacement reactions and redox reactions are not allowed.

- (1) If no reactions can occur, the calculations summarized in (j) 1 to 5 are carried out.
- (m) If reactions are possible, the following calculations are made:
 - 1. The mass of each constituent present is determined.
- 2. Identical species are identified and their masses are summed.
- 3. Using a table of allowed reactions and their priorities, the following calculations are made:

The equivalent concentrations of the reactive species are calculated.

It is assumed that the reaction goes to completion. The final concentration of the reactant in excess is calculated.

The product is identified and its concentration is calculated.

- (n) The remaining reactants are checked in a return to steps (k) through (m).
- (o) After all reaction calculations have been completed, the program steps to the routines described under (j).
- (p) The information on deposit species and concentrations is stored in condensed fashion by octal masking to reduce array size requirements.

Program P4270 calls Subroutines A, FAULT, REPLACE, LE, RETRHØ and MNØ3.

(2) Subroutine MNØ3

This subroutine checks whether more than one type of base nitrate is present, and if such are detected, they are combined as the mixed fuel nitrate deposit species.

Subroutine MNØ3 calls Subroutine A.

(3) Subroutine REPLACE

This subroutine is used to move the structural and deposit layers in array SURDES upward or downward when this is required because of removal of a layer or formation of a new deposit.

Subroutine REPLACE calls Subroutine A.

1. Overlays (11,0) to (11,2)

Overlays (11,0) to 11,2) are called at this time. See their descriptions in Section D. 3. i.

m. Secondary Overlay (10, 2)-Program P42102

This program checks whether the post exposure conditions input in Program P4240 are to be used as the base condition prior to further exposure calculations, or that they should be discarded and the initial conditions reinstated. The flag ISTST is used. This is read from TAPE9(=CDATA), and then NAMELIST NAST5 is checked for any changes. If ISTST iz zero, the initial conditions are to be used, and they are brought back in from TAPE4, where they were stored during operation of P4240. If ISTST is greater than zero, the data on TAPE4 are ignored and the program returns to (0,0).

Program P42102 calls Subroutine A.

n. Primary Overlay (5,0)

The sequence of routines called in this region of the program calculate the effects of plume exposure on the satellite exterior. The sequence includes calls to Overlays (7, 0), and (11, 0) to (11, 2), all previously described.

The primary and secondary level 5 overlays will be described first, then the associated subroutines. Following these will be other primary overlays which are used.

(1) Program P4250

This overlay controls the sequence of calculations for plume exposure effects and performs basic calculations related to the timing of the events. The program expects to find information about the plume, supplied by TCC or KINCON on TAPE30. It first checks for the thrustor "on" time, TOTIME, the selected time increment, DELTIME, and the ID number of the

thrustor which is operating, NTR. If these values are not found on TAPE30, the data input in Overlay (0,0) supplies TØTIME, and TAPE10(=DDATA) is read for DELTME. Then, NAMELIST NAST8 is checked for updated values of TØTIME and DELTME. If no value for TØTIME is supplied, the run is terminated after printing an appropriate message. If DELTME is not supplied, a default value is used.

The number of calculation cycles, NUMCYC, is determined by the relation

NUMCYC = TØTIME/DELTME.

Preliminaries being completed, the clock is started:

TIME = TIME + DELTME

where TIME was previously equal to zero.

Data on the plume configuration is now required from KINCON.

*The first data needed are the latus-rectum of the plume, assuming it has a paraboloid shape. A tape is made upon which is listed each time interval and a zero value of P, which is one-fourth the latus rectum at that time. This is written out on TAPE 31.

*The program later reads TAPE31, expecting to find nonzero values for P. If P is found equal to zero, and there was no previous P, the program reads DDATA for a value of P.

If a previous value for P exists which was not supplied by TAPE31, DDATA is read to find DELP, which is the change in P per time increment, unless DELP has already been input.

*A check is also made to be certain that a value for NTR was read in from TAPE30 earlier. If the value of NTR is zero, a value is read from DDATA (TAPE10).

If the calculation is in the first exposure cycle it is assumed that the available value of P corresponds to the current plume shape. When DELP has been input, implying that SURFACE must calculate the plume shape, the change in P for each subsequent cycle is calculated at the end of the calculations for the current cycle, as described later.

The thrustor location is read from storage, using the thrustor ID number, NTR, as the key.

The next step is to determine which segments of the satellite surface are impinged by the plume, knowing the thrustor location and plume configuration. Program P4251 (Overlay (5, 1)) is called in to do this calculation, and the results are stored in the Array AFFSEG (affected segments) and listed in the Output. The program will actually accept several inputs of P and NTR and calculate and list the plume-surface intercepts. If this feature is used, the program assumes that the last such input is the one to be used for calculation of plume effects.

Because KINCON and upstream subprograms use a coordinate system based on the thrustor, and do not relate to the satellite on which the thrustor is mounted, routines to transform coordinates from thrustor-based systems to satellite-based systems and vice versa are needed. These are available in Overlays (5,2) and (5,3). Because the transformation requirements may vary from run to run, there is a flag, ITRNCL, which selects the needed routines. If ITRNCL is negative, only satellite-based to thrustor-based are computed; if ITRNCL is zero, none are computed; if ITRNCL = 1, both types will be run, and a value of 2 limits the calculations to thrustor-based to satellite-based. The value of ITRNCL is tested; if zero, a new value is sought on TAPE10. Note that input of a 'new' value of zero is allowed.

For demonstration or testing purposes of thrustor-to-satellite based coordinates, TAPE10 is read to provide a thrustor ID number and an index equal to number of locations which will be transformed in the demonstration.

Then the thrustor based coordinates are input from TAPE10, and Overlay (5,2) = Program P4252 is called to calculate the transformations. On return, the results are listed. Then a check is made for further requirements for these transformations. When completed the program moves to the inverse calculation, if it is required.

Here, the data in AFFSEG on actual plume-satellite intercepts are used for the calculation, done by calling in Overlay (5, 3) = Program P4253. Again, the results are listed.

At this point, NAMELIST NAST4 is checked for changed value assignments for P, DELP and NTR.

*When the above sequences are completed, it a set of P values at different times are available from \underline{KINCON} , as discussed earlier, then TAPE31 is read at each time interval to find P, and Overlays (5, 1) and (5, 3) are called, and the thrustor based coordinates for all affected segments are determined at each time interval, and immediately written out on TAPE32. This procedure is done until all the affected segment locations for time slices have been written to the file.

*The file Tape 32 is actually an enquiry sent to \underline{KINCON} : the program will expect to get data on the plume characteristics for each location and time interval on TAPE14 when overlay (12,0) is processed downstream.

The program then checks SURDES to determine whether the surface temperature has changed since the last time cycle, and calculates a new average temperature. Those material properties for structural and previous deposits which affect further plume-surface interactions are identified, and the current values, as determined by the surface temperature, are calculated in Overlays (5, 4), (5, 5), and the final steps in this primary overlay. Representative listings are made and then the program returns to the principal (0,0) Overlay.

Subprogram SURFACE, because of its core size requirements, has been modified to run independently of the remainder of \underline{CONTAM} , instead of in conjunction with the other subprograms under the control of

the CONTAM executive. All portions of SURFACE which were designed to communicate directly with other CONTAM subprograms have been rendered inoperative. They are still present in the program, but they have been changed to COMMENT cards. These inactive portions can be found at the following locations:

Overlay (5,0) - Program P4250

Cards 250 to 261 Card 268 Cards 302 to 326 Card 349 Cards 571 to 582

Overlay (12,0) - Program P42120

Cards 207 to 219

Overlay (13,0) - Program P42120

Cards 147 to 159

The descriptive paragraphs above which have been marked with an asterisk do not apply when this modification is in effect.

Because direct automatic communication between <u>SURFACE</u> and other subprograms of <u>CØNTAM</u> is now impossible, the execution of <u>SURFACE</u> has been modified to facilitate indirect, manual communication. This has been done by setting up three modes of operation, with the mode controlled by data in the physical punched card file INPUT (TAPE17) of the control flag STPFLG (an integer).

If STPFLG = 0, the program runs as a complete unit. Default data on plume characteristics for calculating impingement effects is read from input file EDATA (TAPEII). Note that if the currently deactivated interprogram communication routines are reactivated, plume characteristics data from other subprograms will be read from file TAPEI4, and EDATA will be ignored.

If STPFLG = 1, the program runs only as far as calculation of the plume- a ellite intercepts in Overlay (5,0). At this point, all data in Common and variable storage is copied to file TAPE30 and TAPE30 is confirmed, the coordinates of the intercept locations are calculated in terms of the thrustor based coordinate system and listed. Then the program stops. Note that TAPE30 must have been established as a RESERVE physical tape or a common or permanent disc file because the data stored on it will be required in the third mode of operation.

The listed output data on the coordinates of plume-surface intercepts are used to determine the characteristics of the plume at these locations by some manual method of interrogating the other program in CØNTAM. These characteristics are then coded into punched cards in the format described for

file EDATA (see Section D. 4. n) and this punched card file is input to replace the default file which is currently supplied with the program. If all data inputs being used in the form of physical cards, the EDATA decks are exchanged. If the data input are card images on magnetic tape or disc, the file corresponding to default EDATA is not copied from the master file (SKIPF, fid, 1, 17, B. is is inserted in the control card file where appropriate) and the new card deck is copied to disc by use of the control card CQPYBR, INPUT, EDATA.

If STPFLG = 2, <u>SURFACE</u> reads the stored values of Common and variable storage from file TAPE30, then resumes execution in Overlay (5,0) at a location immediately following the location of the STØP in the second mode. When plume characteristics information is required for calculations (Overlays (12,0) and (13,0)), the data will be sought on file EDATA; presumably the program will find the data prepared manually as described above.

Program P4250 calls Overlays (5,1), (5,2), (5,3), (5,4), and (5,5) and subroutines A, FAULT, and C.

(2) Secondary Overlay (5.1)-Program P4251

This program calculates the intercepts of the thrustor plume with the surface of the satellite and with projections from the surface as a function of the plume configuration. The location of the center point of each segment is the reference point for the calculation; if the center is on or inside the plume boundary, the whole segment is assumed to be affected by the plume.

The method is based on the calculation of the intercepts of two circles, the planar cross sections of the satellite and plume. Each segment is tested to see if it is "inside" the boundaries of the plume by the following sequence of steps:

- 1. The segment x coordinate is checked to ascertain that it is aft of the thrustor.
- 2. The radius of the plume is calculated at a position corresponding to the x coordinate of the segment:

$$R = \sqrt{2 p (x - a)}$$

where a is the x coordinate of the plume vertex.

$$AR = SQRT (2. * P * (SR2 - TX))$$

- 3. The plume radius is checked against the thrustor standoff and the local satellite radius to see if the plume contacts the satellite surface.
- 4. The limits of plume contact with the satellite on either side of the line joining the satellite and plume central

axes are calculated as angular displacements from that line.

Plume contact limit angle = Cos-1 ((Thrustor offset)² + Satellite Radius)² - (plume radius)²/2(Thrustor offset) (Satellite Radius))

$$ALP = ACØS (TR**2) + (SR3**2) - (AR**2))/2.*TR*SR3)$$

- 5. If the diameter of the plume is so large that it impinges beyond the midline of the satellite, the impingement is limited to the midline.
- 6. The 0 coordinate of the segment is tested to see if it lies in the range defined by the 0 coordinate of the thrustor plus/minus the plume contact limit angle.
- 7. If the segment meets the above tests, its ID is entered into the array AFFSEG (affected segment).

An analogous computation is used to find projection segments impinged by the plume; the intercept calculated is that of a paraboloid with a plane at a known angle to the axis of the paraboloid.

Program P4251 calls Subroutines A, FAULT, and LT.

(3) Secondary Overlay (5,2)-Program P4252

This routine, called from \emptyset verlay (5,0), transforms the coordinates of any point reported in cylindrical coordinates based on a specified thrustor to a cylindrical coordinate system based on the satellite.

If X_S, R_S, and θ_S are the required satellite-based coordinates of the point, and Z_T, R_T, and θ_T are the known thrustor-based coordinates of the point, with the thrustor coordinate Z parallel in 3-space to the satellite X, and if the satellite-based coordinates of the origin of the thrustor-based system are X_O, R_O, and θ_O , then the transformation equations are:

$$X_{S} = X_{o} + Z_{T}$$

$$R_{S} = \sqrt{\left(2\left(R_{o}^{2} + R_{T}^{2} + 2R_{o}R_{T}\cos(\theta_{o} - \theta_{T})\right)\right)}$$

$$\theta_{S} = Tan^{-1} \frac{R_{o}\sin\theta_{o} + R_{T}\sin\theta_{T}}{R_{o}\cos\theta_{o} + R_{T}\cos\theta_{T}}$$

Program P4252 calls Subroutine A.

(4) Secondary Overlay (5, 3) = Program P4253

This routine is the converse of P4252, and calculates thrustor-based coordinates of a point from the satellite-based coordinates.

If X_S , R_S and θ_S apply to the satellite system

 X_0 , R_0 and θ_0 are the origin of the thrustor in the satellite system

 Z_{T} , R_{T} and θ_{T} are the required thrustor-based coordinates

$$Z_T = X_S - X_o$$

$$R_T = (R_S^2 + R_o^2 - 2 R_S R_o Cos(\theta_S - \theta_o))$$

$$\theta_{\rm T} = {\rm Tan}^{-1} \frac{{\rm R}_{\rm S} \sin \theta_{\rm S} - {\rm R}_{\rm o} \sin \theta_{\rm o}}{{\rm R}_{\rm S} \cos \theta_{\rm S} - {\rm R}_{\rm o} \cos \theta_{\rm o}}$$

The program uses data in AFFSEG for impinged segments as the source of X_S , R_S , and θ_S .

Program P4253 calls Subroutine A.

(5) Secondary Overlay (5,4) = Program P4254

This program supplies physical and mechanical properties of structural and deposited materials which are needed to calculate the effects of impingement by the plume. The properties are corrected to the current surface temperature if the temperature factors have been input.

The methods for calculation deposit properties are those discussed earlier in Overlay (2,0). The properties of interest for deposits are the enthalpy of the deposit and of the vapor, the enthalpy of deposition, the specific gravity, the viscosity, and the surface tension.

The materials properties at the specific temperature are evaluated by using the temperature coefficients for the equation:

Property Value (T) = A + BT + CT² +
$$\frac{D}{T}$$
 + $\frac{E}{T^2}$

The coefficients have been entered in the input array MATERAL and stored on TAPE1. The properties of interest are the melting point, Vickers hardness, bulk modulus, surface energy, heat capacity, and thermal conductivity.

If temperature coefficients are not available, room-temperature data are used.

The properties data are stored in the array SEARCHD and buffered out on TAPE19.

Program P4254 calls Subroutines A, FAULT, IRNDR, RETRØD, RETVIS, RETRHØ, and HCØNT.

(6) Secondary Overlay (5, 5) = Program P4255

This program changes the values of the constants in the abrasion rate in response to temperature changes on the surface.

A temperature factor is calculated by the equation

Factor = 1-0.25 (Surface Temperature-520)/520

This factor is also applied as follows:

Upper Velocity Limit_T = Lower velocity limit_o x (Factor) 1.2

Hyper Velocity Limit_T = Upper velocity limit_o x (Factor) 1.25

Lower Velocity Limit_T = Lower velocity limit_o x (Factor) $^{0.8}$

These formulas were devised for this program from available data; no such formulas were found in the literature. The lower velocity limit figure is related closely to the effect of temperature on hardness and surface energy.

The modified values for the upper and hypervelocity limits are stored in VUP1A and VUP2A. The lower limit is placed in SKRPTE (I, 3), other SCRPTE constants are passed to SKRPTE unchanged.

Program P4255 calls Subroutine A.

(7) Subroutine LT

This subroutine tests the total number of segments impinged by the plume against the maximum the program can handle, which is 50.

If more than 50 have been identified in Program (5, 1), this subroutine truncates the number to 50 and stops the calculation of affected segments. A message is printed warning of the truncation.

Subroutine LT calls Subroutine A.

o. Primary Overlay (12,0) = Program P42120

This program is designed to acquire data about characteristics of the plume at the locations on the satellite where Program P4250 calculated that impingement occurs. The primary data input occurs on TAPE14, which is supposed to have been written by <u>KINCØN</u> in response to TAPE32. If data are not available on TAPE14, there is a default to reading data from cards (file EDATA = TAPE11). These default data are then stored on TAPE14.

The data required consist of the following:

	- 10	3		
Diameter of Small Droplets (from condensate)	=	PLMCHR(1)	=	SPRPDM
Diameter of Large Droplets (from incomplete combustion, etc.)	=	PLMCHR(2)	=	LDRPDM
Velocity of SD	= .	PLMCHR(3)	=	SDVEL
Velocity of LD	=	PLMCHR(4)	=	LDVEL
Angle of Impingement, SD	=	PLMCHR(5)	=	SDAIMP
Angle of Impingement, LD	=	PLMCHR(6)	=	LDAIMP
Flow Velocity, SD	=	PLMCHR(7)	=	SDMAPS
Flow Velocity, LD	=	PLMCHR(8)	=	LDMAPS
Plume Concentration for 25 mass fraction species of plume constituents		PLMCHR(9) PMCMP (1)	to to	(33) (25)
Chemical Composition of Small Drops, Species	=	PLMCHR(34) by 2's (1, 1) to (10, 1)	to =	(52) SDCMP
Concentration	=	PLMCHR(35) by 2'3 (1, 2) to (10, 2)	to =	(53) SDCMP
Chemical Composition, LD, Species	=	PLMCHR(54) by 2's (1, 1) to (10, 1)	to =	(72) LDCMP
Concentration	=	PLMCHR(55) by 2's (1, 2) to (10.2)	to =	(73) LDCMP
Plume Temperature		PLMCHR(84)	=	РМТМР
Plume Pressure		PLMCHR(85)	=	PMPRS
Plume Density		PLMCHR(86)	=	PMRH Ø
Plume Viscosity		PLMCHR(87)	=	PM VIS

(Unused at present, PLMCHR(74) to (83) and (88) to (90).

These data are required for each impinged segment and each time slice. If data are not input for certain time intervals or segment locations, the program will use data for an earlier time segment or a lower segment number of both.

Program P42120 prints a listing of representative plume characteristics.

Program P42120 calls Subroutines A, FAULT, and D.

P. Primary Overlay (13,0)

(1) Program P42130

This program and its subroutines calculate the results of the plume-surface interaction in terms of surface abrasion, condensation on the surface from plume vapors and deposition on the surface of plume-carried droplets.

A simplifying assumption is made--if the conditions of the plume as it impinges a satellite segment lead to abrasion, neither deposition nor condensation can occur during the same time element on the same segment.

The primary overlay buffers in the data required for calculations and storage then it calls subroutine RANDAL to perform the surface effect calculations. RANDAL in turn calls MADDY and RØBERT.

If deposition or condensation occur, it is necessary to calculate the results of chemical interactions between depositing species, and between new and previous deposits. When this is done, data are stored, a flag called MCFLG is set and a return is made to the principle overlay for calls to Overlays (15,0) and (7,0) which perform the chemical calculations. When these two routines are finished, the flags cause (13,0) to be recalled and the computation reenters at the location from which the return to (0,0) was made. This sequence is necessary due to the core size requirements of (13,0), (7,0) and (15,0). The latter two cannot fit in as secondary overlays to (13,0).

Calculations are performed for all segments during a single plume operating time element. Then the total time (sum of all time elements computed) is compared to the thrustor duration. If they are equal, the exposure is complete and the programs prints a note of this and returns to (0,0) for calculation of the surface state. If the exposure has not reached completion, the program checks to see if the plume characteristics input was from cards or from KINCON. A check is also made as to whether there will be a change in the plume configuration in the next cycle. Depending on the source of the data and the shape of the plume, the flags JMP2 or JMP3 are set and a return is made to (0,0) to get the data needed for the next time element.

The main overlay calls in Overlay (5, 0 et seq) for input from \underline{KINCQN} or if there is no change in plume geometry for card data input, the calculations go to (5, 4) and (5, 5) and the last part of (5, 0) to set up the current surface mechanical conditions before redoing the plume effects. If

the plume configuration changes (for card input), Overlays (5, 1) and (5, 3) are called to calculate the new set of affected segments, then the surface mechanical conditions are determined. The program then returns to (0, 0). If there are new segments affected, the calculation calls in P42120 to check for new card data on plume characteristics, then returns to (0, 0). In either case, P42130 is then recalled to calculate the plume effects during the next time element.

Program P42130 calls (in effect) Programs P4250, P4270, and P42150, and Subroutines A. FAULT and RANDAL.

(2) Subroutine RANDAL

This routine calls in information from TAPE12, TAPE14, and TAPE19 on the properties of the plume, the surface, and the current state of the surface. The plume characteristics are loaded to the variables described under Overlay (12, 0).

The calculations of the effects of the plume on the segment surface are then initiated.

The first step is calculations of the abrasion wear, using the fatigue wear term from the Nielson-Gilchrist equation (Reference D-2).

The wear for the two particle sizes is summed, and the abrasion depth is calculated by dividing by the area.

The abrasion depth is then compared with a standard value, which was either input to the program, or the default value of 10^{-10} inches. If the depth worn away is less than this value, it is taken as negligible, and the program turns to calculating deposition and condensation by means of a call to Subroutine RØBERT.

If abrasion is sufficient, the surface finish resulting from the abrasion is calculated, based on the assumptions:

- (a) The surface roughening due to contact by a drop is proportional to the drop radius and a stepped function of the drop velocity.
- (b) Once a local region has been roughened by an impact of certain size and velocity, additional impacts of similar size and velocity do not cause additional roughening.
- (c) The probability of impact by a droplet is independent of the previous history of the surface location.

From these assumptions and the data on the plume characteristics, the final finish is calculated as follows

SURFIN = Roughening factor = drop diameter

If velocity is greater than upper velocity limit,
roughening factor = 2.5 * diam

If velocity is greater than hypervelocity limit,
roughening factor = 5 * diam

The total mass of droplets per second per unit plume area (per-pendicular to the plume):

(PIMP) = Mass/second-ft² * time slice.

The area of the segment impacted:

(AIMP) = PIMP * π * drop diam. * Cos (impingement angle)

The fraction of the segment area impacted:

(FAMP) = AIMP/Segment area

The final finish is then:

After determining the surface resulting from abrasion, a call is made to a special entry in Subroutine RANDAL (CALL TYLER) to set up the cycle through Overlays 15,0 and 7,0 which determine the chemical effects of the abrasion.

As mentioned earlier in this section, if abrasion is not signified ant, condensation from plume vapors is calculated in Subroutine RØBERT. After the return from RØBERT, the present subroutine, RANDAL calculates the extent of deposition of material from droplets in the plume.

The weight deposited is proportional to the ratio of the difference between the droplet velocity and the lower limit divided by the velocity lower limit. This velocity ratio, multiplied by the droplet concentration, the time, and the sine of the impingement angle, provides the mass depositing. Separate calculations are made for the large and small drop distributions. After each a call is made to subroutine MADDY, where the composition of the deposit is determined from the plume characteristics data, followed by the CALL TYLER route to (7,0) and (15,0) to calculate the chemical composition of the deposited layer.

The routine sets some counters for use by overlay (13,0) and returns to (13,0).

1

Subroutine RANDAL calls subroutines A, FAULT, RØBERT, and MADDY and the special entry TYLER to RØBERT.

(3) Subroutine MADDY

The subroutine looks up the data on droplet composition and calculates the mass of each species deposited, then selects the 4 (or 5, if water is one of the deposits) most abundant species and enters their ID numbers and masses into CQMMQN/10/for use by Overlays (7, 0) and (15, 0) in calculating the final composition of the deposits.

Subroutine MADDY calls Subroutine A.

(4) Subroutine RØBERT

This routine calculates the condensation of plume constituents from the plume vapor onto the satellite surface. The relations used assume an identity of heat transfer and mass transfer processes. Because of this, mass transfer coefficients are not required; heat transfer coefficients are sufficient.

The discussion by Trebal upon which this algorithm is based assumes only one species condenses (or evaporates) although the equations are generalized for an indefinite number of species. However, the degrees of freedom are too great for the number of data for the plume, so the simplifying assumption has been made that each condensable species acts independently of all others during all steps of the process. Only at the end, when the temperature of the condensed film of each substance is calculated, are mutual effects considered. After a film temperature for each species is obtained, the various temperatures are averaged, and this average is used for the final cycle of deposit quantity calculations. For certain species, not all of the data needed for calculation of deposition is available. Certain factors, such as liquid viscosity, are hard to obtain for species such as carbon. The program maintains running averages of certain variables, and these are available as approximations when specific data are not supplied or calculable.

In the first step, the molecular diffusivity of each species at the plume temperature is calculated from the boiling point, molecular radius, molecular weight, plume temperature and pressure, and a collision function. The calculation assumes that the species being treated exists in an environment of nitrogen gas--none of the other condensable species are part of the calculation.

The mean specific heat and the mean molecular weight of the plume are calculated from those properties of the plume constituents and their concentrations. The plume viscosity is also calculated if not supplied by KINCQN, by calculating the viscosity of each species at the plume temperature and correcting the nitrogen viscosity for the species at their concentrations.

The plume Reynolds number is then derived

in which a characteristic length of 1 foot is assumed, as appropriate for typical satellite dimensions.

The coefficient of mass factor CØFMT is calculated from the Reynolds number, using the vertical plate correlation, with corrections for the angle of impingement. The data used to derive the equations are found in the ASHRAE guide and Data Book and the ASHRAE Handbook of Fundamentals.

The Prandtl number of the plume is calculated as

The Schmidt number for each species is calculated from the formula

and then used to determine the formal mass transfer coefficient, ZLS, for each species

$$ZLS = \frac{3600. * CØFMT * Plume Velocity * Plume density}{Mean Molecular Weight}$$

$$* \left(\frac{1}{SCHMDT}\right)^{2/3}$$

where $C\Phi$ FMT is a mass transfer factor calculated from the Reynolds number and the angle of impingement,

$$CQFMT = \frac{(1 + 0.07 \text{ Sin (impingement angle)} * 0.664}{\text{REN}Q}$$

The heat transfer coefficient of each plume species is calculated and the values are summed. The relation used is

Plume Velocity * Plane Density * Specific Heat * Mole Fraction * Mass transfer coefficient
Schmidt 2/3 * Prandtl * Mass flow rate

The rate of heat transfer into the satellite structure is then calculated. A thermal conductance is calculated on the assumption that existent deposits and coatings conduct toward the interior of the satellite, using input data thermal conductivities and known thicknesses. The bottom layer, usually aluminum, is assumed to conduct "radially" along the satellite surface, through one half of its area in contact with other segments, and the contacting segments are assumed to be at the prevailing satellite structure temperature.

It is then assumed that the species depositing from the plume will be at a temperature intermediate between the plume temperature and the satellite wall temperature. The temperature will be such that at the rate of condensation of condensing species, which is controlled by the vapor pressure of the condensing species at the film temperature, the heat input from cooling the species and releasing the heat of condensation must balance the heat flow into the satellite structure from the film temperature to the main satellite temperature.

To calculate this, a film temperature is assumed, and the deposition rate and hence the heat flux rates are calculated. If the heat flows do not balance, the temperature is corrected in the direction which will bring them closer to balance, and the process repeated until a balance is struck. This film temperature is stored, and the process repeated for all condensing species.

When all species have been calculated, the various film temperatures are averaged, and this mean value is taken as the most probable temperature. New deposition rates are calculated for each species at the mean temperature, and then the mass of each material and total quantity of heat deposited is calculated.

The data are stored for use by Overlays (7,0), (15,0) and (13,0).

Then flags are set, and the previously described return to (0,0) and calling of the chemical reaction overlays occurs. After (7,0) and (15,0) have been completed, the calculation returns to Subroutine RØBERT for wrap up and return to RANDAL.

q. Primary Overlay (15,0) = Program P42150

This program determines whether there are any chemical reactions which can occur between species currently depositing, and if such reactions are possible, calculates the products and their concentrations. It is a simplified version of Overlay (7, 0); the latter also considers the reactions between new and previous deposits. The logic followed and the chemical reactions calculated are the same as in (7,0), but fewer checks, comparisons and calculations need to be made; see (7,0) for a description of the calculations and logic.

Overlay (15,0) calls Subroutines A, LE, FAULT, and MNØ3, and Function RETRHØ.

r. Secondary Overlay (10, 1) = Program P42101

This routine calculates accumulated exposure time for the whole program after completion of each exposure cycle. It is for checking purposes after the calculations are all completed. It also sets the loop counter back to zero.

Overlay (10, 1) calls subroutine A.

s. This completes the sequence of calculation routines for plume exposure, which started with the initial call to (5,0), and the program returns to (0,0).

The next sequence is the calculation of space exposure effects. This is a simpler set of routines, because of the initial assumption that conditions external to the spacecraft remain constant at a pressure and temperature originally input in Overlay (14,0). It is important to note that although a and are known for the satellite, no accounting is made of radiant heat transfer to or from the satellite or of heat input to the satellite shell from internal processes. All temperature changes in the shell are assumed due to mass transfer processes on the satellite exterior. It is beyond the scope of SURFACE to include radiant and convective heat transfer, although the importance of such processes are acknowledged. There are a number of effective computer programs to calculate radiant heat transfer to and from satellites and the plume, space, the sun, and the earth, and it would be an important step to integrate CONTAM with such a program.

t. Primary Øverlay (6, 0) = Program P4260

This program and its subroutine PATANN have the function of determining the effects of exposure to a space environment on deposits originating from plume exposure. The program does not consider the effect of space conditions on structural materials, even when it is known that such materials can change. The steps used are the following:

- 1. The array SURDES is screened for segments with deposits and all calculations are limited to such segments.
- 2. Chemical and physical chemical properties related to volatility are called in from TAPE3.
- 3. Surface heat loss by evaporation of volatile species must balance heat inflow from the satellite structure, resulting in an equilibrium surface temperature. These heat fluences are calculated in, the same fashion as is done for the evaporation of intra-thrustor deposits in program TCC.
- 4. With the surface temperature established the rate of evaporation becomes constant.
- 5. The amount of material lost of each volatile species is calculated for each time slice.
- 6. The equations used are described fully in TCC. (See Appendix A.)

The assumption is made that drastic chemical changes are not likely to be caused by evaporation of volatiles, so it is unnecessary to calculate whether or not new chemical reactions can occur after each time slice. Instead, the complete space exposure is calculated in (6,0).

At the end of the exposure time, Subroutine PATANN is called to store the exposure results, then (6,0) returns to (0,0).

Overlay (6,0) calls Subroutines A, FAULT, LE and PATANN, and function RETRHO.

u. Following completion of the space exposure calculations, the main overlay calls in Overlays (7,0) to check for chemical changes, then (11,0 et. seq) use called to determine the current surface state, and finally (10,1) is called to accumulate the total exposure time.

The main Overlay (0, 0) then sequences plume and space exposures until all instructions have been fulfilled. It then calls the termination routine.

v. Primary Overlay (16,0) = Program P42160

This Overlay copies appropriate data from array SURDES to array EFFECT so that EFFECT contains the latest data for all segments. EFFECT is then copied to the PUNCH file in the same format as the input from CDATA in Overlay (4,0). This allows the results from one set of calculations to be used as the initial conditions for a subsequent set.

w. Primary Overlay (10,0) = Program P42100

This routine wraps up the program by printing a listing of the number of calls to all overlays and subroutines, the total exposure times accumulated, and a closing logo.

D. 4 PROGRAM USER'S MANUAL

a. General

The SURFACE program is the fourth link of the CONTAM computer program. It may be run as a subprogram to CONTAM under control of subroutine EXEC, or as an independent program. It was developed on a CDC 6500 computer using FORTRAN IV language. The SURFACE program requires 275,0008 words of core storage. Conversion to another computer system should be straightforward, providing that sufficient core is available. Two subroutines written in the CDC assembly language COMPASS are not essential to the program and can be eliminated readily.

b. Input Data Descriptions

Six named files plus the standard INPUT card file are used for basic formatted data input to <u>SURFACE</u>. Use of several files is necessary to allow a variety of sequences of exposure calculations, all of which require data inputs, without simultaneously requiring major reshuffling of the data input file whenever a different sequence of calculations is to be carried out.

The input files are identified as follows:

External fid	Program (Internal) fid	
INPUT	17	
ADATA	16	
INDATA	5	
BDATA	8	
CDATA	9	
DDATA	10	
EDATA	11	

In addition, when SURFACE is run under \underline{CQNTAM} EXEC in conjunction with the other subprograms, unformatted data supplied from these programs are read from two files

External fid	Internal fid
TAPE31	31
TAPE14	14

If TAPE14 is not supplied, or if the data on it are incomplete, SURFACE generates a complete set of data and stores it on TAPE14 for later use. The input file DDATA is used instead of TAPE31 when <u>SURFACE</u> is run without the other <u>CØNTAM</u> subprograms.

Data statements are used for some values, but they are primarily initializations to zero or else default values.

. c. Data Categories

The following inputs are the major categories required for SURFACE:

Propellant Properties and Thrustor Characteristics

External Materials

Satellite Configuration

Projection(s) Configuration(s)

Structural Materials

Propellant Temperature (Test Case)

Optical/Thermal Properties

Segment Structure

Program Option Selection

Modified Surface Conditions (Test Case)

Program Option Selections

Pulse Characteristics

Program Option Selection

Coordinate Transformation

Thrustor to Satellite Satellite to Thrustor

Molecular Weight

Wear Constants

Plume Characteristics

Velocity Limits

Heat Transfer Coefficients

Plume Physical Properties

Plume Configuration Change

d. In the following sections, the inputs and their functions are described, the input format is given where required, and typical inputs are illustrated. The order of presentation in this manual is DATA statements, program ASSIGNMENT statements, and then the input files in the order in which they are listed in the two tables in (b). Note that this order for the files is not necessarily in the same order in which the program will access the files, because the latter is determined by the data themselves.

e. DATA Statements

Only data which are used in computation are presented. Data used for variable $F \emptyset RMAT$ statements, Hollerith data, control of files and buffers, and program sequencing etc., are omitted here.

Variable	Use	Initial Value
	LINK 0,0	
ACP(I), I = 1, 25	Constant in denominator of Calingaert-David equation	43.
PI	Trigonometric calculations	3. 141592
R	Universal Gas Constant	8.31436×10^7
CMETL	Convert inches to centimeters	2.540005
CMETA	Convert square inches to square centimeters	6. 4516258
CMETV	Convert in 3 to cm 3	16. 387162

Variable	Use	Initial Value
CMETM	Convert pounds mass to grams	453. 5924
CMETF	Convert pounds force to dynes	444822.
CENTCG	Convert cal/gm to Btu/lb	1.8
CMETP	Convert psia to millibars	6.8947×10^4
CRAD	Convert degrees of arc to radians	0.0174532925
CKTR	Convert Kelvin to Rankine	1.8
CSGTED	Convert specific gravity or g/cc to lb/ft ³	62. 427 961
CVPTE	Convert poise to lb(m)/ft-sec	6.72×10^{-2}
CSTTE	Convert dyne/cm to lb(m)/in	5.71015×10^{-6}
TØTIME	Duration of plume or space exposure cycle (sec)	0
L		0
DELTMA	Duration of pulse slice used in computation	0
EXTYPE	Type of exposure	0
	<u>Values</u> <u>Meaning</u>	
	-1 End of program 0 Initial conditions 1 Assigned conditions 2 Plume exposure 3 Space exposure	
IPXCT	Number of plume exposure cycles completed	0
ISXCT	Number of space exposure cycles completed	0
P	Quarter latus rectum of plume	0
DELP	Change in P for next time slice	0
ITRNCL	Flag to control type of coordinate transformations to be computed	0

*

Variable		Use	Initial Value
IF6	Flag to conconditions	ntrol listing of initial data	6
		lists all data stops listing of initial conditions	
	SUBF	RØUTINE WEBBER	
REDPRP(I), $I = 1, 2$	500 In: <u>P</u> I	itialize <u>RED</u> UCED R <u>OP</u> ERTIES array	-1.
		LINK 2,0	
A	Variables	used as parameters	0
В	in call to V	used as parameters ication of materials /ISEST	0
		LINK 6,0	
SURH(I), SURI(I), I	va co	itial zero values for riables used to carry ded thickness of posits for listing	20 * ₺

Typical DATA assignments are shown in Figure D-5.

f. Assignment of Constant Values to Variables

Variable	Use	Assigned Value	At Card
	LINK 0,0		
STPFLG	Controls mode of execution of SURFACE	0	0/0.770
	Value Effect		
	 Run complete program. Stop after calculating plume intercepts. Start at location immediately after above stop. 	_	
NTR	ID number of thruster currently operating	0	0/0.842
IGØTST, IPXTST, ISXTST	Total number of all cycles, plume exposure cycles and space exposure cycles, respectively.	0	0/0.845, 847

Figure D5. Typical DATA Assignments

<u>Variable</u>	<u>Use</u>	Assigned Value	At Card
RDWRT	Flag to allow/stop listing of NAMELIST correction inputs	1	0/0.941
	<u>Value</u> <u>Effect</u>		
	0 No listing 1 Listing produced		
NP1S	Total cycles to be calculated; reset to maximum value of 10 if data input would result in a greater value.	(10)	0/0. 9 <u>6</u> 9
TIME	Time since start of latest exposure cycle calculation. Set to zero prior starting the cycle	0.0	0/0.1028
16F	Resume listing of results from calculations if currently turned off. I6F is the "unit number" for certain write statements. If set to 7, listing is turned off.	6	0/0.1125
NØJ	Used in GØ TØ NØJ state- ment in certain sub- routines and overlays to cause jump back to state- ment 231 in Link 0,0	ASSIGN 231	0/0.1218
	LINK 1,0		
DEDPTH	Default value for lower limit for effective deposit depth	$(1 \times 10^{-9} \text{ in})$	1/0.306
SURDES(I, 7)	Default value for surface temperature	529. 2K	1/0.668
	LINK 14,0		
DATA(119)	Default value for thermal conductivity of liquid $N_2^{\phi}_4$	6 x 10 ⁻⁴	14/0.627
DATA(139)	Default thermal conducti- vity of MMH	0.279×10^{-4}	14/0.657
DATA(140)	Default accommodation coefficient for boiling deposit	0.1	14/0.659

SCRPTE(I, 3), $I \le 25$	Default value for velocity lower limit for structural materials	400	14/0.1160
SCRPTE(I, 3), I > 25	Default value for velocity lower limit for deposits	650	14/0.1163
SCRPTE(I,4)	Default value for wear rate exponent	-3.19	14/0.1164

LINK 7.0

Starting at Statement Numbers 1460 (Card 7/0.789) arbitrary factors for calculation of final surface finish are assigned on the basis of surface temperature, deposit thickness, and deposit melting point. No good general method to calculate these factors in available.

Starting at Statement Numbers 1670, (Card 7/0.941) and at 1720 (Card 7/0.969) the variables ACIDR and BASER are assigned constants which depend on the specific acid and base for which calculations are being made. The values assigned to ACIDR and BASER are the equivalent weights of the acid and base.

Starting at Statement Number 1910, (Card 7/0.1078) the molecular weights of the products are assigned to the variable PRØD.

LINK 15,0

Identical equivalent weight and molecular weight assignments to those in Link 7.0 are made starting at Statement Numbers 600, 625, and 710. (Cards 15/0.501, .529, and .638, respectively)

Subroutine TRANS

At several locations, the amount of energy transmitted through transparent interfaces is assigned for cases where measured values are not available. Transmittance values of 0.96 are assigned for all cases except for infrared and visible light from vacuum to condensed phase, which is assigned a value 0.98.

g. Punched Card Input to File TAPE17 = INPUT

This data input is primarily a set of NAMELIST inputs for supplying a few changes to the standard data inputs. The latter are normally resident as card images on a set of files, and if only a few changes are required it is often easier to use this NAMELIST capability rather than change the basic data files. See Section D-3.g for a description of the eight NAMELIST's used. When NAMELIST input is to be used, the card SWITCH, 1. should be included in the control deck.

The card file is also used input the Mode Control flag, STPFLG, which is described in Section D. 3. n. (1).

The card file deck should be arranged with the STPFLG card first, followed by the NAMELIST cards in any order. STPFLG must be present for the program to run.

Note that Link (2, 1) of <u>SURFACE</u> actually reads in the NAMELIST portion of the INPUT file as a unit, and then stores it unchanged on TAPE18. The links which use NAMELIST read from TAPE18.

Variable	Read Format	Notes
STPFLG	10X, I5	Values of 0, 1, 2 are accepted (Negative values are treated as 0)
	NAMELISTS	Standard CDC NAMELIST formats with \$ in column 2 at start, etc., are used.
NP1S1	NAST1 (Called in 0,0)	Corresponds to NPIS, the total number of exposure cycles to be calculated. Maximum value is 10.
NPSEQ1(I, J)	NAST 1	Corresponds to NPSEQ and details the exposure details. Maximum value of I is 10, of J is 3.
		NPSEQ1(I, 1) is cycle number (I, 2) is type of exposure (I, 3) is exposure duration
ISXTS1	NASTI	Corresponds to ISXTST, the total number of space exposure cycles
IPXTS1	NAST1	Corresponds to IPXTST, the total number of plume exposure cycles
IG ØTS)	NAST1	Corresponds to IGØTST, the total number of cycles, including initial set up and assigned exposure conditions
11S1	NAST2 (Called in 4,0)	Corresponds to IIS, the number of affected segments on the satellite. The maximum value is 175.
EFFEC1(I, J)	NAST2	Corresponds to EFFECT(I, J) I is IIS J corresponds to the particular condition to be entered, its maximum value is 11. It is used as follows:
		J = 1 is Segment X Coordinate 2 is R coordinate 3 is 9 coordinate 4 is segment ID 5 is temperature 6 is top layer material 7 is top layer thickness 8 is top layer surface finish

Variable	Read Format	Notes
		9 is 2nd layer material 10 is 2nd layer thickness 11 is 2nd layer surface finish
Т ØТІМ1	NAST3 (Called in 5, 0)	Corresponds to TØTIME, the total duration of the current motor pulse. It is the same as NPSEQ(I, 3) discussed above, but if only the duration of exposure is to be changed, this is a more convenient input.
DELTM1	NAST3	Corresponds to DELTIME, the "time slice" used for calculations of exposure effects. Reset to 1/2 of TOTIME if greater than TOTIME.
P1	NAST4 (Called in 5,0)	Corresponds to P, the 1/4 latus rectum of the thruster plume
DELPI	NAST4	Corresponds to DELP, the amount that P decreases for each time slice during start-up transients
NTRI	NAST4	Corresponds to NTR, the ID number of the active thruster during the current plume exposure. Program will stop if the value is greater than the total num- ber entered during input of the satellite description.
ISTS1	NAST5 (Called in 10, 2)	Corresponds to ISTST, the flag which controls whether assigned conditions data are to be discarded or used as the basic conditions for further exposures.
		ISTST = 0 - discard > 0 - use as base
IP11	NAST6 (Called in 12,0)	Corresponds to IP1 and identifies the calculation cycle for which the data to be input will apply
IP21	NAST6	Corresponds to IP2, the ID of the affected segment
IP31	NAST6	Corresponds to IP3, the specific plume property of interest.

Input of IP11 or IP21 is used to change the time slice and segment ID assignment of plume characteristics data which is already input in the regular data file, TAPE11(=EDATA.) If the characteristics are to be changed, or new ones entered, NAST7 must be used.

Variable	Read Format	Notes
ЛЛ	NAST7 (Called in by 12,0)	Corresponds to IP1, the time slice or cycle number. Used as J1 in the program to avoid resetting IP1
KIK	NAST7	Corresponds to IP2, the segment ID. Used as K1
PLMCH1(IP3)	NAST7	Corresponds to PLMCHR(IP3), the plume characteristics. These are assigned as follows:
		PLMCHR (1) small drop diameter (in) (2) large drop diameter (in) (3) small drop velocity and gas velocity (ft/sec) (4) large drop velocity (ft/sec) (5) angle of tangent of path of small drop measured relative to plume axis. Also for gas (radians) (6) tangent angle for large drop (radians) (7) mass flow rate for small drops (lbs/sq ft-sec) (8) mass flow, large drops (lbs/ft-sec) (9) N2Ø4 concentration in plume gas (mass fraction) (10) HNØ3 (11) MMH (items 10-31 in units of mass fraction) (12) MMH. H2Ø (13) MMH Nitrate (14) MMH Nitrate (14) Ammonia (17) Ammonia (17) Ammonia hydroxide (18) NH4NØ3, hydrated (20) Mixed fuel nitrates, hydrated (21) AL2Ø3 (22) HCL (23) Hydrochloric acid azeotrope (24) Iron oxide (25) Carbon (26) NH4CL (27) NH4CL, hydrated (28) N2H4 (29) N2H4. H2Ø

Note: Items 10-31 have the units of mass fraction

variable	Read Format	Mores
Note: Odd numb	ered items from 35 t	30 N ₂ H ₄ . HNØ ₃ 31 N ₂ H ₄ . HNØ ₃ hydrated 32 Unassigned 33 Unassigned 34 to 53 are the composition of the small drops; up to 10 components can be handled. The ID of each component is entered into the even numbered locations (34, 36 52) and the corresponding weight fraction concentration is entered in the next, odd, location (35, 37, 53). 54 to 73 are the composition of the large drops, entered in the same way as the small ones. o 73 have the units mass percentage.
		74 to 83 are unassigned 84 plume temperature (OR) 85 plume pressure (PSIA) 86 plume density (lb/ft³) 87 plume viscosity (lb/ft-sec) 88 to 90 Unassigned
NPRFL	NAST8 (Called in by 11,0)	Corresponds to NPRFLG, the flag which indicates that absorptivity and transmissivity ratios are to be calculated for projections and/or limited region of the satellite body. Integer NPRFLG = 0 Do not determine for other regions ≥ 1 Calculate for special regions
MS11	NAST8	Corresponds to MSS1, the lower segment ID for special regions. Integer
MS21	NAST8	Corresponds to MSS2, the upper segment ID. Integer
PRJNM	NAST8	Corresponds to PRNAME, the title for the listing of the special region. Maximum of 8 Hollerith characters.

Notes

h. DATA Files

Variable

Read Format

These files are typically used in the form of disc files, copied from the multifile magnetic tape of the program <u>SURFACE</u> and all its auxiliaries. There also are UPDATE $\slash\hspace{-0.4em}PL'$ s of the DATA files on the tape. These $\slash\hspace{-0.4em}PL'$ s can be used to make corrections to the DATA files -- the output $\slash\hspace{-0.4em}C\slash\hspace{-0.4em}PMPILE$ file is

then named, for example, ADATA by means of the UPDATE parameter C = ADATA. When UPDATE is used to change the DATA files, it is essential to use the D parameter to ensure that data in columns 73-80 is not destroyed. A typical control card sequence for updating one of the DATA files is illustrated in Figure D-6.

The listing given in the figure shows only directives for updating the specific data file, and does not include other processes which would usually be done at the same time, such as executing the program with the new data, and generating a new magnetic tape on which the new DDATA replaces the old, and DDNPL replaces DDØPL.

The UPDATE deck names for the various data files are

File Name	Deck Name
ADATA	AD
BDATA	BD
CDATA	CRCD1
DDATA	DD
EDATA	ED
INDATA	IND

i. File ADATA = TAPE16

This formatted file contains a description of the satellite configuration and the physical, chemical, and mechanical properties of the materials of construction and of plume/deposit constituents.

A maximum of 175 separate locations on the satellite can be identified; 25 each structural materials and plume/deposit constituents can be handled.

ADATA also contains the instructions for the computing sequence, and several program control flags.

In the description which follows, the discussion is separated into sections which correspond to the various devisions of the data on the file

(1) Initialization

Variable	Format Structure	Comments
None (in 0,0)	(///)	Skip file title cards
IERTST (in 0,0)	(5X, I5)	Maximum number of minor errors before program self-aborts. Defaults to 10.
IT1; FACTR(K), K = 1, IT1	5X, I5, 5X, 9F7.0	FACTR controls the duration of the time slices during space exposure in (6,0) as a function of temperature

######################################	2000 2000 2000 2000 2000 2000 2000 200	COPY, A.B. COPIES COMPLETE HULTIFILE FROM TAPE TO DISC. FREES THE TAPE DRIVE.	A 18 18 18 18 18 18 18 18 18 18 18 18 18	KIPF, 8, 9, 17, 8, SKIP TO OLDPL OF DDATA, PARE OLDPL AN INDEPENDENT FARE THE DISC SPACE.	FL:40000. PDATE(D:E.F.U.P=DEOPL:N=DDNPL:C=DDATA	00 Y 8		SFOURNCE DD	
----------------------------------------	-------------------------------------------------------------	-------------------------------------------------------------------------------	------------------------------------------	-------------------------------------------------------------------------------------------	----------------------------------------------------	--------	--	-------------	--

Figure D-6. Directives for Correcting File DDTA:

IN 12-1123

Variable	Format Structure	Comments
DECPTH, DEDPTH (in 1,0)	7X, F9.0, 7X F9.0	DECPTH is used in (13,0) to test whether abrasion is signif- icant or whether deposition is the primary effect
		DEDPTH is used in (13,0) to test whether deposition is significant
NAME (in 1,0) Four calls: NAME (I,1), I = 1,13	13 A 6	NAME carries the titles used for materials, constituents, and projections from the satellite
NAME (I, 1), I = 14,15, I = 1,3; NAME (I,2), I NAME (I,2), I = 8,19 NAME (I,2), I = 20,23		NAME (I, 1) is materials NAME (I. 2) is projections NAME (I, 3) is constituents
		Unused locations are set to UNASND.
		The 2-D array NAME (25,3) is equivalenced to the linear array ALNAME(75) for computations in which the single dimension will save time.

In addition the linear array NAAM(5) is equivalenced to negative locations of ALNAME by (NAAM(5), ALNAME(1)) and the values

NAAM(1) = 6HERØDED (2) = 6H HØLE

(3) = 6H NØNE

(4) = 6HUNASND

(5) = 6H (blank)

set by a DATA statement in Link (0,0). These latter names are used to print the effects of abrasion.

The materials and constituents currently used in SURFACE and stored in array NAME are shown in Figure D-7. A copy of a listing of the first section of ADATA is presented in Figure D-8 to show the appearance of the file. (The names are abbreviated in the program.)

(2) Model Satellite Configuration

The structure used in the standard data is a model satellite. The features of its structure are shown in Figure D-9.

The description of the basic satellite is read from ADATA and is entered into the array SURDES (SURface DEScription), but it only partially

***************************************	• 1/0 • 1/0	31 31
LIST OF MATERIALS	1/0	31
***************************************	• 1/0	31 31
NAMES CORRESPONDING TO THE VARIOUS STRUCTURAL P PLUME DEPOSITS. AND SURPACE-ATTACHED COMPONENTS READ IN AND ASSISHED TO THE VARIABLE ARRAY - NAME NAME(I,1) STRUCTURAL MATERIALS; NAME(I,2) SDEPOS NAME(I,3) SPROJECTION/COMPONENT TYPES	ARE 1/0 100 1/0 1178 1/0 1/0	31 31 31 31 31 31
NAME(1,1) IS ALUMINUM NAME(2,1) IS WINDOW	1/0 1/0	- 31 31
NAME(4,1) IS INFRA RED PORT NAME(5,1) IS ULTRA-VIOLET PORT	1/0 1/0 1/0	
NAME(6,1) IS TEFLON NAME(7,1) IS GOLD PLATE NAME(8,1) IS BLACK COATING	1/0 1/0	31 31
NAME (10,1) IS KAPTAN NAME (10,1) IS KAPTAN NAMES (11,1) - (29,1) ARE UNASSIGNED	1/0	-31 31
NAME (1,2) IS N204 NAME (2,2) IS NITRIC ACID (HYDRATED N204) NAME (3,2) IS MMH	1/0	31 34
NAME (4.8) IS HYDRATED MMM NAME (5,2) IS MMM2.NITRATE NAME (6.2) IS HYDRATED MMM2.NITRATE	1/0 1/0	3/
NAME (8,2) IS AMMONIA NAME (9,2) IS AMMONIUM HYDROXIDE	1/0 1/0	
NAME (10.3) IS AMMONIUM NITRATE NAME (11.2) IS HYDRATED NH4HO3 NAME (12.2) IS HIXED FUEL NITRATES (HYDRATED)	1/0 1/0	34 31
NAME (13,2) IS ALUMIAA NAME (14,2) IS HYDROEN CHLORIDE NAME (19,2) IS HYDROCHLORIC ACID AZEOTROPE	1/0 1/0 1/0	31
MAME (16.2) IS IRON OXIDE NAME (17.2) IS GARBON NAME (18.2) IS AMMONIUM CHLORIDE	1/0	31
NAMÉ (10,2) IS HYDRATED AMMÓNIUM GHLORIDE NAME (20,2) IS HYDRAZINE NAME (21,2) IS HYDRAZINE HYDRATE	1/0 1/0	31 31 31
NAME (23,2) IS HYDRAZINE NITRATE NAME (23,2) IS HYDRATED HYDRAZINE NITRATE NAMES (24,2) AND (25,2) ARE UNASSIGNED	1/0 1/0 1/0	36
NAME(1,3) IS THRUSTOR NAME(2,3) IS RADIO ANTENNA NAME(3,3) IS SOLAR CELL ARRAY	1/0 1/0 1/0	36 36 36

Figure D-7. Structural and Plume/Deposit Constituents

1,54-1,53

DECPTHS 1.E-FIEFPINE 3.E-11
ALUMINNINDCASOLCELIAPORTLYPORTTEFLONGOLD BLACK HHITE KAPTANUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNASNDUNA AMONIANHAGH NATICIANS, ABPXFLABAL203 HCL NZH4A2N3F5C3AQFVNT

Figure D-8. File ADATA - Initial Section

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Figure D-9. Satellite Configuration

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fills the array. The materials present on the exterior and their current condition will be entered later. In addition, data on projection from the basic surface are read in separately.

Variable	Format		Comments
M1S, M2S (in 1, 0)	5X, I5, 5X, I5	= 116, M2S is th	ber of segments to be input e number of data points in array. M2S = 11
TØTAR (in 1, 0)		10X, F10.0	Total area of basic satellite.
SURDES (M) (read in M15 (in 1, 0)	l, M2), M2 = 1, M2S 5 times)	10F8.0/F8.0	Satellite configuration data.

Figure D-10 explains the assignment of various data values to locations in the array SURDES.

A part of the current input for the above variables is illustrated in Figure D-11. Note particularly the presence of values other than -1. in columns 8-11 for segment with ID numbers (column 1) of 5 and 116. These values indicate the presence of projections from these particular segments of the satellite. Subsequent inputs will supply further data about these projections. In this and subsequent inputs, and in program execution, values of -1. for data usually indicate that data are lacking.

(3) Projections from Satellite Surface

In the next set of reads, data about projections from the satellite surface are input. At present, the projections can consist of thrustors, antennas, and solar cell arrays.

The current satellite model has eight ACS thrustors, four aimed aft and four forward, located 90 degrees apart around the girth of the satellite 3.6 feet from the apex, and a ninth thrustor for translation centered on the tail of the satellite. There is also one antenna and one solar cell array.

The information about the projections is input in the following fashion:

The eighth column of the array SURDES is read for each segment, starting at the one with the lowest ID number. When a value different from -1. is found in column 8, a READ of TAPE16 is made to input data about the projection located on that segment.

Variable Format Comment

PRJN, TRX, TRR, TRTHET 10X, A6, 5X, F10.0, 2 (5X, F5.0) (Read in 1,0)

C H2S #2 15 THE X COORDINATE OF THE CENTER OF THE SEGMENT 1/0 M2S #3 15 THE A COORDINATE OF THE CENTER OF THE SEGMENT 1/0 M2S #4 15 THE ANGLE (THETA) COORDINATE OF THE SEGMENT CENTER 1/0 M2S #4 15 THE ANGLE (THETA) COORDINATE OF THE SEGMENT CENTER 1/0 M2S #4 15 THE ANGLE TO THE VERTEX OF THE NEAREST RIUME, PRO1/0 C JECTED ON THE X AXIS (A NEGATIVE VALUE MEANS THAT THE SEGMENT1/0 C W2S #7 15 THE SUBFACE TEMPERATURE OF THE SEGMENT 1/0 M2S #7 15 THE SUBFACE TEMPERATURE OF THE SEGMENT 1/0 M2S #7 15 THE BIRECTION COSINES OF THE NORMAL TO THE PLANING CONTACT IS TANGENT TO THE PROJECTION ON THE SEGMENT 1/0 M2S #7 15 TANGENT TO THE PROJECTION ON THE SEGMENT 1/0 M2S #7 15 TANGENT TO THE PROJECTION ON THE SEGMENT 1/0 M2D M3D M3D M3D M3D M3D M3D M3D M3D M3D M3	4	M2S R 1 15 SEGMENT ID	1/0	111
TASS 42 18 125 42 18 125 42 18 125 18 18 18 125 18 18 18 125 18 18 18 125 18 18 18 125 18 18 18 125 18 18 18	v		\$70	
H2S #4 15 H2S #4 15 H2S #4 15 H2S H2 15 H2S H2 15 H2S H2 15 H1CH 15 H2S H2 15 H2S H2 15 H2S H2 15 H2S H2 15 H2S H2 15 H2S H2 15 H2S H2 15 H2S H2 15 H2S H2 H2 H2S H2 H2 H2S H2 H2 H2S H2 H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2S H2 H2 H2 H2 H2 H2 H2 H2 H2 H2 H2 H2 H2 H	u	S #2 15 THE X COORDINATE OF THE CENTER OF	1,0	:
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Figure D-10. Storage Locations in SURDES for Satellite Configuration Data

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Figure D-11. ADATA Input of Satellite Configuration

The word read is as PRJN is compared with the word 6HTHRSTR. If the words match, the information read in is stored in the form input, where TRX, TRR and TRTHET are the thruster X-, R- and 9- coordinates. One set is indexed and stored for each input of this type. The implicit assumption is made that the plume direction is aft. If the R coordinate for the thrustor location is larger than the satellite radius, a second thrustor firing forward, with the same configurational constants, is assumed and entered in the file.

If the word read in by PRJN is not 6HTHRSTR, the other data read in are transformed as indicated

AREA = TRX M11S = TRR M22S = TRTHET

Then another set of variables is read in; there are M11S separate reads for these new data, which consist of configurational information similar to that read in for the original segments on the satellite body.

Variabl	<u>le</u>	Format	Comment
SURDES(MM1, MM2),	MM2 = 1, MM22S	F7.0, F9.0, F8.0, F9.0, F6.0, F9.0, F8.0	

If the seventh column (temperature) is null, a default value of 529. 2°R is assigned. No data for columns 8 to 11 are expected or accepted; that is, there are no projections on top of projections.

The process of reading in data on projections is continued until all of the original segments in SURDES have been checked for the presence of projections. Current inputs are shown in the excerpt from ADATA in Figure D-12.

(4) Material Properties

The next set of inputs consists of information about the properties of structural materials. The data are read into the array MATERAL (26, 20, 7). After input is complete, the data are stored on the unformatted file TAPE1 to save storage space in core. (Other data are read into an equivalenced array later.) The structure of the properties input is explained in the program segment shown in Figure D-13.

The level MT3 = 7 is not currently assigned.

Variable	Format	Comment
MT1S, MT2S (In 14,0)	10X, I5, 10X, I5	MT!S = 10, MT2S = 9
MATERIAL (I, J, K), J=1, MT2S read in K=6 times, read in I=MT1S times (in 14,0)	F3.0, F7.0, 2E10.7, F6.0 2F8.3, E10.7, F5.2	

Figure D-12. ADATA Input of Satellite Projections

C THE DATA CURRENTLY ENTERED IN THE PROGRAM FOR PROPERTIES ARE	14/0 369
C MATERIALS OF INTEREST, PROPERTIES OF SPECIFIC MATERIALS C (ALLOYS, COATINGS, CHRAMICS, ETC.) SHOULD BE UNTERED IF SPEC	14/0 371
C C	14/0 374 14/0 379 • 14/0 324
C HECHANICAL STRUCTURE HATERIALS	14/0 377 14/0 378 14/0 379
C HATERAL IS THE ARRAY OF PROPERTIES OF STRUCTURAL MATERIALS	14/0 300
C IT IS STORED ON TAPE 1 C HTS CORRESPONDS TO THE NUMBER OF MATERIALS IN THE TABLE C HTS IS THE SPECIFIC PROPERTY	14/0 383 14/0 384 14/0 388
C HT2 # 1 IS THE CODE NUMBER(CORRESPONDS TO NAME(1,1) C HT2 # 2 IS THE MELTING POINT C HT2 # 3 IS THE VICKERS MARDINGS	14/0 307
C HTZ = 4 IS THE BULK MODULUS C HTZ = 5 IS THE SURFACE BRERGY = SUMFACE TENSION C NIZ = 4 IS THE MEAT CAPACITY	14/0 300 14/0 300 14/0 301
C HT2 = 7 IS THE THERMAL CONDUCTIVITY C HT2 = 8 IS THE VIELD STRENGTH C HT2 = 8 IS THE DENSITY	14/0 393 14/0 393 14/0 394
C MTS CONTAINS THE CONSTANTS FOR THE PARAMETRIC EQUATION RELATED CHANGE OF MATERIAL PROPERTIES WITH TEMPERATURE	14/0 304
C PROP(F(T)) B A + B+T + C+T+2 + D/T + E/(T++2) C MTS a 1 18 THE VALUE FOR THE PROPERTY AT BOOK TEMP	14/0 300
C ERATURE (\$40 R, 88 F) C MT3 m 2 18 A C MT3 m 3 18 B	14/0 401 14/0 408 14/0 403
C MT3 # 4 18 C C MT3 # 5 18 D C MT3 # 4 18 E	14/0 494 14/0 408 14/0 406



Figure D-13. Storage Locations in MATERAL for Mechanical Properties

The data used were obtained from a variety of standard handbooks, including the Handbook of Chemistry, Handbook of Chemistry and Physics, Handbook of Applied Engineering Science, Chemical Engineers Handbook, Plastics Encyclopedia, Space Materials Handbook, ASHRAE Guide and Data Book, Military Handbook 5, Materials Selector issue of Materials Engineering, Mechanical Engineers Handbook, etc.

A single equation for thermal effects was selected to avoid carrying separate equations for each property, or even for each material and property combination. The constants were calculated from data reported at several temperatures, and least-squares fitted the equation by a small interactive computer program.

As shown by the copy of this part of the ADATA input (Figure D-14) for some materials insufficient data were found to calculate the correlations coefficients. In these cases, the room temperature constants are stored in MT3 = 2, and zeroes are left for MT3 = 3 to 6.

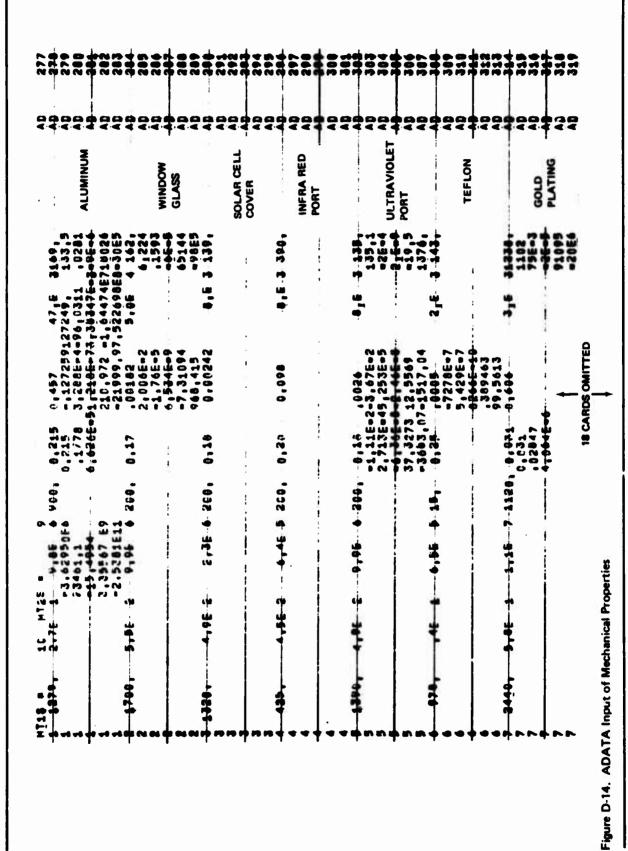
(5) Chemical Properties of Propellant/Plume Constituents

The next data input in sequence is from the file INDATA, read in by the INPUT1 input editor. This input will be described in detail later, after completing the description of the ADATA file. It basically consists of certain physical chemical properties of $N_2 \phi_4$ and MMH.

Following the above input and subsequent to the assignment of the input values to specific variables, formatted input of similar data from ADATA for other plume/deposit constituents takes place. The inputs alternate between reading how many items will be read in on the next line, and the actual values read in. The input of number of items is shown only once in the following table.

<u>Variable</u>	Format	Comments
NUDA (Read in 14/0)	5X, I5	Number of NU DAta in next line; maximum value is 5
I, CP(I) J=1, NUDA (two lines)	5(I5, F10.0)	I corresponds to the code for the con- stituents in NAME(I, 2). CP = vapor specific heat
In subsequent lines, the following variables are input:		
EPCRIT (1 line) FREZT (1 line) FRZEN (1 line) RHØL (3 lines) TENS (2 lines) SHS (1 line) TBØIL (1 line) VAPEN (1 line) VISL (2 lines)	same	Critical pressure, psia Freezing point, ^O K Latent Heat of Melting, cal/g Specific gravity, liquid Surface tension, dynes/cm Specific heat, solid, cal/g- ^O K Boiling point, ^O K Heat of evaporation, cal/g Viscosity, poise

CRS



611

This portion of the ADATA input file is shown in Figure D-15.

Note thatif two successive lines of data have the same number of items, it is not necessary to read in a new value of NUDA (cards 349 and 350), but that no harm is done if a duplicate value of NUDA is read (cards 344 and 346).

The next input consists of a number of additional physical-chemical properties of plume/deposit constituents, read in as the array CHEMIC (26, 20, 7), equivalenced to MATERAL, and then stored, unformated, on TAPE3. Figure D-16 illustrates the assignment of types of data to specified locations in CHEMIC.

The method to input the CHEMIC data is basically similar to that used for MATERAL.

Variable	Format	Comments
JC1S, JC2S (Read in 14,0)	10X, I5, 10X, I5	JC1S = 23 (number of constituents) JC2S = 13 (number of properties) Note JC3S = 6; the seventh layer is not used at present
CHEMIC(I, J, K), J = 1, JC2S (Do K = 6 times) (Do I = 23 times)	10F8.4/4F8.4	Note 2 cards for each read

The data were obtained from a variety of standard sources; Gmelin, Beilstein, Chemical Abstracts, Batelle: Propellant Handbook, CPIA: Propellant Manual, Handbook of Chemistry, Handbook of Chemical Engineering, Handbook of Chemistry and Physics, Manufacturers' literature, Kirk-Othmer, JANNAF Tables, etc. The constants for the variation of solid properties with temperature were calculated in the same fashion as described above for structural materials.

A few portions of the ADATA input to CHEMIC are illustrated in Figure D-17.

(6) Molecular Weight

The next input is that of vapor state molecular weights (MØLWT) of the plume/deposit constituents (see Figure D-18).

<u>Variable</u>	Format		Comments
MØLWT(I), I = 1,7 (Read in 14.0) 8,12 13,19 20,23	7(5X, F5.0)	4 cards	

MØLWT is equivalenced to AMW because parts of the program were written with one name for these data, and parts with the other.

Figure D-15. ADATA Input of Phys-Chem Properties

********************************	14/0	
	14/0	
•	14/0	
• PHYS=CHEM •	14/0	
• PROPERTIES OF	14/0	
- sascific deadaire -	-14/4	_
<u>.</u>	39/0	
	14/0	
	14/0	
CHEMIC IS THE ARRAY OF PHYS-CHEM PROPERTIES OF DEPOSI	714/0	
17 IS STORED ON TARE 3	-144	_
JC18 CORRESPONDS TO THE NUMBER OF MATERIALS IN THE TABLE	14/0	
JC2 CONTAINS THE SPECIFIC PROPERTIES	14/0	
JC2m 1 18 THE MATERIAL CODE NUMBER (SAME AS NAME(1,2)		
JC20 2 IS THE FREEZING POINT JC20 3 IS THE SOLID DENSITY	14/0	
JC20 4 18 THE CRITICAL TEMPERATURE	14/1	
JC20 5 18 THE NORMAL BOILING POINT	14/0	
JCZO 6 18 THE HEAT OF FUELON	14/0	
- JC2= 7 IS THE HEAT OF VAPORIZATION -	84/8	
JOSH & IN THE THERMAL CONDUCTIVITY OF THE BOLID PHAS	E14/0	
JOZ- 9 IS THE THERMAL CONDUCTIVITY OF THE LIQUID PHA	A74\0	
JC2= 10 18 THE HEAT CAPACITY OF THE SOLIS PHASE JC2= 11 IS THE HEAT CAPACITY OF THE LIGUID PHASE	14/0	_
JC2≈ 11 IS THE HEAT CAPACITY OF THE LIGUID PHASE JC2≈ 12 IS THE REFRACTIVE INDEX	14/0	
A SEC SEC SECULAR RADIUS	14/4	
JČŽ D 14 IS THE LĪGUID DENģIŤY	14/0	
JC3 CONTAINS THE CONSTANTS FOR THE PARAMETRIC SQUATION	14/0	1
RELATING BOLLD DEPOSIT PROPERTIES TO TEMPERATURE	14/0	ť
BARRIFIELL . A . BAR A FARAR A RIV A BITTE-AL	14/0	
PROP(F(T)) # A + B+T + C+T++2 + D/T + E/(T++2)	14/0	
JC3 a 1 IS THE VALUE AT \$40 R(80 F)	14/0	
JC3 a 2 18 A	14/0	
UC1 0 3 18 0	444	_
ŬÇ3(# 4 18 C	14/0	
jča » s is d	14/0	
.7c2 a ♦ 18 E	14/0	
ARAPPETICO AP I TAMER REBACTO AL PHAROTANE AF CEMBODIGHE	34/0	
PROPERTIES OF LIQUID DEPOSITS AS FUNCTIONS OF TEMPERATURE ARE CALCULATED FROM REDUCEDICATIONALLY PROPERTIES.	14/0	- 3

Figure D-16. Storage Locations in CHEMIC for Chemical Properties

JC15.	23 471.37	JC25 = 116.6	13 776,4	529.2	68,51	178,2	.000252	.002509	0,295	AD AD	366 367
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_							11317E=		.407426	AD AD	349
1.							1364,5-	y	9.0>3E-	. 1	371
1,							-209E-1	1	-7,00E-	7AD N2O	-372 373 -374
1,							.100453		-200,80	6AD	375
1,							-51,126	ý	24459,5	AD AD	376
2.	416.2	118.0	-1.	569,	12.9	204 2			A	AD	378
0.42	1.402	4, 75	•	7071	7217	2,905	-1.	.00334	0.658	AC AD	379
2,										AC	381
2,			•							AD HNO.	382
2.			•							AC -	-384
-										AD	385
2,										AC AC	387
2,										AD	388
				4 00						AC	390
£.	364				CARDS ON						
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8.	•1,	-1.	-1,	-1,	-1.	849,38	-1,	27875F-7	-1.	AC	453
7,4877	-1.	-1,						-3004E-9	-1.	AC MU	454
-1,62t-2								1.5	•	AC 1413	456
8, 1,543k-5								2551E-13	-1:	AD TA	457
6. •248.17								-,218468	-1.	AD	459
8.								31,6037	-1.	AC AC	460
•156523.				A						AD	462
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23,										AC AC	633
23,										AD N2H4	634
23,										AE HNO	+636
										ADH ₂ O	637
23,							•			AC	639
23,										AD AD	640
•										AC	642

Figure D-17. ADATA Input of Chemical Properties

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Figure D-18. ADATA Input of Molecular Weights and of Wear Equation Constants

(7) Wear Rate Equation Constants

Following the molecular weights, constants for the Nielson-Gilchrist wear rate equation are supplied from ADATA for both structural materials and solid deposits. However, there are few data for the latter substances, and approximations are supplied in the program. The constants were obtained by plotting the observed wear rates from published rain drop and dust erosion tests and observations as functions of input velocity (log-log scale) and then reading off or hand calculating appropriate constants.

The data (see Figure D-18) are read into the array SCRPTE.

SCRPTE's dimensions are 50×5 . The material code column in card images for SCRPTE corresponds to the value of J in array NAME(J, 1) for structural materials and (J, 2) for deposits; thus there can be two cards with the same values in the first column. The earliest such card in the input file contains data on structural materials, the second card contains data on deposit materials. Figure D-19 illustrates the locations in SCRPTE for specific constants

<u>Variable</u>	<u>Format</u>	Comment

SCRPTE(I, J), J = 1.45X, F5.0, 10X, G10.0, Repeat I = 1.50 times 5X, G10.0, 5X, F10.0

If the value for SCRPTE(1, 1) is zero, input ceases even though 50 cards have not been read.

When the example of the inputs from ADATA in Figure D-18 are examined, it may be seen that no data have been entered for a number of the deposit constituents. Values internally calculated by the program from the hardness are entered, if hardness is available. If not assignment of default values is done.

If 50 cards are read for SCRPTE, there is a skip read to bypass the "end of SCRPTE" card with zero in the first location.

The two velocities at which a change in form of the Nielson-Gilchrist equation occurs are next read. These are used for all substrates.

<u>Variable</u>	Format	Comment					
VUP1, VUP2	5X, F10.0, 5X, F10.0	VUP1 is less than VUP2. The values were obtained by inspection of data on high velocity micrometeoroid impact effects. When more data available, different values for different substrate materials will probably be found.					

14/01112	14/01115	14/01121
SCRPTE (SCRIPT E) STORES THE DATA USED TO DETERMINE FOR THE WEAR EQUATION.	TION DEFEXP TV RATIO IS LESS THAN CNE, (IF IS CONSTANT)	FI/SEC)
C THE ARRAY SCRPTE (SCRIPT E) STORES C CONSTANTS FOR THE WEAR EQUATION.	CRPTE INDICES FOR THE EQUAT BREATVEL/VEL-LOW-LIMITA HICH APPLIES IF THE VELOCIT ATIO IS ONE OR GREATER, E 11	C F S IS WEAR RATE TERM E F 3 IS VELOCITY LOWER LIPIT(FIXSEC)

Figure D-19. Storage Locations in SCRPTE for Wear Rate Constants

(8) Optical/Thermal Properties

The next batch of cards images in ADATA contain optical and thermal properties for the structural materials and deposit constituents. The data are read into the array PRØPTY(51, 20, 2) which is also equivalenced to MATERAL. After approximation of missing data by means of several routines, the data are stored in unformatted fashion on TAPE2. Figure D-20 describes the assignment of particular properties in array PRØPTY.

In this portion of the ADATA file, the materials to which the properties apply are not identified in the input. (For the benefit of the user, the name is given on a portion of the card which is not read.) The data are presented in the order of the materials in the array NAME, and if NAME contains the word UNASND, input is bypassed. The program supplies the material ID numbers as it proceeds, and puts them in the array.

There are two read statements associated with each material. The first read reports to the program on how many data cards and how many columns of data on each will be found by the second read. The first read can report that no data cards exist; in such cases the second read is skipped. The lack of data is not surprising in some instances; few people have considered that hydrochloric acid is a likely coating for a satellite, and studied its absorptivity and emissivity, etc. The ADATA file is illustrated in Figure D-21.

<u>Variable</u>	<u>Format</u>	Comments
N1S, N2S (Read in 14,0)	5X, I5, 5X, I5	N1S = number of data cards, max = 5. If zero, the read of PRØPTY is skipped. N2S = number of columns of data, max = 20.
read N1 times (N3 is either 1 or structural or depo	N3), N2 = 2, N2S + 1 2, corresponding to osit materials, set in the program)	if N2S. LE. 8 Format = 8F10.4 if N2S. GT. 8 Format = 8F10.4/ 8F10.4

(9) Temperature Effects

For a demonstration of the capability of calculation of properties of plume/deposit constituents at various temperatures by Subroutine WEBBER, (called by Link 2, 0), two temperatures are read in from ADATA and then used in the calculation.

Variable	Format	Comments				
TTANK(1), TTANK(2) (Read in 2.0)	10X, F10.0, 10X, F10.0	These inputs are in ^O K				

CNDS TO THE THE TANK THE THE TANK THE THE TANK THE THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TANK THE TA	/0126 /0126	0127	0127	22222	/0128 /0128 /0128 /0128 /0128	100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 100115560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 10011560 1001
PROPTY IS THE ARRAY OF OPTICAL/THERMAL CONST OF THE NAME OF ENTERIES IN THE TABLE AND CORRES OF THE NAME OF ENTERIES IN THE TABLE AND CORRES OF THE NAME OF ENTERIES IN THE TABLE AND CORRES OF THE NAME OF TABLE AND CORPES OF THE NAME OF TABLE AND CORRES OF THE NAME OF TABLE AND CORPES OF THE NAME OF TABLE OF THE THERMAL FOR THE THERMAL ENTERIES OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVITY OF THE THERMAL CONDUCTIVIT	PROPTY IS THE ARRAY OF OPTICAL/THERMAL CONSTANTS	NIS IS THE NUMBER OF ENTERIES IN THE TABLE 14 N2 - 1 IS THE CODE NUMBER OF THE MATERIAL, IT IS 14	CONSTANT THROUGHOUT EACH TABLE AND CORRESPONDS TO 14 THE SAME NUMBER IN NAME.	MATERIALS AND COATINGS), OR THE SURFACE TEMPERATURE14 (FOR DEPOSITS) 14 N2 e 3 is alpha(S), the solar absorptivity N2 e 4 is epsilon(TH), the thermal emissivity N2 e 5 is diffuse reflectivity	NZ = 7 IS THE THERMAL CONDUCTIVITY NZ = 8 IS THE THERMAL CONDUCTIVITY NZ = 9 IS REFRACTIVE INDEX (VISIBLE) NZ = 10=17 ARE TRANSMITTANCES (PER INCH OF THICKNES14 NZ = 10 AT 15 MICRON (IR)	N2 = 12 AT 2 HICRON (IR) N2 = 13 AT 1 HICRON (IR) N2 = 14 AT 0.5 MICRON (VS) N2 = 15 AT 0.1 MICRON (UV) N2 = 17 AT 0.05 MICRON (UV) N2 = 17 AT 0.05 MICRON (UV)

Figure D-20. Storage Locations in PRØPTY for Optical/Thermal Constants

VPU1=	3.634PU2= 29 N2S= 16	9.805					ALUMINU	DA DAMI	681
	4rb 4rest.	÷104.		-1.0	1.9	01407		AB	665
C.		J.	3,	0.	0.	Ŏ•	0,	AD	084
-1.0	0,	C.C5	0,93	-1.0	0,98	-1.0	-1.0	AD AC	689
	1.0 [.015	0.05	-1.0	0.92	-1.0	-1.0	-1.0	AD	686 687
-1.0	-1.0							AD	
-4 0	5.0	-1.0	0.85	-1.0	0196	-1.0	-1,0	AD	489
-1,0	3,0. C.030	-1,0	-1.0	-1.0	-1.0	-1 0	-1 0	AD	600
-1.0	-1.0	-413	41.0	-2.0	-1.0	-1.0	-1.0	AD AD	691
	4.0 0.050	-1.3	2,73	7.85	0,95	-1.0	-1,0	AD	693
-1.0	1.3			1 14	- 1			AC	694
-1,0	5,0 0,070	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	AD	695
	6,0 0.100	-1.7	-1.0	-1.0	-1.0	-1.0	-1.0	AD AD	696 697
-1.0	-1.5	- 4 .	-210	- 410		-210	0	AB	
	7,6 0.123	C. 25	1,53	-1.3	0,58	-1.0	-1,0	AD	699
-1,0	-1.5							AD	700
•1,U	•.0 2.140 •1.0	0.05	-1.0	-1.0	-1.0	-1.0	-1.0	AD	701
-210	9.0 0.160	J.06	-1.0	-1.0	-1.0	-1.0	-1.0	AD AD	702 703
-1,0	-1.0		•••	•••		410		AD	704
	0.0	-1.7	-1.0	-1.0	-1.0	-1.0	-1,0	AD	705
-1,0	-1.0	-4				. 39		AD	706
-1,0	.1.0 0.180 -1.0	-1.0	-1.0	-1.0	-1.0	-1.0	-1,0	AD	707
1	2.0 :.196	0.08	-1,0	-1.0	-1.0	-1.0	-1.0	AC 2A	70 8 70 9
-1,0	-1.0		•••					AD	710
	3,0 0.200	-1.7	-1.0	-1.0	-1.0	-1.0	-1.0	AC	711
-1,0	-1.0 4.0 0.216	-1.3	-4 5		2 8			AD	712
-1,0	-1.0	-1.0	-1.0	-1.0	-1,0	-1.0	-1,0	AB	713
	5,0 3.220	-1.0	-1.0	0.76	-1.0	-1.0	×1,0	AC	714 715
-1,0	-1.3							AD	716
	0.2 0.240	0.15	-1.0	-1.0	-1.0	-1.0	-1,0	AC	717
-1,0	5,0 0,200	-4 -	-4 4				4	AC	718
•1,0		-1.7	-1.0	-1.0	-1.0	-1.0	-1.0	AD	719
3	0.0 0.330	-1.0	J.50	-1.0	0,20	-1.0	-1,0	AD AD	720 721
-1.0	-1.5		-,-,-		• 4 - •	-, v	10	40	722
	5.3 0.360	-1.3	-1.0	-1.0	-1.0	-1.0	-1,0	AD	723
-1.0	-1.0 0.0 0.385	- 4 .				• -		AC	724
-1.0	0.0 0.38J	-1.0	-1.0	-1.0	-1.0	-1.0	-1,0	AD	725
-410	0.0 5.420	0.21	-1.0	-1.0	-1.0	-1.0	-1.0	AD AD	726 727
-1,0	-1.0		-410	1 4		-110	4410	AD	728
	0.0 0.450	-1.0	-1.0	n . 50	-1.0	-1.0	-1.0	AD	729

Figure D-21. Part A. ADATA Input of Optical and Thermal Constants

	C
-	_

•1.0	-1.0							AD	730
-1,0 -00,0	-1.6	-1.9	-1.0	-1.0	-1.0	-1.0	-1,0	AD	731 732
100,0	0.567	-1,3	-1.0	-1.0	-1.0	-1.0	-1,0	AC AG	733
130.0	3.670	0,54	-1.0	-1.0	-1.0	-1.0	-1.0	AD.	735
150.0	-1.C 0.750	-1.7	-1.0	-1.0	-1,0	-1.0	-1.0	AD	736 737
160.3	-1.5 3.830	-1.0	-1.3	-1.0	-1.0	-1.0	-1,0	AD AD	738 739
A15= 2	-1.0 '25= 16						MINDOW	AD AD	740 741
0,1	j.62	0.69	3,9A .53	9.0018	0.0	0,001 0 2	1,547	AD	742
10.3	0.09	1.85	0.10	9.0018	0.0	-1.	0,0 -1,	AC	744
0.0 415 = 1	0.0 V250 10	0.6	.44	P • ₩	3.0	O _I n Solar ci	0.0 ELL COVER	AD AD	745
10.0	3.1	0.25	-1,	•1,	•1,	.00242	2,45	AD	747
4150 L	v250 16	•	, 0.5	. 45	٥,٥	0 t 0	0.0 IN PORT	AD AD	74 8 74 9
2,5 0.305	1.07	J.\$ J.75	-1. 0.75	•1, 0,15	-1. 1.00	0,98 0.0	2,72	AC AC	750 751
N184 1	4254 16 2,075	0.51	-1	-	4	0.004	UV PORT	AC	752
0.0	C	0.0	-1. .05	•1,	41, ,85	.0026 .25	1,4585	AD AD	753 754
N150 1	-250 10 .28	. 67	-1.	•1.	-1.	.00192	TEFLON	AC AC	7 55 756
0. N150 2	425= 1e	£ •	٥,	٨.	9.	0,	0. GOLD	AD.	757 758
v . 1	0.1	n.c2	3.90	9.607	-1,	0,606	•1.	AC	759
C, 1,0	9.26	8.04	°, °,85	0,607	-1,	0, -1.	0.	AD AD	760 761
0: N15: 2	i. ∷25= 16	ο.	≎.	r•	U e	0. BLACK	CCATING	AC AC	762 763
10,6	0.64 0.	0. 65	0.05	r. -1 .	-1.	.0572	0.00	AC AC	764 765
03,0	₽.89	1.77	0.02	. •1·	-1.	-1,	0.00	AD	766
N150 2	₹. :25= 16	U ,	0,	Α,		.0. HHITE	CCATING	AD.	767- 768
10,6	C • 12	C.93	0,05	-1.	-1,	.0067	0.00	AE	769
9016	0.21	(.45	0.02	°,	•1,	-1.	0,00	AD AD	778
416e		C	G. 	0.	0.	- · · · · · · · · · · · · · · · · · · ·	0.	AD	772
10.	-1.	.7	-1.	•1.	-1.	.000299947	1,78	AD	774
N1501-	.0668 H2807- -	. 	.0009	.0006	· · -	01	0,	AD.	775 774
560,C	•1.	-1.	-1.	0.0755	-1.	,00239		AD	777

Figure D-21. Part B

859.

3600

(10) Exposure Sequencing

The next data on ADATA are the controls for the exposure sequences to be computed. There are two reads; the first, NPIS, is the total number of cycles to be run, where this number includes initial conditions, assigned conditions, and both space and plume exposures. Subsequently, the details of the exposure sequences are read in.

Variable	Format	Comment
NP1S (Read in 0, 0)	5X, I5	Maximum value allowed is 10
NPSEQ(I, J), J = 1, 3 (read NP1S times in 0, 0)	3(10X, F5.0)	J = 1 is ID number of cycle J = 2 is type of exposure, EXTYPE J = 3 is exposure duration for plume on space exposure cycles = TØTIME

Figure D-22 shows the values associated with EXTYPE.

The final data controls preparation of a punched deck which contains the latest version of the array EFFECT. This array contains the information calculated in the program describing the state of the satellite surface after the last exposure cycle. This deck of punched cards can be used as the file CDATA (TAPE9) and when input in this fashion, the information can be used as the assigned initial condition prior to additional exposures.

Preparation of the punched deck is controlled by the flag NEFPN.

<u>Variable</u>	Format	Comments
NEFPN	10X, 15	Zero inhibits punching . 1 or greater causes punching. Normal setting is 1.

The data currently used for the control input reads, described above, are illustrated in Figure D-23.

Following this last data input from file ADATA = TAPE16, the routine DRØPIT is called to return the file and release the disc space it occupied. Note that this action does not release the core space assigned to the FET (File Environment Table) or to the buffer for the file, thus the core requirements for the program are unchanged.

j. File INDATA = TAPE5

This file is used to supply properties of $N_2\mathcal{O}_4$ and MMH. The INPUT1 editor is used, which requires special formatting of the data cards. The following information explains the requirements for using INPUT1.

(1) Description of INPUT 1

INPUT1 clears a specified area of core and reads REAL or INTEGER numbers from INPUT1 cards into this storage area.

EXTYPE DECLARES TYPE OF EXPOSURE EXTYPE = "1 INDICATES END OF COMPATYPE = 0 INDICATES INITIAL CLEXIVES = 1 INDICATES ASSIGNED CL

EXTYPE

ASSIGNMENT OF EXPOSURE SEQUENCES

Figure D-23. ADATA Input of Program Sequencing Control Parameters

(2) Use

CALL INPUT1 (X, Y, Z) where:

- X is the first element of an array where the input data is to be placed; DATA(1) in <u>SURFACE</u>.
- Y is the last element of the above-mentioned array; DATA(202) in <u>SURFACE</u>.
- Z is the array dimensioned large enough to hold any data which goes into X through Y. This Z array will hold the reference run data. This is DATBAG in SURFACE.

The arrays X and Z must be dimensioned at the start of the program. The dimensions of Z are not mentioned in the call to INPUT1.

(3) Restrictions

- 1. Within each reference run (including the zero reference run), the case numbers must be in increasing order.
- 2. After a program calls INPUT1, no data may be input from the TAPE5 except by INPUT1.

(4) Error Indications

X

All INPUT1 data cards will be examined. If an error is found which is clearly in a certain case, that case will be not executed; if that case number = 0, that is, it defines a reference run, then no case in that reference run will be executed. If the reference run or case number changes illegally, no case will be executed after the illegal change. This subprogram may not find more than one error per card.

The following error messages are possible:

MINUS PUNCH IN SECOND COLUMN OF FRACTION BAD PUNCH IN EXPONENT REF RUN NUMBER HAS CHANGED INCORRECTLY REF RUN AND CASE NUMBERS ARE BOTH ZERO NO REF RUN STORAGE HAS BEEN PROVIDED CASES NOT IN INCREASING ORDER BAD PUNCH IN FIRST COLUMN OF FRACTION LOCATION IS IN ERROR ILLEGAL CHAR IN DATA BELOW

(5) Method

Data numbers to be read by INPUT1 must be on cards punched as shown in Figure D-24.

Each card must have a one punched in column 1, and the reference run and case numbers punched in columns 69-73. Punches in columns 66-68 and 74-80 are not examined. Each card has four fields, and data numbers

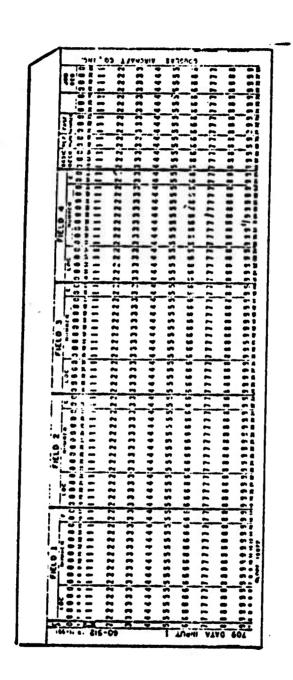


Figure D-24. INPUT1 Input Card Format

may be entered one per field. A completely blank field is ignored, while a field used to enter a data number must contain no blanks. Each field consists of three subfields:

- 1. LØC,
- 2. NUMBER
- 3. E

For purposes of explanation assume i is punched in the LQC subfield, n is punched in the NUMBER subfield, and e is punched in the E subfield. The number i denotes which element of the x array is to be set equal to the data number in this field. (This is X, the first INPUT1 argument.) Thus i is a subscript of X, so i must be greater than zero and equal to or less than the subscript associated with the INPUT1 argument Y. The only acceptable value of e is XX or a number in the range $-38 \le e \le 38$. If e is XX, the data number in this field is interpreted as an INTEGER whose value is n. If $-38 \le e \le 38$, the data number in this field is interpreted as a REAL whose value is n times 10° . (Here n is interpreted as though a decimal point preceded the first digit.) All other values of e are unacceptable and will result in error printouts. Either e or n, or both may be negative numbers. The minus sign is indicated by punching both a minus punch and a number punch in the first column of the subfield.

The data supplied to the INPUT1 editor were first prepared for an early version of the <u>TCC</u> subprogram, and contained a number of entries pertaining to thrustor design and operation that are not required by <u>SURFACE</u>. The unneeded entries have been removed from the INDATA deck, but some of the spaces for these entries are still reserved in the array DATA.

The card images in INDATA are shown in Figure D-25.

The data supplied by the INDATA file are the following:

Item Number	Description	Units
9	External Pressure	psia
10	Surface Temperature	· o K
	MMH Properties	
109	Boiling Point	°K
110	Freezing Point	°K
111	Critical Temperature	oK
112	Critical Pressure	psia
113	Vapor Cp	$cal/g_{-}^{o}K = Btu/lb_{-}^{o}R$

* # # # #	7 7 7 7	000000000000000000000000000000000000000	######################################
11	INI	01000	
	171	01000	
U (2.	00000	<u>::1332950cc03600013437830:0c00cc135295000363:0c01364609520900</u>
		00070	2929460606666300130262601060136013143200010000
			TO THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERT
•	11 X	1000	1.1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
11	7	61039	1111721005050505305118675063
•		00030	<u>11139956666660006611469000006000011500000000000011046674000</u>
"	100		179109306666366036630611022200006063691115940000600036011211990000004
~ •	2	01000	

Figure D-25. INDATA Cards for INPUT1 Edited Input

Item Number	Description	Units
114	Liquid Specific Heat	$cal/g-{}^{o}K = Btu/lb - {}^{o}R$
115	Not used	
116	Molecular Weight Vapor	g/g-mole = lb/lb mole
117	Latent Heat of Evaporation at Boiling Point	cal/g
118	Latent Heat of Fusion at Freezing Point	cal/g
119	Liquid Thermal Conductivity	cal/sec-g- ⁰ K
120	Vaporization Accommodation Coefficient	none
121	Reference Temperature for Physical Properties	°K
122	Density, liquid	g/cc
123	Viscosity, liquid	poise
124	Surface Tension, liquid-air	dyne/cm
	N204 Properties	
129	Boiling Point	°K
130	Freezing Point	°K
131	Critical Temperature	°K
132	Critical Pressure	psia
133	Vapor Cp	$cal/g^{O}K = Btu/lb^{O}R$
134	Liquid Specific Heat	$cal/g^{O}K = Btu/lb^{O}R$
135	Not used	
136	Molecular Weight, Vapor	g/g mole = lb/lb-mole
137	Latent Heat of Evaporation at Boiling Point	cal/g
138	Latent Heat of Fusion at Freezing Point	cal/g
139	Liquid Thermal Conductivity	cal/sec-g-OK

Item Number	Description	Units
140	Vaporization Accommodation Coefficient	none
141	Reference Temperature for Physical Properties	°К
142	Density, liquid	g/cc
143	Viscosity, liquid	poise
144	Surface Tension, liquid-air	dyne/cm
	MMH·N ₂ Ø ₄ ·H ₂ Ø Deposit	
185	Density	g/cc
186	Specific heat of gas evolved when deposit is pyrolyzed	$cal/g^{O}K = Btu/lb^{O}R$
187	Latent heat of pyrolysis	cal/g
	Operations	
201	Default pulse duration	sec
202	Default time slice width	sec

k. File BDATA = TAPE8

This file contains the assignment of structural materials to specified satellite segments, and their initial surface condition. The data are assigned to the array SURDES (175, 66) corresponding to the segment ID numbers. (See Section D. 4.i for initial input to SURDES.) The new data are located in SURDES in the following locations

SURDES (M1, 12) = Code number of material into player

13 = Thickness of top layer

14 = Surface finish of top layer

30 = Code number of material in second layer

31 = Thickness of second layer

32 = Surface finish of second layer

Positions 48, 49, and 50 are corresponding locations for third layer data, but none are currently input. Later in the computation, condensate or deposits form the top layer and the structural materials are pushed down one layer. The material ID numbers correspond to those given in Figure D-26 and in array ALNAME.

4.

3/0 127 3/0 128	1 0/2	10 1	3/0 1	3/0 1	29 3/0 1	2.075	31 370 1	2 3/0 1	3/0 1	/0 13	81 0/8	A1 0/8	41 0/8	At 0/8	3/0 14	1 0/	3/0 14	#T 078	3/0 14	10 14	10 07	19 19	15 07	1 07	
PLUME CONST.CODE			2 NZDA AD	TEX	H HYDRAT	H NITRAT	NIT. AG	188	AMMONIA	.0	Z412	LEZS NITANON.	ED ABNO	0	NCI	HÇLIVOKAZI	0	CARBON	7	NH 4 CL SHSD	4102	I.	2 2 2	ž	MINE TON
MATERIAL CODE		3	3	OLA!	P PO	<u> </u>	EFLO	9 970	ž	THITE COAT	ZALAY	Z													

Figure D-26. Material Code Assignments

Variable	Format	Comments
None (Read in 3,0)	(//)	Skips the file leader cards
SURDES (M1, 12), M1 = 1, M1S	20F4.0	Material code input. A value of -99, will terminate the input.
Unnamed	(R)	Skip the -99, if M1S values found, Repeated after every SURDES read.
SURDES (M1, 13), M1 = 1, M1S	8E10.7	Thickness
SURDES (M1, 14), M1 = 1, M1S	16F5.0	Finish
SURDES (M1, 30), M1 = 1, 9	20F4.0	Second layer material. To save file space, Ml values
SURDES (M1, 30), M1 = 41, M3S	20F4. 0	are limited to those segments which have second layers.
SURDES (M1, 31), M1 = 1, 9	8E10.7	Second layer thickness
SURDES (M1, 31), M1 = 41, M3S	0110.1	become layer threehess
SURDES (M1, 32), M1 = 1, 9	16F5.0	Second layer finish
SURDES (M1, 32), M1 = 41, M3S	101 3, 0	becond layer linish
KTEST	5X, I5	Flag to control further input of segment materials KTEST = 0: no further data KTEST 1: more data, number of inputs = KTEST
MZ3, MZ2, MZ1, (SSR(MZ), MZ = 1, MZ2) (Read in 3, 0)	3(5X, 15) 5G10.0	

The function of KTEST and the variables MZ1, MZ2 and MZ3 are summarized in Figure D-27.

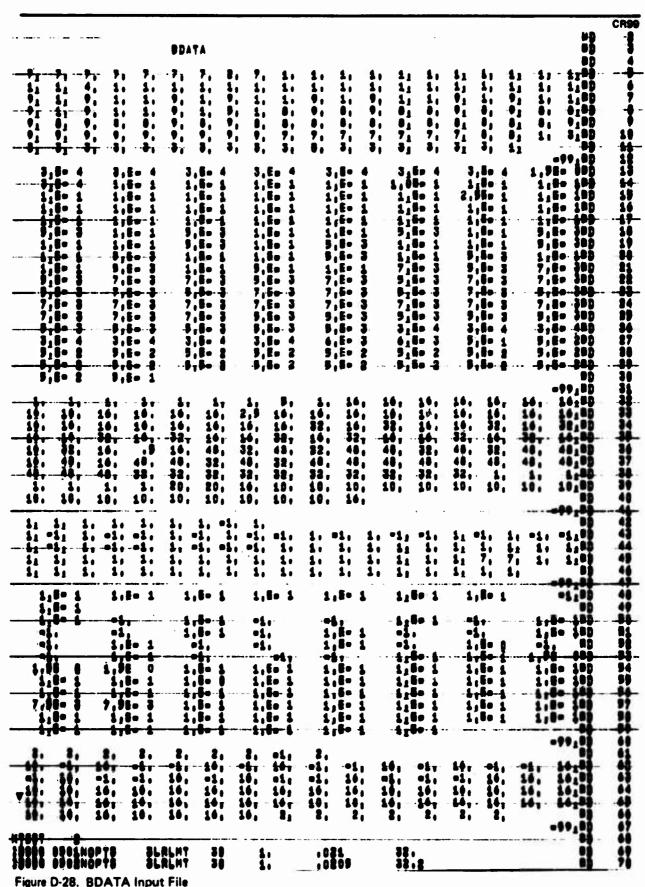
After appropriate tests and modifications of indices, SURDES (MZ3, n), where n is a function of MZ1 and MZ2, is set equal to SSR (MZ).

The current input supplied by BDATA is shown in Figure D-28.

1. File CDATA = TAPE9

This file is used to supply descriptions of the surface state coresponding to exposure to plume or space. It inputs such information as deposits, thickness, surface finish, temperature, etc., which are usually calculated by the program. These special inputs are then used as the basis for calculating the final surface condition--optical and thermal coefficients, etc. The input can be derived from previous <u>SURFACE</u> runs, from test data, or from other sources. This input can be used for a demonstration calculation or can be used as the basic state prior to further exposures. This selection is controlled by the flag ISTST, input either on NAMELIST NAST5 or later in file CDATA.

Figure D-27. Operation of Flag KTEST



igule D-20. DDATA iliput i lie

The data on CDATA is read into the array EFFECT (175, 15). The assignment of specific variables to locations in EFFECT is detailed in Figure D-29.

Variable	Format	Comments
None (Read in 4, 0)	(//)	Skips the file header cards
IIS, I2S	5X, I5, 5X, I5	IlS is the number of affected segments I2S is the number of data points for each segment.
EFFECT (I1, I2) I2 = 1, I2S Do I1S times (Read in at 4, 0)	5G10. 6/3G10. 6	If I2S is less than 9 file will not read properly
ISTST (Read in 10, 2)	10X, I5	Flat to select whether surface condition calculated from assigned input is used as basis for further calculations, or whether original pristine input is used.
		ISTST = 0; use original conditions
		ISTST 1; use conditions resulting from assigned state input.

The complete CDATA file is shown in Figure D-30.

m. File DDATA = TAPE10

This input shown in Figure D-31, contains the primary data defining the operating conditions for plume exposure and also some data used to demonstrate the capabilities of the coordinate transformation routines.

Variable	Format	Comments
None (Read in 5,0)	(//)	Skips file header cards.
DELTIME	10X, F10·0	Time increment for thrustor calculations
DELP or DUM	5X, F10·0	DELP = Change in latus rectum, between time increments during startup and tailoff. If DELP already input, this read is to a dummy variable.
NTR, P	10X, I5, 10X, F10·0	NTR is thrustor ID number P2 is 1/4 latus rectum

Figure D-29. Locations in EFFECT for Various Surface Conditions

C

			CDATA						
\$25 1.600 1.57060 52. 650. 1. 0.095 250. 1. 0.095 250. 1. 0.00 1.57060 52. 650. 1. 0.095 250. 1. 0.095 250. 1. 0.00 1. 0.096 250. 1. 0.00 1. 0.096 250. 1. 0.00 1. 0.096 250. 1. 0.00 1. 0.096 250. 1. 0.00 1. 0.096 250. 1. 0.00 1. 0.096 250. 1. 0.00 1. 0.096 250. 1. 0.00 1. 0.096 250. 1. 0.00 1. 0.096 250. 1. 0.00 1. 0.096 250. 1. 0.00 1. 0.096 250. 1. 0.00 1. 0.096 250. 1. 0.00 1. 0.096 250. 1. 0.00 1. 0.096 250. 1. 0.00 1. 0.096 250. 1. 0.00 1. 0.096 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0.005 250. 1. 0	4,325	-	700	42.	.009	-	\$60.0	1001	1000
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Figure D-30. CDATA Input File

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	4 N			•	•		88	22
8 ZZ 6ZZ	2.5					 0,09	888	133
SOP: O	70000		95.0	*			3222 3222	14434

Figure D-31. DDATA Input File

		
		These data are used to calculate plume-satellite intercepts. Several sets can be calculated, because the program returns and reads new NTR and P until a value of 999 is found in NTR.
ITRNCL	10X, 15	ITRNCL is a flag which selects which coordinate transformation routines will be computed.
		ITRNCL = -1: Satellite to thrustor
		= 0: None
		= 1: Thrustor to satellite and satellite to thrustor
		= 2: Thrustor-to- satellite only
		If <u>SURFACE</u> is computing the effects of a second plume exposure, ITRNCL is automatically set to -1. and the read is skipped.
N, NR	3(10X, I5)	This input supplies the thrustor number (NR) and the number of data sets to follow (N) for the calculation of thrustor coordinate system based coordinates
ZZ, RR, THE (read in 5,0)	4(10X, F10·0)	This input supplies demonstration data consisting of coordinates of locations in the thrustor-based coordinate system. The values are used to calculate the coordinates of the same locations in the satellite-based coordinate system. The read is only made if ITRNCL is positive, and it is made N times.

Format

Variable

Comments

After the data set for ZZ, etc., is exhausted, the program returns and reads N, NR again. This sequence continues until a value of -1 is read for N. No further data are read from DDATA on the plume exposure currently being computed.

For each additional plume exposure cycle, four more cards will be read; the inputs required are DELTME (one card), DELP (one card), NTR and P (one card), and a NTR = 999 card to end the input for the cycle.

The copy of file DDATA in Figure D-31 has input for two cycles; the second cycle input starts at card DD. 16.

n. File EDATA = TAPE11.

File EDATA is used to supply information about the characteristics of the thrustor plume at the locations where it impinges satellite surfaces. The data in the file were selected from typical output of subprograms KINCON and MULTRAN as recorded in the 1972 final report on CONTAM.

The <u>SURFACE</u> program needs one set of plume descriptive data for each impinged segment for each time interval, and will accept any number of inputs up to that quantity. If fewer are input the program will use the inputs actually available for locations/time intervals which were not supplied.

Variable	Forr	nat	Comments
blank (Read in 12,0)	(//)		Skip past file heading
NPMCR (Read in 12,0)	10X, I5		Total number of sets of plume characteristics to be read in at this time.
IP1S, IP2S, IP3S (Read in 12,0)		3(10X, 15)	IP1S = Time slice number IP2S = Affected segment number in array EFFECT (not segment ID) IP3S = Number of individual
			data points. (This variable is not operative.)
PLMCHR (IP3), IP3 = (Read in 12,0)	1, 90	10F8.0 (9 cards)	Specific plume characteristics

A single set of 9 cards is first read in. Then the program checks to see if more data are to be available (is NPMCR greater than 1?). If not, the values of IPIS and IP2S are ignored and the array PLMCHR is filled for the

range IP1 = 1, N1; IP2 = 1, N2; where N1 = number of cycles and N2 = number of affect segments, generated from the values of PMLCHR and written on TAPE14. If more than one set are to be input, the program writes PLMCHR to the tape up to the input values of IP1S and IP2S. Then the program reads EDATA for new values of IP1, IP2, and PLMCHR. This cycle is repeated until a value of -1 for IP1 is found, which indicates that no more PLMCHR values are available. The current value is written out to fill the file.

The copy of file EDATA shown in Figure D-32 has two sets of input for the first plume exposure cycle, and a single set for the second cycle.

The variables that are assigned to specific locations in PLMCHR are given in detail in Section D. 4. g.

o. Summary of File Assignments in SURFACE

Functions of all the files assigned to SURFACE are summarized below.

Number (Internal)	Name (External)	Function
1	-	Storage for array MATERAL
2	-	Storage for array PROPTY
3	-	Storage for array CHEMIC
4	-	Storage for array SURDES
5	INDATA	INPUT1 data input
6	ØUTPUT (Line printer)	Listing results
7	-	Dummy file when listing not desired
8	BDATA	Standard data input (Formatted)
9	CDATA	Standard data input (Formatted)
10	DDATA	Standard data input (Formatted)
11	EDATA	Standard data input (Formatted)
12	-	Buffered storage for common blocks \emptyset ED and \emptyset EF
13	-	Buffered storage for common block R6
14	-	Storage for array PLMCHR. Generated in <u>SURFACE</u> or other <u>CØNTAM</u> subprograms.
15	CØUNT	Buffered storage of count of calls to over- lays and subroutines.

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8	ED.	8	20			В		<u> </u>		~	04		2	15	39	0	ED H	9	1	0	F. 12		ש ה ה	ED	<u>B</u>	0.3	τ. π 3 Ξ
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Number (Internal)	Name (External)	Function
16	ADATA	Standard data input (Formatted)
17	INPUT (Card reader)	NAMELIST card input of changes
18	-	NAMELIST inputs from 17 are stored on 18 and read from 18
19	-	Buffered storage of array SEARCHD
20	FLG	Overlay (2, 1) (Object module)
21	FLG	Overlays (4,0), (10,2) (Object modules)
22	FLC	Overlays (5,0), (5,1), (5,2), (5,3), (5,4), (5,5), (12,0), (13,0) (Object modules)
23	FLE	Overlay (6,0) (Object module)
24	CHEEP	CHECKPOINT save of program
25	FLØ	Overlays (0,0), (1,0), (2,0), (3,0), (14,0) (Object modules)
26	PUNCH (Card punch)	Card deck of array EFFECT (Formatted)
30		Buffered storage of variables when program is to be stopped after calculating intercepts and input of same variables when restarted
31*	-	Communication with other subprograms in CONTAM about thrustor operation
32*	-	Inquiry to other CØNTAM subprograms about plume characteristics
-	FLA	Overlays (11,0), (11,1), (11,2) (Object modules)
-	FLB	Overlay (7,0) (Object module)
-	FLD	Overlay (10,0), (10,1), (16,0) (Object modules)
-	FLZ	Overlay (15,0) (Object module)
-	DAYFILE (Not named on ' Program Card)	Listing of file releases, RFL's and termination causes.

^{*}These two files are referenced in sections of the program which are currently not operational.

Figure D-3 is a copy of the Program Card showing the assignments by parameters. Three files are not mentioned on the Program Card. These are TAPE13, TAPE14 and TAPE 19. These three files are equivalenced by EQUBUFF to other files which have finished their functions at the times EQUBUFF is called.

p. Operation of SURFACE

In the following sections, it is assumed that <u>SURFACE</u> and its auxillaries are stored in a master file on magnetic tape or in disc Permanent Files.

(1) No Modifications/Corrections

This is the most straightforward method, with defaults in their standard conditions, and all that is required is to copy the needed individual program and data files from the Master File. The program will most likely be executed in this fashion for test or demonstration runs. Of course, the punched data card input for STPFLG must be supplied. The deck to operate in this manner is shown in Figure D-33. The example is based on the assumption that <u>SURFACE</u> is stored on a magnetic tape file. If, instead, it is stored on a disc Permanent File, the REQUEST card shown should be replaced with an appropriate ATTACH card.

The cards marked with circled numbers are included for certain special reasons:

Number \bigcirc , COPYSBF the first file to listing output, is included so the user will obtain the Table of Contents to the multi-file master file for more complex operations in the future. If this listing is not wanted, the COPYSBF card can be replaced with a SKIPF, A, 1, 17, B card.

Number ②. The file named PX4 consists of a small program to print a count of the calls made to each of the overlays and subroutines during execution, and a statement of what routine was being processed when execution stopped. It uses the information stored on File CQUNT by Subroutine A of SURFACE. PX4 is designed to be used after an EXIT card so that it is only called if the run is aborted for some reason. A similar routine included in SURFACE is called when normal termination occurs.

The advantage of the count of calls to subroutines and overlays is that it is easier to determine what part of the execution had been reached. The normal error statements on the CDC computers will tell at what address an error occurred, but if the address is in a routine called 100 times, it is often difficult to track the cause of the problem when it is not known at what stage in the execution that the call occurred. If use of PX4 is not desired, the COPYBF can be replaced by SKIPF, A, I, 17, B. and the short deck after the EXIT(S) card can be omitted. Figure D. 34 is a typical output from PX4.

Number 3. The file RØMTM contains a small program which prints the total CP and PP times used prior to initiating the execution of the main program. It then calls in the main program for execution. The first action in the main program is to set internal clocks and again give the totals for CP and PP time. It is sometimes found that there is an appreciable time interval between the two sets of CP and PP times. If use of RØMTM is not desired, the CØPYBF should be replaced by a SKIPF card, the LQAD of

```
JOH CARD.
               IFL = 10000, MAXFL = 275000.
REQUEST, SUR, HY.
                           (NUMBER)
 PEWINDIGUR.
COPY, SUR, A.
RETURN, SUR,
REHIND, A.
+COPYSRF, A, OUTPUT,
                                   TABLE OF CONTENTS
SKIPF, A, 3, 17, B.
SOPYBFTATABATAT
COPYSF.A.BDATA.
COPYRF.A.CDATA.
•COPYBE,A.DDATA,
COPYBF.A. EDATA.
COPYBE, A, INDATA,
<del>8K1PF,A,B,17,B,</del>
                                   LUAD HODULE(ROM)
COPYBF, A, LGOPR,
                                   LOAD MODULE (ROM)
COPYBF.A.ROMTM.
                                   LOAD MODULE (ROM)
RETURN, A.
REWIND, ADATA, BOATA, CDATA, DDATA, EDATA, INDATA, COURT,
<del>RFL+275000,</del>
SETÇORE,
LOAD, LGOPR,
NOBO.
BETCORE,
LOAD, ROMTH.
NOGO,
RETURN, LGOPR, RCMTM,
REDUCE.
RFL,60000.
<del>revindicount.</del>
PAP.OFF.
SETCORE,
REDUCE.
                       END CF RECURD
            EXECTION YOU CECK
                0
7/8/9
                       END OF RECORD
                   END OF JOB/FILE
4/7/8/9
```

Figure D-33. Control Cards to Run SURFACE From File Copy with No Modifications

UARCO FINAL

OVERLAY (1+0)	ENTERED	1	TIMES.	
OVERLAY(2.0)		1	TIMES.	
DVERLAY (3.0)			TIMES.	-
OVERLAY(4+0)		1	TIMES.	
OVERLAY 5.0	ENTERED	1	TIMES.	
OVERLAY 7.0	ENTERED		TIMES.	
OVERLAY 11.0	The second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second secon	2	TIMES.	
OVERLAY 10+2		1	TIMES.	
OVERLAY 2.1	ENTERED	<u></u>	TIMES	
OVERLAY 14+0			TIMES.	
VISCEST	ENTERED	2	TIMES.	
WEBBER	ENTERED	2	TIMES.	
ZES	ENTERED	3	TIMES.	
HCONT	ENTERED	65	TIMES.	
RETRHO	ENTERED	57	TIMES.	
RETROD	ENTERED	46	TIMES.	
RETVIS	ENTERED	44	TIMES.	
OVERLAY 11+2		148	TIMES.	
OVERLAY 11.1		5	TIMES.	
HOTPAR	ENTERED	5	TIMES.	
INTER	ENTERED	80	TIMES.	
INTRA	ENTERED	291	TIMES.	
MAXSUR	ENTERED	331	TIMES.	
MINSUR	ENTERED	319_	TIMES.	
OVERLAY (5+1)		3	TIMES.	
OVERLAY (5+3)		1 3	TIMES. Times.	
OVERLAY (5.2)	ENTERED		TIMES.	
WARD	ENTERED	551	TIMES.	
OVER	ENTERED	6960	TIMES.	
SUB A	ENTERED	1	TIMES.	
SUB C	ENTERED	zi	TIMES.	
SUB LT	ENTERED	59	TIMES.	
SUB TRANS	ENTERED	44	TIMES.	
SUB TK	ENTERED	311	TIMES.	
SUR	ENTERED	551	TIMES.	
SUB LE	ENTERED		TIMES.	
ALTER	ENTERED		TIMES.	
FORM	ENTERED		TIMES.	
INPUT1	ENTERED			
J		_		
CRETURN AFTE	R A CALL T	O ANOT	HER ROUTINE	18
NOT INCLUDE	D IN THE C	DUNT D	F ENTRIES)	
LAST ROUTINE	ENTERED W	AS OVE	RLAY 5.0	
PENULTIMATE	ROUTINE WA	S SUB	C	

Figure D.34. Calls to Subroutines and Overlays Listed by Auxillary Program PX4.

RØMTM at Number 4 should be omitted, and the card TP1 at Number 5, which initiates execution, must be replaced by a card reading "FLØ." The execution initiation cards mentioned (TP1 and FLØ) are the file names on which the principal Overlays reside. They are defined on the cards QVERLAY (file name, 0,0).

Number 6. After loading the programs, it is advisable to RETURN the program RØM's because they are no longer needed and they are simply using up disc storage space.

(2) SCOPE Card Changes to Default Operations

There are two changes in operation which are controlled by the addition of SWITCH cards in the SCOPE control card deck. If the card SWITCH, 1 is inserted, NAMELIST input will be expected from the Input card deck. If SWITCH, n., with "n" having any value from 3 to 6 is inserted, a Checkpoint-Restart tape file will be made at periodic intervals. It is necessary to supply a magnetic tape named CHEEP for the checkpoint file. Both SWITCH cards can be included, but be certain that two SWITCH cards with values greater than 1 are not present.

The cards illustrated in Figure D-35 are typical examples. The three cards indicated by Number (7) will turn on both the checkpoint option and a read of NAMELIST input. They should be placed in the control deck (Figure D-33) at the location marked with a Number (7A) in that figure. Typical NAMELIST cards are shown at Number (8) of Figure D-35, these should be inserted at location Number (8A) in Figure D-33.

The particular Checkpoint SWITCH, 5, card will cause the Checkpoint file to be written after completing five ØVERLAY calls, and then after every subsequent six ØVERLAY calls. The NAMELIST cards illustrated will prevent calculation of space exposure effects by overriding standard input, and cause evaluation of average solar absorptivity and thermal emissivity on the solar cell array, a location which is normally bypassed in the evaluation.

(3) Modification of the Program

When modifications to the program are required, such as turning off portions of the Output listing by assigning IF6 = 7, the UPDATE processor should be used, with the QLDPL drawn from the third file in the program master file. Unless the modification are very extensive and involve a large fraction of the 62 decks which make up the QIDPL, the "Selective" ("Normal") or the "Quick" mode should be used. The decision between "Selective" or "Quick" modes should be made on the basis of whether or not the NEWPL generated during UPDATE processing is to be saved. If it is to be saved, the "Selective" mode should be used. In either mode, only those decks actually modified will be written to the COMPILE file by UPDATE. The COMPILE file should then be compiled by the RUN compiler, and the resulting ROM of the changed routines merged into the current ROM by the COPYL processor.

If the changes include introduction of a new program unit (external function, subroutine, or overlay), COPYL will not function successfully, and COPYN must be used.

HS24 - 125, PRUNH -6H50LARY (RESERVE) By SNAST1 IPXTS1 = 05 REQUEST, CHEEP, HY, CK.

Figure D.35. Additional Control Cards to Initiate a Checkpoint File and Namelist Input

The control card listing in Figure D-36 will modify a few cards in two routines, prepare a new program library and a new RQM, and then generate a new master tape. The LISTIT routine is used to check the contents of the new tape.

In the series of operations controlled by the deck shown in Figure D-36, the following points may be noted. Number 1 The new tape file is called using the RE\$ERVE parameter with the \$ for S, and it is not made permanent by CONFIRM until after the LISTIT is performed successfully at Number 2.

Number 3 The file Table of Contents is updated to reflect the dates at which the new versions of the program were prepared.

Number 4 The new Master File is first generated as a multifile disc file named STRCP and later copied to the tape. This reduces the number of times that the magnetic tape unit is accessed. At facilities which do not restrict loading of magnetic tapes to the first steps of the job, the request for the new tape can be delayed until Number 5, thus leaving a drive unit available for other users.

Number 6. SURFACE should be compiled with minimum length buffers. The RØM supplied by MDAC was compiled with all tape buffers set at 1001g words. If larger buffers are desired, the whole program must be recompiled, and the load and execution field lengths must be increased by 30000g for each 1000g increase in buffer size.

Number 7. The load module or RØM generated at this operation has the same field name (fid) as the one described in Section D. 4. b. (1) above (see Figure D-33), and can be called into load and execution in the fashion illustrated. However, both PX4 and RØMTM would have to be copied from Master File A by the additional control cards Number 8 and 9 in Figure D-36, which should be inserted at locations Number 8 and 9A. The job deck which includes the STPFLG card will then follow the last UPDATE job deck.

Figure D-37 is the data on the structure of the ØLDPL of <u>SURFACE</u> when the latest version was generated. Figure D-38 is the output from a LISTIT of the master file for SURFACE.

(4) Modifications to DATA Files

This was discussed in Section D. 4.

(5) Temporary Replacement of DATA Files

If a completely new deck of data cards equivalent to one of the DATA files is to be used, but if the use is for only a few runs and the original file is not to be replaced, the DATA file from the master is not copied, and the COPYBF card is replaced by a SKIPF and a COPYBR, INPUT card which causes a deck to be read in from the input stack. Warning, do not use COPYBF on the input stack unless all cards up to the end-of-job 6/7/8/9 are to be copies to a single file. For example, if a replacement file for DDATA has been punched by a previous run of the program, then Number (11) in Figure D-33 would be replaced by

```
JDO CARD. IFLU1000. MAXI
ID GARD. (FACILITY OPTIONS)
RECUEST. RUR, MY. (RUI
MEGUEST. REMBUR, MY. (REI
REMIND. SUR, NEMBUR.
COPY, SUR, A.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 CRSS
                                                                                                                                                                                 HAXFLETOODS,
                                                                                                                                                                                      (AUMBER)
                                                                                                                                                                                       (RESERVE)
                           RETURNAL PRESS THE TREES THE TREES THE TREES THE TREES THE TREE OF COLOR PROPERTY OF THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES THE TREES T
                                                                                                                                                                                                                                    PREES THE TAPE SRIVE
                                                                                                                                                                                                                                     TABLE OF CONTENTS
                                                                                                                                                                                                                                     OLDPL FOR T-0-C
(3)
                               COPYERF, STRCP, CUTPUT,
                                                                                                                                                                                                                                    TABLE OF CONTENTS FOR NEW TAPE
                           COPYOF, STRCT2, STRCP.

RETURN, STRCT1, STRCT2.

GOPYOF, A. OLDR1,

RFL, 40000;
                                                                                                                                                                                                                                    OLDPL FOR SURFACE
                              VPDATELO-RU, PROLERI, NOMENRI, COOMPN;

RFL, 10000.

SOPYOF, NEWRI, STRCP.

REPTOR, SHPN, STROP.

RETURN, OLDRI, NEWRI.
                                                                                                                                                                                                                                    NEMPL FOR SURFACE
                               THE OLD COMPILE FILES

DATA FILES AND PX4 (OLDPLE AND COMPILES)
                       PYSF, A, STRCP, 19,
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                                                                                                                                                                                                                                    LOOP HAS ONLY THE CHANGED ROUTINGS
                                                                                                                                                                                                                                    HERGE TO CREATE COMPLETE, MODIFIED ROM
                                                                  SURRESTION SET FOR TABLE OF SONTENTS -- UPDATE JOB DECK
                             ODELETE STREET
                                                                                                                         (NEWBATE)
                             •D 50
                                                                                                                                                                                                               (NEWDATE)
                             +1 ++
                                                                                                                                                                                                               (NEWDATE)
                             ·D 76
                                                                                                                                                                                                               (NEMDATE)
                             *$69U$NÇ$ STR
7/8/9
                                                                   SORREGISON OFF FOR SURFASE --- UPDATE -USB DECK
                            OF STREET
                                                                    107 . 7
                             -D 1/0.005
[F(TAR(NTR).LE.0) TRA(NTR) = SURDES(M1.3) + 0.79
                             */0/*
                             6/7/8/9
                         BKIPB.A.1.17.B.
                            COPYOF, A. ROMTH,
                      48K170, A, 1, 17, B,
```

Figure D-36, Control Cards for Modification of SURFACE

(1) CØPYBR, INPUT, DDATA. SKIPF, A, 1, 17, B.

and the new deck including a 7/8/9 card, would be placed at Number (11

(11A)

If the new deck is to replace the previous deck of the same name in the master file, UPDATE must be used to generate the program libraries (NEWPL and OLDPL) and the sequenced COMPILE file. For this case, the file named in the COPYBR should not be DDATA. Figure D-39 shows the appropriate set of cards for this operation. These also would replace Number (1) in Figure D-33. The UPDATE job deck directives in Figure D-39 must then be placed immediately following the new DDATA deck in the job stream, which is located at Number (11) (Figure D-33).

D. 5 TEST CASE

Subprogram <u>SURFACE</u> was executed independently of the <u>CONTAM</u> executive. Hence, the data for running the sample case were modified from that used for other subprograms. First: certain parameters were arbitrarily assigned. This was done for either of one of two reasons:

a. The parameter is not meaningful for other subprograms, and so was not used earlier.

Parameter	Program Variable	Assigned Value	Input Method
Active Thrustor Identification	NTR	2	NAMELIST/NAST4 Input of NTR1
Plume Exposure Duration	TØTIME	2 x 10 ⁻³ sec	NAMELIST/NAST3 Input of TØTIM1
Plume Exposure Time Increment	DELTIME ·	2 x 10 ⁻⁴ sec	NAMELIST/NAST3
Space Exposure Duration	тØтіме	6 x 10 ⁵ sec .	ADATA Input
Space Ambient Pressure	PE	1 x 10 ⁻⁶ psia	INDATA Input

b. The parameter is not specifically defined in other subprograms, so an estimated value was developed by inspection of the related output from the other subprograms.

SE SC SD HCN ERRY SERTION SE ST. ST. ST. ST. ST. ST. ST. ST. ST. ST.
SE EFF 1470 571
14/0
HCT WARD
15/0 NERE FOUND
2 2 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3
- W.
EXT
808
FILE
200
4 2
375
2
E E
1870

THIS UPEATE REDUINED 3410CE NORES OF CONE. Figure D-37. Structure of OLDPL of Surface (1 of 2)

UNLABELED ULDEL

PASTER AUDIT, IDERT CARD TOTAL

UPDATE V1.2

06/07/73 13.55.50-

PAGE 3

SUBMARY OF UPDATE IDENTIFIERS WITHIN DECK - TOTALS IDENTIFIER TOTAL ACTIVE

IDENTIFIER	TOTAL	ACTIVE
YALKSER		
C/C	1342	1342
36	19	18
	119	118
SD.	29	29
<u> FCR</u>	33	
MHO	17	17
₩0₽ 	14	14
HAD	40	39
PHP	29	28
VI	137	135
NBP	245	241
4\$	19	
EXP	113	113
#FL 1/0	44	44
14/0	1555	1534
FRI.	15	15
1A	12	- 11
5/0	508	495
3/1	169	165
4/6	200	193
3/0	675	654
9/1		
17.	19	15
1/2		59 51
9/4	363	350
3/5	51	49
- 1/2	346	350
7/6	1391	148

PPI.	18	18
10/6	105	
-10/1	**	92
13/2	470	448
►DT	• • • • • • • • • • • • • • • • • • • •	9;
IEP		28
PAX	20	74
×10	22	21
EAR		
SUF THUS	21	21 90
NAKD	21	X
11/1	343	313
11/2	196	168
12/4		418
13/6 HA!	200	103
F01-	497	460
15/0	886	886
***		-11
171	288 166	286 166
	14116	13755
		-3177

Figure D-37. Structure of OLDPL of Surface (2 of 2)

4.

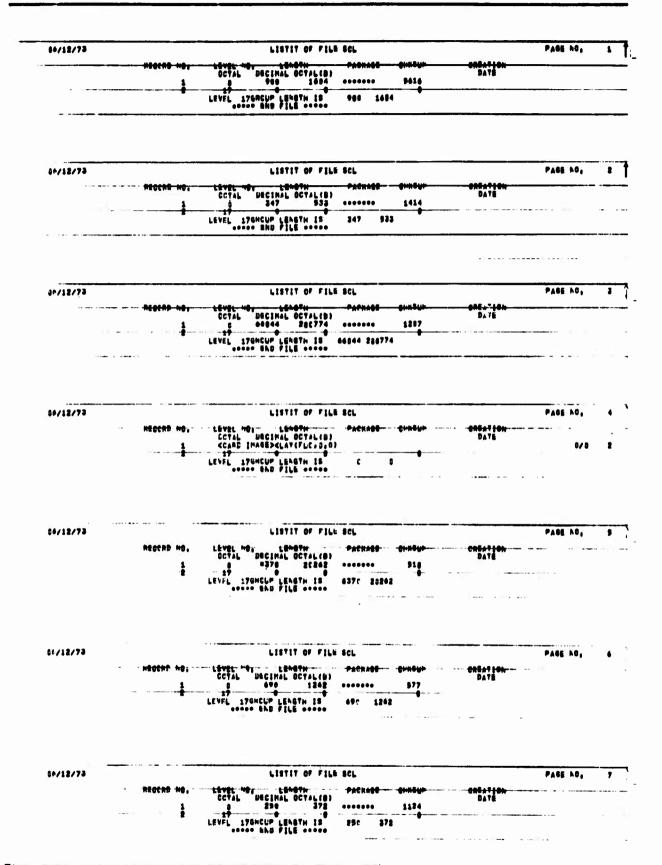


Figure D-38. LISTIT Output from SURFACE Master File (Page 1 of 5)

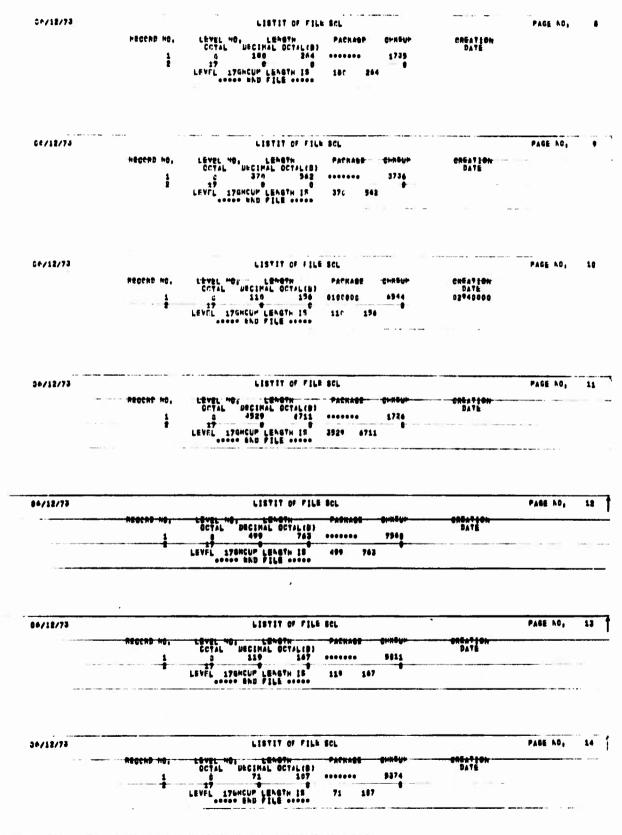


Figure D-38. LISTIT Output from SURFACE Master File (Page 2 of 5)

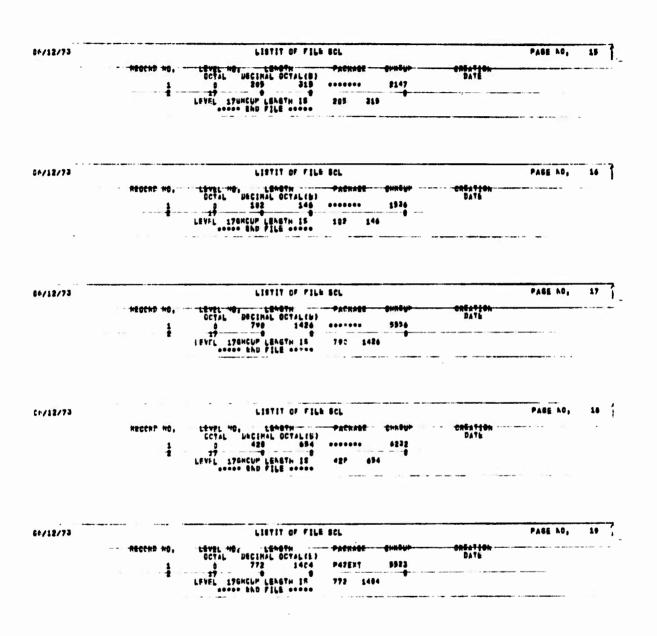


Figure D-38. LISTIT Output from SURFACE Master File (Page 3 of 5)



St/12/73		LISTIT OF FILE SCL		er - to er itte - stee te steen e ere seit	PAGE NO. 20 '
	HERFHE NO.	LEVEL HO, LEASTH PACHAGE CCTAL DECIMAL OCTALIED	CHEUP	CREATION DATE	
	1>	RC HAGE><(CVEHLAY(FLO.0.0) > 1 8881 4807 P1942	4397		
	3	0 02 122 A c 91 43 F	4432 7034		
	:	e 719 18+8 e 0 247 355 C	-907 9432		
	Ž	g 111 197 FCCAT C WG 180 RETHMC	3348 4118		
	1	0 40 132 RETHICE 0 102 146 RETVIS	4441 4875		
	11 12	1 144 - 200 RADA 1 171 FAULT	9446 4372		
	i	6 93 65 Lf	7752 4432		
	15	: 4597 ZC5A7 WEFEE	7945 272		
	17	. 47 197 694	9855	00/12/73	
	18	RE TAGES CVENLAY (FLO. 1.C) >	2376	06/12/73	
	20 21)	RE [HASE> <cvehlay(flo:14.0)></cvehlay(flo:14.0)>	1310		
	22 23	9 4946 19504 P42140 0 785 1895 18951	975 9767		
	74 2 1	0 04 100 FORE	4275 4440		
	76> 27	#D [MAGE> <cverlay(flo.p.g)> 1691 3273 F4220</cverlay(flo.p.g)>	5310		
	76) 29	RC [MAGE>< OVERLAY(FLG,2,1) >	1295		
	30> 31	RE [MAGE>(CVEHLAY(FLO,3,0) >	£141		
	32> 33	" FL 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	5324		
	34> 39	RE IMAGENCEVENTATIFLE (910)	2241		
	34> 37	OF THATES/CHELLASTEL C. S. 41	2847		
	34	43 193 19	5003		
	39> 40	RE [MAGE>CCVENLAY(FLG,9,2) > 130	2046		
	41>	AD 1448 PACE BLAY (FLC, 5, 3) > 146 822 14253	7755		
	43> 44	## IMAGE > C	8490		
	45) 46	RC [MAGE> <cyenlay(flc,5,5)> G 120 17C F4255</cyenlay(flc,5,5)>	3625		
	47> 48	0 1125 5145 P4260	5366		
	49 50)	## A44 A74 PATANT	2507		
	51	3667 7123 F4270 84 174 F603	7213 1122		
64/12/73		LISTIT OF FILE SOL			PAGE AD, 21
	HECCHD +0.	LEVEL NO. LENGTH PATRAGE	CHASUP	CREAT Oh	
	43	CCTAL DECIMAL OCTALIGN	2465	DATE	
	54>	#F [MAGE> <cv&hlay(fld,10,0)> 5 642 1204 P49100</cv&hlay(fld,10,0)>	4313		
	56> 57	#S IMAGE>COVERT AT (FLD +10 +11 > 2 06 108 P4F101	1197		
	58>	## 1 #################################	5612		
	40>	FE THABEPCCYBHLAV(FLA:11.8)	1227		
	41 42 42	G 200 321 EXTRAF	7442 7613		
	64 65	216 370 147FR	4453 4772		
	14 A A A 7	j 204 434 [PYFA 9 79 106 MADSUN L 74 112 MIPSUN	5776		
	≜A	7 206 414 FVER	31C1 44CJ	•	
	40 25	1 492 7.4 TEARS	9617 7607		
	71 72>	: 134 232 FARI RC Thagedecybellay (Flats): 13 3	4794		
	73 74>	AT IMAGENCEARMY (LTV 11'5) >	5503		
	75 76>	T PAGE > COVERLAD (FLG. 12.0) >	1475		
	77 7 4 >	C 1337 2471 P4;12; AD [MAGE>COVERLAN(FLC:13:11 >	6703		
	7 0 80	. 613 1145 P4713C - 1242 55:6 RAIDAL	142		
	#1 #2	a 1/6 2A3 KABDÝ : 1347 2565 MOTERT	1613 3243		
	P3> 84	FL IMAGE > COVERLAY (FLZ, 15, C) >	3150		
	P5 PA>	t 2444 4676 P42150 H4 174 FECS FL [FAGE>CEVERLATIFED-16.() >	1223		
	A7 68	n 108 274 \$47160	5577 C		
			•		

Figure D-38. LISTIT Output from SURFACE Master File (Page 4 of 5)

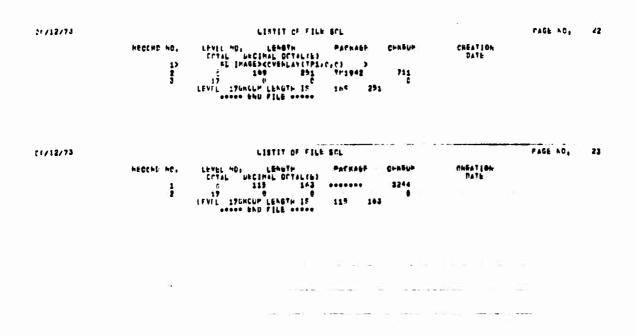


Figure D-38, LISTIT Output from SURFACE Master File (Page 5 of 5)

CR99

•DECK	DD	PDATE	JOB	DECK	FOR	INSERTION	OF	HEW	סמ
**EAD 7/8/9	INDD		·						

Figure D-39. Control Cards for Replacement of a DATA Deck

4 1

Parameter	Program Variable	Assigned Value	Input Method
Plume Shape Specification 1/4 Latus Rectum	P	1.0	NAMELIST/NAST4 Input of Pl
Plume Shape Change During Startup and Tailoff; Change in Latus Rectum	DELP	0.001	NAMELIST/NAST4 Input of DELP1

Second: Using the input data files described earlier with the above specific values, <u>SURFACE</u> was executed with STPFLG = 1. This resulted in execution proceeding only as far as calculation of the satellite surface segments which are impinged by the thrustor plume, and their coordinates in the thrustor-based coordinate system. These data were listed, current values of all variables were stored on a magnetic tape, and then the program stopped automatically.

The list of locations of impinged segments was then used as the basis of a manual search of <u>KINCQN</u> and <u>MULTRAN</u> output listings to find the plume characteristics at the locations and times corresponding to plume time slices. The method of determining these data is described in the main section of the description of the test case.

The plume characteristics data were then punched on cards in the correct format for file EDATA.

A second run of <u>SURFACE</u> was then made with STPFLG = 2, with the new EDATA file replacing the default data supplied with the program, and using the magnetic tape to supply values of variables which had been stored earlier.

In this run, computations started immediately after the location at which the previous run was halted.

The listings from the two runs are discussed separately below:

RUN WITH STPFLG = 1 FOR CALCULATION OF PLUME-SURFACE INTERCEPTS

The option to list all normally-generated output was left turned on (16F = 6); hence the listing is complete. In the illustrations given in this report, some redundant portions are truncated to save space.

Initialization and Timing (Figure D-40)

The first three statements (date, time prior to entry, and rollout warning) are generated in the auxiliary (0,0) overlay, Program TM42. The note about long operating times and rollouts is borne out almost at once; the excessive time for running (2, 1) is due to a rollout, see the section of the dayfile indicating a 14 minute rollout inserted to the left of the listing of the time needed for (2, 1). The remainder of the timing messages are generated in the main program. Overlay (2, 1) is used to input all the NAMELIST modifications to the data.

21/ 6/73 CR99A

CP TIME USED = 37.89 SECONDS PP TIME USED = 278.28 SECONDS SEPORE ENTRY TO PROGRAM

IF ANY OPERATING TIMES SEEM UNREASONABLY LONG CHECK THE DAYFILE FOR ROLLOUTS IN THE SAME CLOCK TIME PERIOD

MICKORY-DICKORY SAYS 8.88 SECONDS AT START OF 8.0 LEVEL ACTIVITY STARTING AT CLOCK TIME 8126128.8

> CP TIME USED = 37,98 SECONDS PP TIME USED = 275.83 SECONDS SINCE JOB STARTED

CHECKPOINT OPTION IS OFF

INSERTEL PROM DAYFILE

18.26.32.ROLLOUT COMPLETED. (PL 130180)
88.40.35.ROLLUN COMPLETED.

MICKORY-DICKORY SAYS 871.20 SECONDS FOR RUNNING OVERLAY 2.1 FINISHED AT CLOCK TIME 81411 0.8

> CP TIME USED 4 38.99 SECONDS PP TIME USED 0 278.90 SECONDS SINCE JOB STARTED

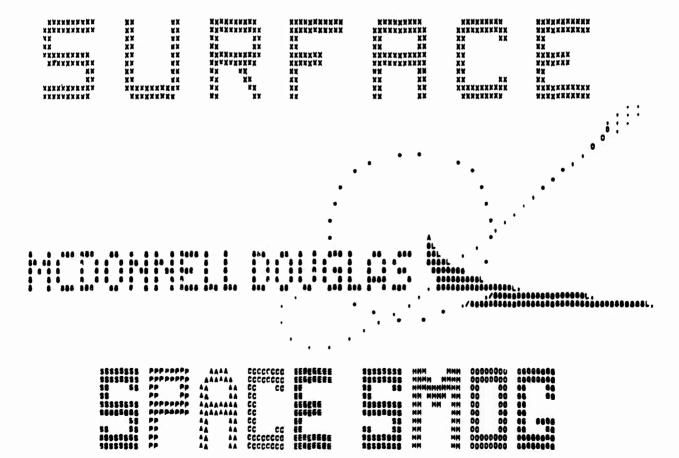


Figure D-40. Initialization Listing

Satellite Configuration Description

The next operation is reading in the configuration of the satellite used in the calculations. The data are read from file ADATA. Figure D-41 presents the input data as tabulations which consist of a description of the satellite exterior.

Note that the presence of 'projections' from the satellite surface is signalled by numbers other than -1. appearing in the column headed "Height of Projection" in the main table "Satellite Configuration." Following the main table, details are given about the projections; the details include the ID number of the segment to which they are attached (compare column 1 of the main table).

Properties of Structural Materials and Plume/Deposit Constituents

The properties of materials and constituents which will form the basis for calculation of exposure effects are next listed. The data were read from ADATA and INDATA. Approximations for data not supplied are performed after the inputs are completed, and the information given in Figure D-42 includes both input and approximated data. The listing is in the order physical properties of structural materials, phys-chem properties of plume/deposit constituents, wear rate properties of all substance, and optical and thermal constants for all materials. The last section of this part lists significant data which were not input and which could not be approximated.

Coefficients and Physical-Chemical Properties of Deposit Materials

The values for representative constants which are internally generated from the input data are listed. The listing is presented as Figure D-43. The listed values can be compared with available literature values for a check of the accuracy of the program computations. Flagrant variation can indicate errors in either the input data or in a computation module.

Data Revisions

The revised data supplied by means of NAMELIST input are listed, because the flag RDWRT is greater than zero. These revisions alter the values input from the data files or the defaults assigned in the program. Figure D-44 shows the revision inputs. Remember that these revisions do not affect data storage or execution paths until the point at which execution tests for their existence and then replaces the old values. Hence the other data file input listings still show the original values as supplied.

Assignment of Calculation Cycles

The basic set of calculation cycles, read from file ADATA, is next interpreted and listed. These data can be superseded by NAMELIST inputs, and in this particular case, they are (see section above on Data Revisions). The listing of calculation cycles is illustrated in Figure D-45.

the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the s

		6 H E	No. 7		K DIST			DIRFC	TION COSINES
••••					SEGMENT	SURFACE		OF	NORMAL TO
ID NO.	COORD.	COORD.	THETA A	1EA	TO PLUME VERTEX	TEMPER	PROJECTION	PROJE	CTION TANGENT PLANE
*****	(FT)	(#7)	(RAD) (90	FT)	(FT)	(DES R)	(FT)	LAMBDA	MU NU
1	0.000	0.000	0.00000	1	-3,600	520.0	-1.0	-1.0	-1.0 -1.0
3	.305	.590	1.04720	1	-3.295 -3.295	520.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0 -1.0 -1.0
4	.305	.590	5.23600	1	-3.295	520.0	-1.0	-1.0	-1.0 -1.0
5	.620	1.200	.62830 1.88490	1	-2.980 -2.980	520.0 520.0	-1.0	1.0000	-1.0 -1.0
7	. 620	1.200	3.14159	1	-2.900	520.0	-1.0	-1.0	-1.0 -1.0
•	.620	1.200	4.39818	1	-2,980 -2,980	520.0 520.0	-1.0	-1.0	-1.0 -1.0 -1.0 -1.0
10	1.325	1.000	.31416	1	-2.275	520.0	-1.0	-1.0	-1.0 -1.0
11	1.325	1.600	.94248 1.57080	1	-2.275 -2.275	520.0 520.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0 -1.0 -1.0
13	1.325	1.000	2.19911	i	-2.275	520.0	-1.0	-1.0	-1.0 -1.0
14	1.325	1.400	2.82743 3.45575	1	-2.275 -2.275	520.0 520.0	-1.0 -1.0	-1.0	-1.0 -1.0 -1.0 -1.0
10	1.325	1.000	4.08407	1	-2.275	520.0	-1.0	-1.0	-1.0 -1.0
17 18	1,325	1.000	4.71240 5.34070	1	-2,275 -2,275	520.0 520.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0 -1.0 -1.0
19	1.325	1.600	5.96902	1	-2.275	520.0	-1.0	-1.0	-1.0 -1.0
20 21	2.325	1.600	.31416	1	-1.275 -1.275	520.0 520.0	-1.0 -1.0	-1.0	-1.0 -1.0 -1.0 -1.0
55	2.325	1.000	1.57000	!	-1.275 -1.275	520.0	-1.0	-1.0	-1.0 -1.0
23 24	2.325	1.600	2.82743	1	-1.275	520.0 520.0	-1.0	-1.0	-1.0 -1.0 -1.0 -1.0
25	2.325	1.000	3.45575	1	-1.275 -1.275	520.0	-1.0	-1.0	-1.0 -1.0 -1.0 -1.0
26 27	2.325	1.000	4.08407	1	-1.275	520.0	-1.0	-1.0	-1.0 -1.0 -1.0 -1.0
85	2.325	1.600	5.34070	1	-1.275	520.0	-1.0	-1.0	-1.0 -1.0 -1.0 -1.0
20	2,325 3,325	1.600	.31416	3	-,275	520.0	1.5		APPLICABLE
31	3.325	1.000	.94246	1 3	275 275	520.0 520.0	-1.0 1.5	-1.0	APPLICABLE
32 33	3.325 3.325	1.000	1.57080 2.19911	i	-,275	520.0	-1.0	-1.0	-1.0 -1.0
34 35	3.325	1.000	2.82743 3.45575	1 3	-,275 -,275	520.0 520.0	-1.0 1.5	-1.0 NOT	APPLICABLE
36	3.325 3.325	1.600	4.00407	1	275	520.0	-1.0	-1.0	-1.0 -1.0
37 38	3.325 3.325	1.000	4.71240 5.34070	3	275 275	520.0 520.0	1.5	-1.0	APPLICABLE -1.0
39	3.325	1.600	5.96902	1	275	520.0	-1.0	-1.0	-1.0 -1.0
41	4.325	1.600	.31416	1	.725 .725	520.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0 -1.0 -1.0
42	4.325	1.000	1.57080	1	.725	520.0	-1.0	-1.0	-1.0 -1.0
44	4.325	1.600	2.19911	1	.725 .725	520.0 520.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0 -1.0 -1.0
45	4.325	1.000	3.45575	1	.725 .	520.0	-1.0	-1.0	-1.0 -1.0
46	4.325	1.600		1	.725 .725	520.0	-1.0 -1.0	-1.0	-1.0 -1.0 -1.0 -1.0
48	4.325	1.000	5.34070	1	.725	520.0	-1.0	-1.0	-1.0 -1.0
50	4.325 5.325	1.600		1	.725 1.725	520.0 520.0	-1.0 -1.0	-1.0	-1.0 -1.0 -1.0 -1.0
51	5.325	1.600	.74248	1	1.725	520.0	-1.0	-1.0	-1.0 -1.0
52 53	5.325 5.325	1.600		1	1.725	520.0 520.0	-1.0 -1.0	-1.0	-1.0 -1.0 -1.0 -1.0
54	5.325	1.000	2.82743	1	1.725	520.0	-1.0	-1.0	-1.0 -1.0
55 56	5,325 5,325	1.600		1	1.725	520.0 520.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0 -1.0 -1.0
57	5.325	1.000	4.71240	1	1.725	520.0	-1.0	-1.0	-1.0 -1.0
50	5.325	1.000		1	1.725	520.0 520.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0 -1.0 -1.0
60	4.325	1.600	.31414	1	2.725	520.0	-1.0	-1.0	-1.0 -1.0
61	4.325 4.325	1.000		1	2.725	520.0 520.0	-1.0 -1.0	-1.0	-1.0 -1.0 -1.0 -1.0
63	4.325	1.600	2.19911	1	2.725	520.0	-1.0	-1.0	-1.0 -1.0
65	6.325 6.325	1.600		1	2.725 2.725	520.0	-1.0 -1.0	-1.0	-1.0 -1.0 -1.0 -1.0
**	6,325	1.600	4.08407	1	2.725	520.0	-1.0	-1.0	-1.0 -1.0
67 68	6.325	1.600		1	2.725 2.725	520.0 520.0	-1.0 -1.0	-1.0	-1.0 -1.0 -1.0 -1.0
6.9	6.325	1.650	5.94902	1	2.725	520.0	-1.0	-1.0	-1.0 -1.0
70 71	7.325 7.325	1.000	.94248	1	3.725 3.725	520.0 520.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0 -1.0 -1.0
72	7.325	1.600	1.57080	1	3.725	520.0	-1.0	-1.0	-1.0 -1.0
73 74	7.325 7.325	1.600		1	3.725 3.725	520.0 520.0	4.0	.9820	.1000 .0500 -1.0 -1.0
75	7.325	1.000	3.45575	1	3,725	520.0	-1.0 -1.0	-1.0	-1.0 -1.0
76 77	7.325 7.325	1.600		1 1	3,725	520.0	-1.0	-1.0	-1.0 -1.0 -1.0 -1.0
78	7.325	1.600	5.34070	1	3,725	520.0	-1.0	-1.0	-1.0 -1.0
74 80	7.325 8.325	1.600	.31416	1 1	3.725 4.725	520.0 520.0	-1.0	-1.0 -1.0	-1.0 -1.0
-1	0.325	1.600	.94248	1	4.725	520.0	-1.0	-1.0	-1.0 -1.0
43	4,325 4,325	1.600		1	4,725	520.0 520.0	-1.0	-1.0	-1.0 -1.0 -1.0 -1.0

Figure D-41. Satellite Configuration Listing (Page 1 of 3)

••	6.325	1.000	2.02743	1	4,725	520.0	-1.0	-1.0	-1.0	-1.0
85	4,325	1.400	3.45575	1	4,725	520.0	-1.0	-1.0	-1.0	-1.0
•	4.325	1.600	4.08407	1	4,725	520.0	-1.0	-1.0	-1.0	-1.0
87	4.325	1.000	4.71240	1	4,725	520.0	-1.0	-1.0	-1.0	-1.0
	4.355	1.400	5.34070	,	4.725	520.0	-1.0	-1.0	-1.0	-1.0
**	4.325	1.600	5.76902		4.725	520.0	-1.0	-1.0	-1.0	-1,0
••	4.325	1.600	.31416		5.725	520.0	-1.0	-1.0	-1.0	-1.0
•1	9.325	1.000	.94246		5.725	520.0	-1.0	-1.0	-1.0	-1.0
*2	4.325	1.600	1.57000	ı,	5.725	520.0	-1.0	-1.0	-1.0	-1.0
•3	9.325	1.000	2.19911		5.725	250.0	-1.0	-1.0	-1.0	-1.0
94	9.325	1.600	2.02743	1	5.725	520.0	-1.0	-1.0	-1.0	-1.0
95	1.325	1.400	3,45575	1	5.725	520.0	-1.0	-1.0	-1.0	-1.0
**	9.325	1.000	4.00007	1	5.725	520.0	-1.0	-1.0	-1.1	-1.0
97 98	*.325	1,600	4.71240	1	5.725	550.0	-1.0	-1.0	-1.0	-1.0
	9.325	1.600	5.34070	1	5.725	520.0	-1.0	-1.0	-1.0	-1.0
•••	9.325	1.000	5.94902	1	5.725	520.0	-1.0	-1.0	-1.0	-1.0
100	10.325	1.400	-31416	ļ	4.725	520.0	-1.0	-1.0	-1.0	-1.0
101	10.325	1.600	.94248	1	4.725	520.0	-1.0	-1.0	-1.0	-1.0
105	10,325	1.600	1.57000	1	4.725	520.0	-1.0	-1.0	-1.0	-1.
103	10,325	1.600	2.19911	ı	4,725	520.0	-1.0	-1.0	-1.0	-1.0
104	10.325	1.000	2.02743		6.725	520.0	-1.0	-1.0	-1,0	-1,0
105	10,325	1.600	3.45575	1	6,725	520.0	-1.0	-1.0	-1.0	-1.
100	10.325	1.600	4.08407	1	0.725	520.0	-1.0	-1.0	-1.0	-1.0
107	10.325	1.000	4.71240	1	6,725	520.0	-1.0	-1.0	-1.0	-1.0
100	10.325	1.400	5.34070		6,725	520.0	-1.0	-1.0	-1.0	-1.0
109	10.325	1.000	5.96902	1	6.725	5.0.0	-1.0	-1.0	-1.0	-1.0
110	10.025	1.195	.52360	1	6.725	540.0	-1.0	-1.0	-1.0	-1.0
111	10.825	1.195	1.57000	1	6.725	550.0	-1.0	-1.0	-1.0	-1.0
115	10.025	1.195	2.61800	1	6.725	520.0	-1.0	-1.0	-1.6	-1.0
113	10.025	1.195	3.06520	1	6.725	520.0	-1.0	-1.0	-1.0	-1.0
114	10.025	1.195	4.71240	1	4.725	520.0	-1.0	-1.0	-1.0	-1.0
:15	10.025	1.195	5.75960	1	6.725	520.0	-1.0	-1.0	-1.0	-1.0
116	10.825	0.000	0.00000	5	6.725	520.0	2.5	NOT A	PPLICAL	ILE
NOT	E VALUES OF	F -1.0 1	HOICATE NO		NTERED. RWAL TOTAL	AREA =125.	OO SOUAPE	FEET		

PROJECTIONS

AERIAL AT SEGMENT NO. 5

••••	8	E G	H		E	N	-					X SEG				SURFACE
10	×					Ť										TEMPER.
NO.	CODR	0.	COO	RĐ								VE	RTE	X		ATURE
******) ====	• • •			-		•	(8		T) ===:	-	FT)			(DEG R)
501	.5	••	5	. 3	80	-6	.21	300		1		-2	. 95	•		529.2
205		41														529.2
. NEGAT	IVE V	ALUE	FO		ANG	LE	816	NAL	8 T	HA"	7 8	EGME	NT	18	ON	
SIDE O	F PRO	JECT	ION	•	ACI	NG	444	YF	ROM	T	HE	LUH	E.	• •		
	RFA D	FAE	RIA	L :	18	2	. 0	\$0 U	ARE	F	ET.	•	•••			••••
THRU	STOP	SEGMI	•••		18	•••	.0 Y (FT	•••	ARŁ	••	R (FT)	••••	 †	 HE T		••••
THRU	1870#	SE GMI	ENT		•••	•••	· · ·	•••	ARE	••		••••				••••
THRU	1870#	SEGMI NI	ENT O.		•••	••• •	Y (FT)	•••	•••	e (FT))		AAC)	••••
THRU	1870# 10.	SEGMI NI	ENT O.		•••	····	¥ (FT)		10	e (FT))	.31	410	0	••••
THRU N	1870# 10.	SEGMI NI	ENT O.		•••	7	Y (FT)	2.	100	PT:)	.31	#A0	0 0	••••

Figure D-41. Satellite Configuration Listing (Page 2 of 3)

3

SOLCEL AT SEGMENT NO. 73

	\$ E	6 # E	N T		¥ 0187	
10	1		THETA	AREA	SESMENT TO PLUME	SURFACE TEMPER-
NO.	COORD.	COORD.	144		VERTEX	ATURE
	(FT)	(FT)	(RAD)+	(SOFT)	(FT)	(DEG P)
••••						
7301		2.502	2.19911		3.244	550.0
7302	6.600	4.161	1.49900	1	3.000	560.0
7303	4.405	4.056	2.19911	1	3.003	560.0
7304	4.596	3.451	5.84400	1	2.998	560.0
7305	6.420	5.122	1.50100	1	2.420	560.0
7306	6.415	5.038	2.19911	1	2.815	540.0
7307	4.410	4,954	2.89700	1	2.410	540.0
7308	4.232	6.100	1.50300	1	2.632	540.0
7309	4.227	0.020	2,19911	1	2.627	560.0
7310	6.225	5.940	2,89500	1	2.622	560.0
7311	6.725	6.200	-2.89500	1	3,123	510.0
7312	6.718	6.119	-2.19911	i	3.118	510.0
7313	6.713	6.030	-1.50300	i	3.113	510.0
7314	6.911	5.235	-2.89700	i	3.311	510.0
7315	4.904	5.137	-2.19911	ĭ	3.306	510.0
7316	0.901	5.039	-1.50100	ī	3.301	510.0
7317	7.099	4.275	-2.89900	ī	3.499	510.0
7318	7.094	4.155	-2.19911	i	3.494	510.0
7319	7.089	4.035	-1.49900	i	3.449	510.0
7320	7.372	2.661	-2.19911		3.772	520.0

^{*} NEGATIVE VALUE FOR ANGLE SIGNALS THAT SEGMENT IS ON SIDE OF PROJECTION FACING AWAY FROM THE PLUME.

THRUSTOR SEGMENT X R THETA

AREA OF SOLCEL IS 20.0 SQUARE FEET.

NO.	NO.	(FT)	(FT)	(RAD)
	PLUME	AFT		
5	110	11.050	1.0000E-05	0.000000
	PLUME	FORWARD		
*	30	3.600	2.100	.314160
	35	3.000	2.100	1.571000
	35	3.600	2.100	3.142000
•	37	3,400	2.100	4.712000

TOTAL SATELLITE AREA, INCLUDING PROJECTIONS, IS 147,00 SQUARE FEET.

MAIN STRUCTURE	PROJECTIONS

BASIC SEGMENTS	THRUSTORS
WINDOWS	RADIO ANTANNAE
INFRA PED PORTS	SOLAR CELL ARRAYS
ULTRA VIOLET PORTS	SUPPORTS

HICKORY-DICKORY SAYS 7.20 SECONDS FOR RUNNING GVERLAY(1.0) FINISHED AT CLOCK TIME 11-24-48.0

> CP TIME USED = 124.18 SECONDS PP TIME USED = 41.35 SECONDS SINCE JOB STARTED

Figure D-41. Satellite Configuration Listing (Page 3 of 3)

	H ,	ATERIALS	AND DEPOSITS	USED IN PROGRAM					
			PIALS	DEPOSITS					
			MIN	N204 MXFL					
		WTN	DOW	HNOS ALZO					
		80L		MMHH50 HCF.	.40				
		UVP	ORT	MMHNO3 FEZO	3				
		GOL	LON	MHN.10 CARE					
		BLA	CK	AMONIA, AGAR	ici.		•		
		HM1 KAP		NHACH NZMA					
		UNA		ANN. AS NIME	03				
				A@MY Unas					
						STRUCTURAL MA			
							THERMAL		
MATERIAL		MELTING	VICKERS	BULK	SURFACE	HEAT	CONDUCTIVITY	YIELD	DENBITY
	NO.	POINT		MODULUS (PSI)	TENSION	CAPACITY	(BTU-IN/ SOFT-DEG-SEC)	STRENSTH (PSI)	(LB/CUFT)
		(968 #)	(46/34)	(597)	(0146764)	(010/[4-056)	*** !	(101)	(60/60/1)
						************	*************		
ALUMIN WINDO	1	1274.00	27.000E+01	48.000E+05	*00.000	.215	.457	47.000E+03 50.000E+03	167.00
BOLCEL	3	1320.00	49.000E+01	23.000E+05	200.000	.180	.002	\$0.000€+02	139.60
IRPORT	\$	1350.00		44.000E+04	200,000	.200	.098	80.0005.02	390.00 135.00
TEFLON	•	978.00	40.000E-01	45.000E+04	15.000	.250	.001	20.0006+02	143.00
OL ACK	7	4700.00		11.000£+06 10.000£+05		.031	.021	30.000E+02 12.000F+03	1230.00
WHITE	•	5270.00	20.0006+01	10.000E+05	190.000	.250 .310	.666 .023 .007	12.0006+03	354.00
KAPTAN	10	1140.00	10.0000+00	51.000E+03	80.000	.310	.002	75.000E+02	88.70
NOTE			O INDICATE NO GENERATED.	DATA ENTERED.					
********	*****		••••		*********	***********			**********
REFERENCE	9U4 ND.	1	CASF NO.	1					
			OPERATING C						
		,((PSIA) 1.000000E-06	URE WALL TEMPER (DEG R) 529.20000					
				T INSTRUCTIONS					
	ſ		\$TOP TIME (8EC) .02000	TIME INTERV. (SEC) .00400	_				
			HMM PROP						
.01	ING POI		EEZING POINT	CRITICAL TEMP.		L PRESS. (PSIA)			
	4 8.000 Vapor c	۰,	397.00000 L10UID CP.	1052.00000	. 401	15.000000 WEIGHT			
	1U/LB-D .9958	••	.090000	(BTU/LB=0E) \$2859;	A4800 0.50	1/LB=MOLE) 16.074000			
((MEAT VA BTU/LB) 78.0000		NT HEAT PUB. (BTU/LB) 97.300000	LIQ. THERM. ((BTU-IN/38FT-DE) .00088	8-8EC)	, COEFF.			
(1	RENCE T DEG #) 40.000		SPECIPIC GRAVITY .870000	VI8CO817 3-8-77-464 9-8-8-8-8-8-8-8-8-8-8-8-8-8-8-8-8-8-8-8	(LD	TEMBION (M)/IM) -000174			
			W204 PROPI						

(1	ING POI DEO 8) 24.2000		EEZING PÓINT (DEG R) 471,370000	ERITICAL TEAP, (DEG R) 776.40000		(P814)	:: : : : : : ·		

Figure D-42. Properties of Materials (Page 1 of 10)

VAPOR CP.	LIQUID CP.	SOLID CP.	VAP. HOL. HEIGHT
(BTU/LR-DEG)	(RTU/LB-DEG)	(8TU/L8-0E6)	(LB/LR-MOLE)
.245000	.374300	,295000	44.005000
LATENT HEAT VAP.	LATENT HEAT FUS.	LIG. THERM. CON!	. ACCOM. COEFF.
(8TU/L8)	(ATU/LE)	(BTU-IN/SOFT-DEG-86	(2)
178.200000	510000	.002509	.100000
PEFERENCE TEMP.		V18C0817V	SURFACE TENSION
(DEG R)	GRAVITY	(LBM/FT-SEC)	(LB(H)/IN)
540.000000	1.434000	.000205	.000140

PHYSICO-CHEMICAL PROPERTIES OF PLUME DEPOSITS

MAME	CODE	FREEZING POINT	SOLID DENSITY	CRITICAL TEMP.	BOILING	FUSIC MEAT	N VAPOR		PMAL CTIVITY LIG.	CAPACI	l T Y	FRACTIVE INDEX LIO.	MOLECULA RADIUS (ANGSTR.	P LIGUID DENSITY } (LB/
		(DEG #1	(LO/CUFT)	(DES R)	(DES #)	(81	U/L8)	(BTU-IN/30F	7-36C-06G)	(BYU/LE	-begi			CUFT)
4204	1	471.4	118.0	776.4	524.2		178.2	2.5208-04	2.515-03	.295	.750	1,420	3,710	49.522
H463	ž	410.2	110.3	851.5	509.0	92.9	200.2	1.17-1-02		. 656	420	1.402	4.005	41.207
MMM	3	397.0	75.3	1052.0	648.0	97.3	370.0	1.6935-03		.350	.700	1.430	4,607	54.312
*****	4	302.0	4.54	1108.0	900.0	-1.0	180.0	2.3496-03		395	.784	1.403	5,250	64.925
MMHN03	5	597.0	4.54	1350.0	900.04	42.5	160.08	-1.0	1.60E-04	.947	.752	-1.0	5,298	-1.0
MMN.40		-1.0	4.50	1350.0	900.04	-1.0	180.00	-1.0	1.13E-04		. 82 1	-1.0	5.644	-1.0
HATER	7	491.0	57.2	1145.0	671.0	143.2	971.3	4.107E-03	1.13E-03	.490	1.000	1.333	3,140	42.347
AMONIA		351.9	51.0	730.0	431.5	194.6	589.3	1.930E-03	9.70E-04		1.064	1.325	2.076	42.543
NHADH	•	349.5	+3.2	989.7	475.0	104.0	640.0	2.450E-03	7.00E-04	1.341	1.015	1.357	3.190	54.937
NHANO3	10	797.0	107.8	1080.0	720.04	32.0	180.08	4.840F-03	1.386-03	.415	.641	1.141	4.100	00.023
AMN. AG	11	598.0	4.54	1029.1		-1.0	180.08	4.710F-03	1.34E-03	.420	.707	1.177	4.348	81.531
MEFUAG	12	-1.0	4.50	1350.0	900.0A	-1.0	180.05	1.4906-04	4.28E-05	.793		-1.0	5.298	42.424
AL203	13	4150.0	247.5	6645.3	4430.0	401.0	100.0	1.3026-03	3.726-04	.198	.333	1.700	0.300	-1.0
HCL	14	291.0	42.4	584.3	338.4	23.4	190.7	4.3186-04	1.236-04	.191	.192	1.411	3,350	102.319
HCL.A0	15	450.0	42.4	875.0	644.4	120.0	177.7	3.2486-03	7.206-04	.753	.570	1.365	3.400	75.725
PE203	10	0.0056	327.0	12960.0	8640.0	1500.n	180.0	4.017E-03	2.50E-03	. 155	.117	3.010	4,540	-1.0
CARBON	17	1000.0	140.0	12075.0	8050.0	-1.0	10.0	1.2006-05	3.43E-03	.170	.129	-1.0	3,000	-1.0
NHACL	1.6	1450.0	45.3	1633.0	1100.08	-1.0	142.0	4.6935-03	1.34E-03	.397	.538	1.371	3.240	-1.0
AGAMEL	1.9	463.0	4.54	1035.3	690.2	135.0	100.0	4.4016-03		.434	.724	1.378	3,495	00.011
4544	20	494.3	71.5	1117.0	696.0	170.0	550.0	1.4146-03		.478	.736	1.464	3.094	62.789
MEHANG	51	395.1	4.54	1057.5	705.0	-1.0	100.0	2.5016-03	7.32E-04	1.030	.780	1.424	3.234	102.019
N3H503	55	618.0	4.54	1115.0	743.84	-1.0	180.08	-1.0	-1.0	-1.0	-1.0	-1.0	3,743	-1.0
THYMBA	53	503.0	4.54	1350.0	900.04	-1.0	180.00	-1.0	1.20E-04	.766	,580	-1.0	4.001	-1.0

NOTE VALUES OF -1.0 INDICATE NO DATA ENTERED.
OR INTERNALLY GENERATED.
(A). DECOMPOSITION TEMPERATURE.
(B). HEAT OF ARLATION OR DECOMPOSITION.
(S). SUBLIMATION.

HEAR	CONSTANTS	0F	MATERIALS	ON	SURFACE
		•		•	

HAME	10	CONSTANT	FORES	EXPONENT (SLOPE)	MOLFCULAR
		(FT-LB/LB)	LIMIT	(86044)	(VAPOR)
		(,,-60,60)	(FT/SEC)		******
	•••		••••••	**********	
		8	TRUCTURAL		
ALUMIN	1	3.0006.00		-3.3125	
#1400=	5	5.000E+0		-3.5025	•••
BOLCEL	3	5.000E+0		-3.5625	•••
IRPORT	4	5.000E+0		-3.5625	•••
UVPORT	5	5.000£+0		-3.5625	•••
TEPLON	•	1.500E+0		-2.5000	•••
BOLD	7	3.000E+0	460	-3.3125	
PLATING					
BLACK		1.2006+0	200	-2.5000	•••
COATING					
BHITE	. •	1.200E+0	200	-2.5000	•••
COATING					
KAPTAN	10	1.500E+0	****	-2.5100	•••
		ē	SF18043		
N204	1	1.0006+04	550	-3.0000	44.0
MN03	ž	1.0006.0		-3.0000	43.0
-	3	1.0006+0	550	-3.0000	46.0
DZHMMM	4	1.000E+0	550	-3.0000	
MMHN03	5	1.0006+0	550	-3.0000	109.0
MMN . AG	٠	1.000E+0	550	-3,0000	154.0
WATER	7	1.000E+0	650	-3.0000	10.0
AMONIA		1.500E+0	178	-3.1900	17.0
NHACH	•	1.5006.05	160	-3.1400	35.0
NHENO3	10	1.500E+0	123	-3.1900	80.0
AMW. AG	11	1.500E+0	101	-3.1400	98.0
MEFUAD	15	1.500E+0		-3.1900	252.0
AL 203	13	1.500E+05	6 41	-3.1900	101.4
HCL	14	1.500E+0		-3.1900	36,5
HCL.40	15	1.500E+0		-3.1900	21.7
FE203	10	1.500E+0	5 70	-3.1900	100.0

Figure D-42. Properties of Materials (Page 2 of 10)

CARBON						-3.1		48.
NHECL	10	1.500	E+85	130			.00	53.
AGAMEL	1.0	1.500	E+05	101		-3.1	***	87.
PHSH	50	1,500	E+05	150		-3.1	900	32.
W2M440	15	1.500	E+05	101		-3.1	400	50.
N3H503	55	1.500	E+85	101		-3.1	900	95.
ABHYNT	5.2	1.500	E+05	161		-3.1	***	113.
NOTE	VAL	JES OF -	-1.0	14010	ATE 40	DATA	ENTER	EO.
	AP TI	STERNAL		THE BAT	631			
•••••	*****					••••	•••••	******
••••	••••	••••••					#41	**************************************
•••••	••••	VELOC1	74 LI		.]#[78 (LL #41	enials.
•••••	••••	VELOC1	TV LI	D#E# (, IMITS (ER	 LL #41	enials.
•••••	••••	VELOC1	74 LI	D#E# (HYPI	ER ITY	 LL #41	Salvra
•••••	••••	VELOC1	TV LI) #3#C	, IMITS (ER ITY	LL #41	'ENTALS
•••••	••••	VELOC1	TY LI) #3#C	HYPI	ER ITY	LL #41	PRIALS

SUPFACE PROPERTIES INPUT

ALUMIN

	485048-					THERMAL			PER INCH		S H I	T T A			
FINISH	TIVITY	IALLA		SPECULAR	THERMAL	BTU-IN/	INDER	15	MEM INCH	S INIC	KME BB AT	0.5	AVELENGT	0.05	
(40-14)	(80584)	(IME HM)	01+1085	378C0C0M		FT-DEG-86	c	(18)	(18)	(1F)	(14)	(VIS)	(04)	(44)	0.01 (UV)
0.0	.0100	.0400	-1.0	. 9900	. 7600	.4570	0.0	0.0	0.0	0.0	0.0	1,1	1,1	0.0	1,1
.50	.0110	.0350	. 9300	.9890	. 9450	-1.0	-1.0	-1.0	-1.0	0.0	0.0	0.0	0.0	0.0	0.0
1.00	.0475	.0500	-1.0	.9525	. 9500	-1.0	-1.0	-1.0	-1.0	0.0	0.0	0.0		0.1	0.0
2.00	.0200	.0400	. 4500	. 9000	. 9000	-1.0	-1.0	-1.0	-1.0	0.0	0.0	0.0		0.0	0.0
3.00	.0300	-1.0	-1.0	. 9700	-1.0	-1.0	-1.0	-1.0	-1.0		0.0	0.0	0.0		0.0
4.00	.1100	.0500	.7300	. 8900	. 9500	-1.0	-1.0	-1.0	-1.0	0.0	0.0	0.0			1.1
5.00	.0700	-1.0	-1.0	. 4300	-1.0	-1.0	-1.0	-1.0	-1.0		0.0		0.0	0.0	
	.1000	-1.0	-1.0	.9000	-1.0	-1.0	-1.0	-1.0	-1.0	0.0	0.6	0.0	0.0	0.0	0.0
7.00	. 1200	. 0850	.5800	. 6800	.9150	.9506	-1.0	-1.0	-1.0	0.0	0.0	0.0		0.0	0.0
8.00	.1400	.0500	-1.0	. 4600	.9500	-1.0	-1.0	-1.0	-1.0	0.0	0.0	0.0		0.0	0.0
	.1600	.0600	-1.0	.8400	.9400	-1.0	-1.0	-1.0	-1.0	0.0	0.0	0.0	0.0	0.0	0.0
10.00	.1700	-1.0	-1.0	. 6300	-1.0	-1.0	-1.0	-1.0	-1.0	0.0	0.0	0.0		0.0	0.0
11.00	.1800	-1.0	-1.0	.0200	-1.0	-1.0	-1.0	-1.0	-1.0	0.0	0.0	0.0	0.0	0.0	0.1
12.00	.1900	.0800	-1.0	.6100	.9200	-1.0	-1.0	-1.0	-1.0	0.0	0.0	1.1	0.0	0.0	0.0
13.00	.2000	-1.0	-1.0	. 8000	-1.0	-1.0	-1.0	-1.0	-1.0	0.0		0.0	0.0		0,0
14.00	.2100	-1.0	-1.0	.7900	-1.0	-1.0	-1.0	-1.0	-1.0	6.0	0.0	1.0	0.0	0.0	0.0
15.10	.2300	-1.0	-1.0	.7700	-1.0	-1.0	-1.0	-1.0	-1.0	0.0	0.0		0.0	0.0	0.0
03.05	.2000	.1000	-1.0	.7400	.9000	-1.0	-1.0	-1.0	-1.0	0.0	0.0	0.0	0.0	0.0	0.0
25.00	.2980	-1.0	-1.0	.7020	-1.0	-1.0	-1.0	-1.0	-1.0	0.0	0.0	0.0	0.0	0.0	0.0
30.00	.3300	.2000	.5000	. 6700	.8000	-1.0	-1.0	-1.0	-1.0	4.0	0.0	0.0	0.0	0.0	0.0
35.00	.3000	-1.0	-1.0		-1.0	-1.0	-1.0	-1.0	-1.0	0.0	0.0		0.0	0.0	0.0
40.00	.3000	-1.0	-1.0	. 6200	-1.0	-1.0	-1.0	-1.0	-1.0	0.0	0.0	0.0		0.0	0.0
50.00	200	.2100	-1.0	.5800	.7900	-1.0	-1.0	-1.0	-1.0	0.0	0.0	1.1		0.0	0.0
60.00	.4750	-1.0	-1.0	.5250	-1.0	-1.0	-1.0	-1.0	-1.0	0.0	0.0	0.0	0.0	0.0	0.0
80.00	.5000	-1.0	.5000	-1.0	-1.0	-1.0	-1.0	-1.0	-1.4	0.0	0.0	1.1			0.0
100.00	.5000	-1.0	.4400	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	0.0	0.0		0.0	0.0	0.0
130.00	.6700	.3400	. 3300	-1.0		-1.0	-1.0	-1.0	-1.0	0.0	0.0	0.0	0.0	0.0	0.0
150.00	.7500	-1.0	.2500	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	0.0	0.0	0.0	0.0	0.0	0.0
100.00	.4000	-1.0	.2000	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	0.0	0.0	0.0	0.0	0.0	
401			INDICATE ENERATED	NO DATA	ENTERE	D.									

-

Figure D-42. Properties of Materials (Page 3 of 10)

80LCEL

(HU-IN) (80	117	IVITY	801	E C T I A R SPECULAR	THERMAL	COND. RTU-IN/ FT-DEG-8	INDEX	(18)			S M I KNESS AT 1 (IR)		A N C	0.05 (UV)	0.0 (UV
NOTE VA	000 LUES	.8500 OF -1.0	-1.0	.9000 NO DATA	.1500	.0024	2.4500	0.0	0.0	,2500	.4500	. 4500	0.0	0.0	0.
•••••		•••••	*******	•••••	•••••	IRPORT		• • • • • •	••••••		•••••	••••••	••••••	•••••	
BURFACE ABS FINISH TIV (MU-IN) (SO	ITY	IALLA	80(AR	THERMAL	THERMAL COND. BTU-IN/ FT-DEG-B	INDEX	15 (IR)				HICRON 0.5 (VIB)	A N C		• . o
NOTE VA	INTER	NALLY G	ENERATED,			D.	2.7200	.0050		.7500	.7500	.1500	0.0	1.0	• •
						UVPORT									
SURFACE ASS FINISH TIV: (MU=IN) (SO	ITY LAR)	IVITY (THERM)	DIFFUSE	SPECULAR	THERMAL	THERMAL COND. BTU-IN/ FT-DEG-8	INDEX	(IR)			S M I HHEBS AT 1 (IR)	WICRON (VIS)	O.2 HAVELENS	THE OF 0.05 (UV)	0.0
HOTE VAL	550 LUES	.9100 OF -1.0	-1.0	.9450 NO DATA				0.0		0.0	.0500	.****	,6500	.2500	.005
						TEPLON							••••••		
SURFACE ASSO Finish Tivi (MU-IN) (SOL	177	IVITY	# O t	AR'	THERMAL 188	COND. BTU-IN/ TT-NEG-8	INDEX	15 (18)		OF THIC		MICRON 0.5 (VIS)	(UV) 0.5 HVAEFEND V V C		
50.00 .26 NOTE VAL	LUES (MALLY 6	INDICATE ENERATED		.1300 Entere	.0019	-1.0	0.0	0.0	0.0	0.0	0.0	•.•	0.0	0.0
••••••						6010	PLATIN								
	177	IVITY	8 D L	. A # 1	HERMAL	COND. ATU-IN/	INDEX	15 (IR)							0.0 (UV
SURFACE AUSO Finish Tivi (MU-In) (SOL						7-DEG-81									

Figure D-42. Properties of Materials (Page 4 of 10)

BLACK COATING SURFACE ASSORD- PHISS- R E F L E C T I V I T V THERMAL REFRACT.
FINISH TIVITY IVITY S O L A H THERMAL COND. INDEX
(MU-IN) (SOLAH) (THERM) OIFFUSE SPECULAR BTU-IN/
SOFT-DEB-SEC (IR) (IR) (IR) (IN) (VIS) (UV) NOTE VALUES OF -1.8 INDICATE NO DATA ENTERED. OR INTERNALLY GENERATED. WHITE COATING SURFACE ABSORD- EMISS- R E F L E C T I V I T V THERMAL REFRACT.

FINISH TIVITY IVITY B O L A R THERMAL COND. INDEX

(MU-IN) (SOLAR) (THERM) DIFFUSE SPECULAR

BTU-IN/

ROFT-DES-DEC (IR) (IR) (IR) (IR) (VIS) (UV) NOTE VALUES OF -1.0 INDICATE NO DATA ENTERED. OR INTERNALLY SEMENATED. KAPTAN SURFACE ABBORNS EMISS OF FLECTIVITY TMERMAL PEPRACT.

FINISH TIVITY IVITY SOLAR TMERMAL COND. INDEX
PRINCH OF THICKNESS AT MICRON NAVELENGTHS OF

(MU-IN) (SOLAR) (TMERM) DIFFUSE SPECULAR STU-IN/ 15 S & 1 0.5 0.2 0.05

BOFT-OBSO-BEC (IR) (IR) (IR) (IR) (VIO) (UV) 10,00 -1,0 .7000 -1,0 -1,0 .3000 .0003 1,7000 0,0 .0008 NOTE VALUES OF -1.0 INDICATE NO DATA ENTERED. OR INTERNALLY GENERATED. N204 DEPOSIT PER INCH OF THICKNESS AT HICRON HAVELENGTHS OF TEMPERE ABBORS- EMISS- R E P L E C T I V I V TMERMAL REPRACT.
ATURE TIVITY IVITY B O L A R TMERMAL COND. INDEX
(ORS R) (BOLAR) (TMERM) DIPPUBE SPECULAR BOPT-DEG-SEC (IR 0070 - 2750, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2750, 2750, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, 2050, .0272 NOTE VALUES OF -1.0 INDICATE NO DATA ENTERED. OR INTERNALLY SEMENATED. HN03 DEPOSIT

TEMPERE ABSONS- FMISS- R E F L E C T I V I T V THERMAL REFRACT. T R A N S M I T T A N C E
ATURE TIVITY IVITY S O L A R THERMAL COND. INDEX PFR INCM OF THICKMESS AT MICRON MAVELENSTMS OF
(DES R) (SOLAM) (THERM) DIFFUSE SPECULAR BTH-IN/ 15 S 2 1 0.5 0.2 0.05 0.01
SOFT-DEG-SEC (ER) (IP) (IR) (IR) (VIS) (UV) (UV) (UV)

500.00 .7177 .9722 50.1111 .2223 .0278 .0034 1.4020 .0290 .0292 .0292 .0295 .0295 .0276 .0274 .0273

800.00 .7177 .9722 50.1111 .2223 .0278 .0034 1.4020 .0290 .0292 .0292 .0295 .0295 .0277 .0276 .0274

MOTE VALUES OF -1.0 INDICATE NO DATA ENTERED.

DR INTERNALLY GENERATED.

Figure D-42. Properties of Materials (Page 5 of 10)

0.01 (UV) -1.0

	******) DIFFUS			STU-IN		15 (IR)	(1R)	(18)	(IR)	(V18)	(UV) 9.8 HAVELENG	e.e
******	.7751 .7769		• 50.1125 1 50.1115			.0017 -1.0	1.4300	.0200	. 0201	16261	.0294 .0294	.0294	.0274	.027
	OR INT	ERNALLY	BENERATE	D.									*******	
						иннн\$0								
(DEG R)	(SOLAR)	IVIT	- R E F (7 3 0 1) DIFFUSI	LAO	THERMAL BO	COND. BTU-IN/ FT-DEG-1	EC	15 (IR)			KNESS AT	MICRON 0.5 (VIS)		THE OF
540.00		-1.	-1.0	-1.0 -1.0			-1.0 -1.0	-1.0		-1.0 -1.0	-1.0 -1.0	-1.0 -1.0		-1. -1.
	OR INTE	RNALLY	O INDICAT				*******			•••••	•••••		•••••	
						PHH403	DEPOSI	₹						
ATURE (DEG #)	(SOLAR)	IVITY (THERM	9 0 9 01FFUSE	BPECULA	THERMAL R 80	COND. STU-IN/ FT-DEG-8	EC	15 (IR)	PER INCH S (IR)	OF THIC	ENEBS AT	PICRON 0.5 (VIS)	(UA) 9°S MTAEFENG.	0.05 0.05 (UV)
500.00			-1.0	-1.0 -1.0	-1.0 -1.0		-1.0 -1.0	-1.0 -1.0	-1.0 -1.0	-1.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0	-1.
	OR INTE	RNALLY	O INDICAT	·•										
						MHN.48	DEPOSI		,				••••••	
ATURE (DEG R)	(BOLAR)	IVITY (THERM) DIFFUSE	SPECULA	THERMAL H 804	COND. 87U-1N/ 77-DEG-8	EC	15 (IR)	PER INCH 5 (IR)	OF THICK	CIR)	#ICRON 0.5 (4'8)	MAVELENGT 0.2 (UV)	46 OF
560.00 800.00		.7500	\$0.2750 50.1990		.2500 .2476	-1.0	3.0030	.0500	.5000	.0000	.0226	.0227	.0195	.0193
40TE	VALUES OR INTE	OF -1.	O INDICAT	E 40 DAT	A ENTERE	·.								

HATER DEPOSIT

ATURE	AUSOPA- TIVITY (BOLAR)	IVITY	8 0		THERMAL	THERMAL COND. RTU-IN/		15 (IR)	PER INCH	OF THIC	KNESS AT	T T MICRON 0.5 (VIS)	O.S MAVELENG	E THE OF 0.05 (UV)	0.01
492.00 560.00 672.00	.7600 .7634 .7764	.9542 .9630	50.1200 50.1103 50.1140	.2400 .2366 .2366	.0500 .0458 .0370	.0042	1.3330 1.3276 1.3186	.0293	.0295 .0295 .0295	.0295 .0295 .0296			.0260		
	OR INTE			E NO DATA	ENTERE	•									

Figure D-42. Properties of Materials (Page 6 of 10)

AMONTA	REBORTS

TEMPERE ABSORD- EMISS- R E F L ATURE TIVITY IVITY S O L	ECTIVITY THERMAL REFRACT. LAR THERMAL COND. INDEX	T R	A N 8 M 1 T T THICKNESS AT MICRO	A N C E
(DEG R) (SOLAR) (THERM) DIFFUSE		15 5	7 1 0.5 18) (18) (VIS	0.2 0.05 0.01 (UV) (UV) (UV)
540.00 .7045 .9876 50.1078 000.00 .7082 .9087 50.1069	.2155 .0194 .0019 1.3250 Sec. 1.0 1.3250			7 .0200 .0279 .0278
NOTE VALUES OF -1.0 INDICATE OR INTERNALLY SEMERATED.				

NH40H DEPOSIT

ATURE	TIVITY	IVITY	8 0	E C T I L A R SPECULAR	THERMAL	INDEX	(IR)	PER INCH S (IR)	OF THIC	8 M I KNE88 AT (IR)	T T MICRON 0.5 (VIB)	(AA) 0'S HVAEFENG.	E THB OF 0.05 (UV)	0.01 (uv)
\$40.00 800.00 NOT		.9794 OF -1.0	50.1091 50.1082 INDICAT	.2182 .2185 E NO DATA	.0228 .0206 ENTEREC	1,3570	. 0293 . 0293	.1294 .0294	,0294 ,0295	.0296 .0297	.0296	.0279	.0277 ,0278	.0276

NHANOS DEPOSIT

TEMPERE ABSORS- EMIS ATURE TIVITY IVI (DEG R) (SOLAR) (THE	TY BOLAR 1	/ I T Y THERMAL REFRACT, THERMAL COND. INDEX BTU-IN/ BOFT-DEG-SEC		T R A N PER INCH OF THIC 5 2 (IR) (IR)	B M I KNESS AT	T T A N C MICRON WAVELENG 0.5 0.2 (VIS) (UV)	E OF 0.05 0.01 (UV)
400.00 .7975 .99	97 50.1017 .2034 60 50.1013 .2025 1.0 INDICATE NO DATA V GENERATED.	.0043 .0048 1.1413 .0032 -1.0 1.1205 ENTERED,	.0300	.0301 .0301 .0301 .0303	.0302	.0302 .0292 2920, 2020.	.0290290 .0290290

AMN.49 DEPOSIT

	717177	IVITY		A R	THERMAL	COND.	REFRACT.		PER INCH	OF THIC	KHESS AT	HICRON	HAVELENGT	HS 07	
(DEG R)	(SOLAR)	(THERM)	DIFFUSE	SPECULAR		87U-IH/ 74-9E6-81	EC	15 (18)	(10)	(1R)	(18)	0.5 (VIS)	6.5	0.05 (UV)	0.01
540.00	-1.0	-1.0	-1.0	-1.0	-1.0	.0047	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	-1.4
600.00	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0

MEFUAG DEPOSIT

ATURE	TIVITY	IVITY	8 0	AR	THERMAL		REPHACT.		PER INCH	OF THIC	S H I	HICHON	A N C WAVELENGT	HS OF	
(DEG A)	(BOLAR)	(THERM)	DIFFUSE	SPECULAR		BTU-14/		15	5	. 2	1	0.5	0.2	0.05	0.01
					80	T-DES-8	EC	(14)	(1R)	(TR)	(IR)	(418)	(UV)	(UV)	(uv)
540.00	-1.0	-1.0	-1.0	-170	-1.0	.0001	-1,0	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0
800.00	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0	-1.0

Figure D-42. Properties of Materials (Page 7 of 10)



4 6

NOTE VALUES OF -1.8 INDICATE NO DATA ENTERED. OR INTERNALLY GENERATED.

4L203 DEPOSIT

ATURE	ABSORB- TIVITY (SOLAR)	IVITY	8 0	E C T I L A R SPECULAR	THERMAL			15 (IR)	PER INCH	OF THEC	R M I KNESS AT 1 (IR)	T T MICRON 0.5 (VIS)	A N C HAVELENGT 0.2 (UV)	E HS OF 0.05 (UV)	(UA) 0.01
540.00 650.00 1570.00 3600.00	.9800 .0880 .9800 .5000	.1100	50.0100 50.4560 50.0100 50.2500	0000. 0000. 0000.	.4000 .8700 .4100 .7200	0.0	1.7000 1.6752 1.6176 1.4552	0.0	0.0 0.0 0.0	0.0 0.0 0.0	0.0	0.0 0.0 0.0	0.0	*.0 *.0 *.0	• • •

NOTE VALUES OF ${\sim}1.0$ INDICATE NO DATA ENTERED. OR INTERNALLY GENERATED.

HCL DEPOSIT

TEMPERE A ATURE T (DEG R) (IVITY	IVITY	8 O L	AR 1	HERMAL	THERMAL COND. GTU-IN/ T-DES-S	INDEX	(18)	PER INCH 5 (IR)	OF THIC	B M I KNESS AT (IR)	(VIB)	(UV)	0.05 (UV)	0.01 (UV)
800.00 NOTE		.9735 S	50.1116 50.1106 Indicate Emerated.	.2212 ATAG ON	.0265 .0265 ENTERED	-1.0	1.3002	.0201	.0292 .0292	,0292 ,0293	.0294 .0295	,0295 ,0295	.0275 .0277	.0274 .0275	.0272 .0274

HCL.AQ DEPOSIT

TEMPERE	ABBURS-	Ewiss-	REFL	ECTI	V I T V	THERMAL	REFRACT.	•	T 1	R & N	8 M E	7 7	ANC	E	
ATURE	TIVITY	IVITY	8 6 1	, A R	THERMAL	COND.	INDEX		PER INCH	OF THIC	KHESS AT	MICRON	MAVELENS'	THE OF	
(DEG R)	(SOLAR)	(THERM)	DIFFUSE	SPECULAR	_	STU-IN/		15	\$	2	1	0.5	5.0	0.05	0.01
					801	T-0E6-8	EC	(IR)	(18)	(IR)	(IR)	(V18)	(uv)	(VV)	(UV)
540.00	.7811		50.1095	.2189	.0237	.0032	1.3650	.0292	.02+3	.0294	.0296	.0296	.0278	.0277	.0275
*****	.7629	. 4786	50.1084	.2171	.0214	-1.0	1.3442	.0293	.0294	.0295	.0296	.4296	.0279	.0278	.0277
			INDICATE ENERATED.		ENTERF										

FE203 DEPOSIT

	ABSORB-	EMISS- IVITY			V I T V		. REFRACT		PER INCH	OF THIC	S M I	T T HICRON	A N C	E HS OF	
(DEG R)	(80LAR)	(THERM)	DIFFUSE	SPECULAR		FT-DEG-		15 (IR)	\$ (1R)	(IR)	(18)	0.5 (VIS)	(44)	0.05 (UV)	(An)
560.00 1370.00 2650.00	.5040 .6000 .7120	.0500	50.2480 50.1600 50.1440	.4960 .3200 .2000	.3700 .1500 .1100		3.0100 2.9436 2.0428	0.0	1.0 0.1 0.1	0.0	0.0	0.0	1.1	1.0 1.0	::
			INDICAT	E NO DATA	ENTERE	٥.									

CARBON DEPOSIT

Figure D-42. Properties of Materials (Page 8 of 10)

CRS																
		HAVELENS		KNESS AT				15			THERMAL	. A R		IVITY	TIVITY	ATURE
(U	(ÚV)	(44)	(vis)		IA)	••••	(1Å)	(141	EC	7-066-8	107					
0	0.0	1.1 1.1 1.1	0.0 0.0 0.0	0.0	0.0 0.0		0.0 0.0	0.0 0.0 0.0	-1.0 0.0 0.0	0.0	.1900	.3520	.8100 0.1760 7.1800 0.1840	.0000	.4480	720.00 170.00 620.00
					•		•				L ENTERED	,	WERATED,	MALLY	OR INTE	
		•				****			DEPOOL	MMECL		*******	******	*****	••••••	
		A N C			A N		•		REPRACT.	THERMAL			.	fulle.	485088-	75 40505
0.	TH S OF 0.05 (UV)	HAVFLENS' 0.2 (UV)	HICRON 0.5 (VIB)	(NESS AT	THIC: 2 18)	H 0f	PER INC (IR)	15 (18)	INDER		THERMAL	A R		IVITY	TIVITY	ATURE
:02	.0276	.0278	4950.	.0294	0294		.0294	.0292 .0292	1.3710	-1.0	.0243 .0221 ENTERED		INDICATE	.9779 S		540.00 800.00 NOTE
				•••••	•••••	••••	•••••	•••••					NERATEN.			
									DEP0811	ABAHCL						
•.		A N C HAVELENST 9.2 (UV)				H DF	PER INCI	15 (IR)	REPRACT. INDEX		THERMAL	A R		IVITY	TIVITY	ATURE
	.0276 .0277	.0277	.0295	.0295	0293 0294	••••	. 0293	. 02-02		.0045	.0520	.5103	0.1100	.9749 9	.7799 .7817	
		*******								•	ENTERED		INDICATE NEWATED.		OR INTER	
									DEPOSIT	N2H4						
• • CU		A N C MAVELENST 6.2 (UV)				H OF		15 (IR)	REPRACT. INDEX		THERMAL	A R		IVITY	TIVITY	
.020	.0271		.0292	.0292	0240		.0249	.0200	1,4644	-1.0	.0353	.5542	::::::	.9647 5	.7717 .7730	100.00
		******		******		••••		•••••					ERATED.	ATTA BE	OR INTER	
									DEPOSIT	MSH410	,					
0.0	HS OF	A N C MAVELENGT 0.2 (UV)	FICRON (H OF		15	REPRACT. INDEX	THERMAL COND. BTU-IN/ T-DEG-BE	THERMAL	C T T	8 0 L	(MISS- V TY THERM)	AUSORU- TIVITY (SOLAR)	TEMPERE ATURE
	.0273	.0275	.0294	.0294	292	••••	.0291		1.4240	.0024	.0304	.2243	.1122	5		40.00
										•	ENTERED	NO DATA			VALUES DR INTER	

Figure D-42. Properties of Materials (Page 9 of 10)

N3H50	3	DEP	TIRC

ATURE	TIVITY	IVITY	8 0	E C T I L A R SPECULAR	THERMAL		INDEX	15 (18)	PER INCH S (18)	OF THIC	S M I KNESS AT (IR)	T T MICRON 0.9 (VIS)	(UA) 0.5 PVAEFEND.	E 1MB OF 0.05 (UV)	0.01 (UV)
			-1.0 -1.0 INDICATI ENERATED	-1.0 -1.0 E NO DATA	-1.0 -1.0 ENTERE	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0

AGHYNT DEPOSIT

ATURE	TIVITY	14114	\$ 0 I	E C T I A R SPECULAR	THERMAL	THERMAL COND. BTU-IN/	REFRACT. INDEX		PER INCH	OF THIC	B M I KNESS AT	T T HICRON	A N C HAVELENGT	E MB OF 0.05	0.01
					501	T-DEG-SE	C	(IR)	(IR)	(IR)	(IA)	(A18)	(UV)	(UV)	(UV)
540.00	-1.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0	-1.0 -1.0	-1.0
MOTE		05 6	THRICATE												

NOTE VALUES OF -1.0 INDICATE NO DATA ENTERED. OR INTERNALLY GENERATED.

UNAVAILABLE SIGNIFICANT DATAR NEITHER INPUT NOR INTERNALLY APPROXIMATED

HYDRATED MONOMETHYL HYDRAZINE -- HEAT OF FUSION.

MONOMETHYLMYDRAZINE NITRATE -- THERMAL COMDUCTIVITY (SOL). REPRACTIVE INDEX (D LINE). BOILING POINT. CRITICAL PRESSURE, VISCOSITY (LIQUID). SURFACE TENSION (LIQUID).

MYDRATED MONOMETHYLHYDRAZINE NITRATE-MELTING POINT. THERMAL CONDUCTIVITY (SOL). REFRACTIVE INDEX (D LINE). SOILING POINT. CRITICAL PRESSURE, VISCOSITY (LIQUID). SURFACE TENSION (LIQUID). MEAT OF PUSION.

AMMONIUM NITRATE --VISCOSITY (LIQUID).

HYDRATED AMMONIUM NITHATE . HEAT OF FUSION.

MYDRATED MIXED MITRATES -- MELTING POINT. REFRACTIVE INDEX (D LINE). VISCOBITY (LIQUID). SURFACE TEMBION (LIQUID). MEAT OF PUBION.

ALUMINUM OXIDE -- BOILING POINT. VISCOSITY (LIQUID). SURFACE TENSION (LIQUID).

MYDROGEN CHLORIDE -- VISCOSITY (LIQUID). SURFACE TENSION (LIQUID).

IRON OXIDE -- BOILING POINT. VISCOSITY (LIQUID). SURFACE TENSION (LIQUID).

CARSON -- REFRACTIVE INDEX (D LINE). SOILING POINT. VISCOSITY (LIQUID). SURFACE TENSION (LIQUID). MEAT OF FUSION.

AMMONIUM CHLONIDE -- BOILING POINT. VISCOSITY (LIGUID). SURFACE TENSION (LIGUID). HEAT OF FUSION.

HYDRAZINE HYDRATE -- VISCOSITY (LIQUID), HEAT OF FUSION.

MYDRAZINE NITRATE -- SPECIFIC MEAT (SOLID). THERMAL CONDUCTIVITY (SOL). REFRACTIVE INDEX (D LINE). BOILING POINT. LIBUID MEAT CAPACITY. THERMAL CONDUCTIVITY (LIQ). VISCOSITY (LIQUID). SURFACE TENSION (LIQUID). MEAT OF FUSION.

MYDRATED MYDRAZINE NITRATE -- THERMAL CONDUCTIVITY (BUL). REFRACTIVE INDEX (D LINE). BOILING POINT. VISCOSITY (LIQUID). SURFACE TENSION (LIQUID). MEAT OF FUSION.

MICHORY-DICKORY SAYS 483.60 SECONDS FOR RUNNING OVERLAY 14.0 FINISMED IT CLOCK TIME 11-32-51.6

> CP TIME USED = 150.94 SECONDS PP TIME USED = 43.94 SECONDS SINCE JOB STARTED

Figure D-42. Properties of Materials (Page 10 of 10)

	KANDSEN-FUNCANIE CO	EFFICIENTS
FLUID	CNSTV	CHSTH
M204	2.9673899015E-05	3.1429169E-05
HN03	3.47268965476-06	3.4373957E-05
MM(44	2.96738990156-05	2.85714296-04
минико	3.50014217406-00	8.2537901E-04
MMH403	4.56781964025-06	1.9784219E-06
MHN. 40	5.4294527618E-06	1.3994642E-04
MATER	1.85023069946-06	2,59695846-06
AMONTA	1.00393190006-00	3.67430526-06
NH#OH	2.58839004426-06	2.4415072E-06
NHEND3	3.91327791606-06	1.71492356-05
AHN.AG	4.33120490016-00	1.45732796-05
MEFUAG	0.9453793094t-06	5.30617636-07
4L203	4.41654632966-06	4.6132859E-06
+CL	2.44327367526-06	1.4441706E-06
HCL.40	2.03810035896-06	1.16599926-05
FE203	5.53421070226-06	3.1949308E-05
CARBON	3.03121203966-06	4.25187646-05
WHACL	5.2001000100E-00	2.1075260E-05
AGAMEL	4.13911236306-06	1.55745756-05
PHEN	2.47497426648-06	2.2200593E-06
NZHAAG	3.09371783306-00	9.07536516-06
N3H503	4.20439574356-06	-1.0
AGHYNT	4.65087770946-06	1.49407286-04
UES OF-1.0	INDICATE THAT THE TER	14
	ATED BECAUSE ESSENTIAL	

NOTE--VAL COULD NOT DATA -ERE

CALINGEART-DAVIS COEFFICIENTS

LN(P) = 4+(8/(T-43.0)) FOR P IN MM. T IN DEG K

NAME	A	100
4204	26,795586	-3254,496353
HNGS	25.070959	-3070,394206
MMH	24.439086	-3363,502319
MMHH50	37.557346	-10844.005645
MMMMO3	-1.7	-1.0
MHN.AG	-1.0	-1.0
MATER	25,498530	-3914.416532
AMONTA	24.130000	-2026.677054
NHADH	21.901157	-1763-122984
NHANC3	25.910304	-4313.145316
AMN.AG	26.682285	-4346.664654
MEFUAD	24.704724	-5004.357300
AL203	51.976900	-43884,439245
HCL	22.901702	-1315.601431
HCL.40	32.504442	-6453.586427
FE203	34,063461	-96257.003310
CARBON	33.844480	-88654.500510
NHACL	24.163152	-5071.134691
AGAHCL	25.757619	-4001.174983
N2H4	26.112522	-4221,462134
NZHGAG	25.034350	-4145.780868
N3H503	26.027094	-4515.057023
ABHYNT	26.704724	-5884.357388
	0-1.0-164	~ **** * 3 7 / 3 4 4

NOTE--VALUES OF-1.0 INDICATE THAT THE TERM COULD NOT BE CALCULATED RECAUSE ESSENTIAL DATA MERE NOT SUPPLIED

	 	 _

	CRITICAL TEMP.	CRITICAL PRESS
	DEG #	PSIA
N204	776.400000	1441.300000
HN03	053.500000	905.723700
MMM	1052.000000	1195.00000
-	1104.000000	1128.642240
MMHH03	1350,000000	-1.00000
MHN. AG	1350.000000	-1.00000
MATER	1105.000000	3226,000000
ATHOMA	730.000000	1635.700000
NH4OH	989.700000	1396.744316
NH4NO3	1080.00000	1125.166714
AHN. AG	1029.150000	1074,362773
MEFUAD	1350.000000	1394,980911
AL203	**45.00000	6689, 395434
HCL	584.300000	1178.700000

Figure D-43. Derived Coefficients and Properties of Deposit Constituents (Page 1 of 5)

```
897.207071
13004.323409
12114.328458
507.00000
1080.504959
2135.00000
1102.486540
1100.77640
1344.480911
                                                   875.00000
1290.000000
12075.000000
1633.00000
1137.600000
1157.500000
                      HCL.40
FE203
CARHON
                     HSH496
HSH496
HSH496
                      43M503
                                                       1115.040000
NOTE--VALUES OF-1.0 INDICATE THAT THE TERM
COULD NOT BE CALCULATED BECAUSE ESSENTIAL
DATA MERE NOT SUPPLIED
                                              VAPOR ENTHALPY AT ABSOLUTE ZERO NAME (BTU/LB
                                    NAME (810-24)
N204 273,023
MN03 403,104
MNH 148,366
                                                                                                               148.366

.1.0

161.622

121.337

1210.59

604.941

1263.56

398.376

-1.0

541.470

213.165

79.2801

-1.0

-1.0
                                                                MMM
OSHMM
EOWHMM
BA-WMM
RATAW
ALNOWA
HOAMM
EOWAMM
                                                                AMN.AS
MXFUAS
AL203
MCL
MCL.AS
FE203
CARSON
                                                                MM4CL
ABAMCL
N2M4
N2M4AB
N3M503
ABMYNT
                                                                                                               -1.0
-1.0
776.821
122.056
NOTE--VALUES OF-1.0 INDICATE THAT THE TERM COULD NOT BE CALCULATED SECAUSE ESSENTIAL DATA MERE NOT SUPPLIED
                                 ------
                                              VAPOR PRESSURE AT THE FREEZING POINT NAME (PSIA)
                                       N204 2.19063
HN03 101570
HMH 3.533781£-03
                                                                  мини20
Еоинии
                                                                                                               4.387449E-17
                                                                                                            4,387449E-17
-1.0
-1.0
8,589199E-02
.740905
.354986
53,5362
1.06927
-1.0
                                                                  MHN.49
HATER
AMONIA
NM40H
                                                                   NHANO3
AMN.AG
MXFUAR
                                                                                                             1.00-27
-1.0
2.0228
5.472401E-05
2.050310E-18
214.307
1.302218E-02
3.053294E-02
1.205017E-08
                                                                   HCL.49
FE203
                                                                  NH4CL
AGAMCL
N2H4
N2H4AG
N3H5O3
AGMYNT
                                                                                                              1.933660E-03
 NOTF--VALUES OF-1.0 INDICATE THAT THE TERM
COULD NOT BE CALCULATED BECAUSE ERSENTIAL
DATA HERE NOT SUPPLIED
                                                                  PARACHOR
                                         N204
73.790115
                                                                                                               HNO3
                                                                                                            124.280390
                                       MMH
127.675718
```

Figure D-43. Derived Coefficients and Properties of Deposit Constituents (Page 2 of 5)

MMMN03

MHN. AD

-1.00000	-1.000000
WATER	AMONIA
52.501303	51.429413
NH40H	NM4ND3
112.931140	178.697982
00.182810 0102810	#XFUAG -1.800000
AL203	MCL -1.00000
HCL.40	FE203
50,893141	-1.00000
CARBON -1,000000	-1.00000
AGAMCL	N2M4
250,90647	91.029603
H2H4AQ	N3H503
89.899934	-1.00000
49HYNT -1.00000	

NOTE--VALUES OF-1.0 INDICATE THAT THE TERM COULD NOT SE CALCULATED SECAUSE ESSENTIAL DATA MERE NOT SUPPLIED

DERIVED WMM PROPERTIES

.300000 315.000 112,985 462,388 5.055018E-00 .988503 109998 .400000 420.800 250.080 507.002 1.513490E-02 .932980 2.881702E-03 .500000 520.000 329,720 671,736 .8203.7 .877397 8.113390E-04	3.234435E-04 2.567667E-04 2.008375E-04
.500000 520.000 32°,724 671,736 ,820367 ,877397 8.113340g-84	2.0083256-04
.000000 631.200 403.366 776.410 10.6515 .021814 2.0803862-04	1.545765E-04
.700000 736.400 477,006 681.084 61.0007 .766231 1.769939E-04	1.100122E-04
.800000 841.600 550.646 485.758 210.066 ,71048 1.162207g-04	0.643056E-05
.900000 940.000 624.286 1090,43 \$63,530 .329241 8.019660g-05	4.2234506-05
.444444 1057.00 647.425 1195.10 1194.49 1.138293E-03 5.754644E-05	4.092751E-15
1.00000 1052.00 944.514 946.516 1195.00 2.910106F-07 1.294791E-05 (CRITICAL TEMP.) (CRITICAL PRESS.)	1.256507E-31
1.01000 1002,52 1205,57 0, 1,3077396-65	

DERIVED NEGA PROPERTIES

REDUCED TEMP	. TEMPERATURE (DEG R)	LIG. ENTHALPY (BTU/LB)	VAP. ENTHALP (BTU/L8)	Y VAPOR PRESSURE (PSIA)	SPECIFIC GRAVITY	VISCOSITY (LBM/FT-SEC)	SURFACE TENSION (LOM/IN)
.300001	232.450	60.7114	341.734	2.746280E-10	1.84410	10000	4.372654E-04
.400000	310.500	91.0152	364.638	7.099040E-05	1.74041	10000	3,4691022-04
.500000	340.200	114.519	307.542	4.096244E-02	1.03073	109998	2.713390E-04
	465,840	137.423	410.445	1.77267	1.53304	109998	2.088439E-04
.70000	543.480	261.647	433,349	21.0645	1.42935	2.594643E-04	1.578216F-84
.00000	021.120	319.077	456,253	131.595	1.32567	1.703739E-04	1.1677302-04
.900000	698.760	374.107	479.157	505.757	.614177	1-1756356-04	5.706452E-05
	774.399	434.334	502,000	1441.29	2.1234102-03	8.4340246-05	1.220495E-14
1.00000	776.400 (CRITICAL TEMP.)	469.199	469.199	1441.30 (CRITICAL PRESS.)	5,4206002-07	1.8980998-05	1.6476312-31
1.01000	784.164		504.351	*******	٥,	1.9170006-05	**********

Figure D-43. Derived Coefficients and Properties of Deposit Constituents (Page 3 of 5)

^	۰	a	0	•

		DERIVED -ATER					
REDIJCED TEMP.	TEMPERATURE (DEG R)	LI9. ENTHALPY (BTU/LP)	VAP. FNTHALPY (BTU/LB)	VAPOR PRESSURE (PSIA)	SPECIFIC GRAVITY	VISCOSITY (LBM/FT-SEC)	SURFACE TENSION (LGM/TN)
.300000	349.500	171.255	1374.61	1.192027E-05	1.09982	109998	4,846761E-04
.400000	466.000	224.340	1435.95	2.803#166-02	1.03798	107778	4.7972768-04
.500000	582.500	475.240	1492.29	1.03643	.976141	4.3787756-04	3.7522346-04
	.44.000	591.790	1548.63	25.0844	.414302	1.554545E-04	2.00015F-04
.700000	815.500	704,240	1004,97	150.111	.452464	4.552362E-05	7,182449E-84
.00000	932.000	#24,790	1001.31	551.534	.790626	6.272435E-05	1.614817E-04
.90000	1048.50	941.290	1717.05	1402.97	, 506294	4.3201846-05	7,4912146-05
. • • • • • •	1165.00	1057.79	1773.99	3225.98	1.2063996-03	3.1057826-05	1.6988358-14
1.00000	1105.00 (CHITICAL TEMP.	1415.89	1415.49	3220.00 CRITICAL PRESS.)	3,2376156-07	6.9879876-06	2.347583E-31
1.01000	1170.05	•••••	1779.62	******	0.	7.057847E-04	••••••

FUEL VAPOR HEAT CONTENT (RTU/LH)	(HTU/LA)	WATER VAROR MEAT CONTENT (RTU/LB)	(AT 540 DEG R)
453.60000	170.10000	1102.30000	

CALCULATED VALUES AT	TEMPERATURES OF HMM	= 1008.000	OFG #. N204 #	491.760 DEG	۸,
PPH PRESS (PSTA)	N2DH PRESS (PSIA)	(18/0	UFT) (LR/	CUFTS	/ N204 DENSITY RATIO
1195.00000	1441.30000	14.027		100	0,00058
.000098	V204 V19C. M (L94/FT-9EC) .0003555	9000, TEN (LBM/IN) .0029053		NS.	

TEMPFRATURE IMPUT DE SAG.000 DEG R TO SUBROUTINE -MEBREM-. CALLED FROM -OVERLAY(2.0)- FOR CALCULATION OF REDUCED MEDIEFO IS NOT SIGNIFICANTLY DIFFERENT FROM A PREVIOUS IMPUT FOR THE SAME SUBSTANCES. THE NEW VALUE WAS REEN MESET TO THE PARVIOUS FRIEN OF SOG.OOD DEG R.

	MMH + 1	4204	
	CONTAMINANT PRO	PERTIES	
	CESTIMATED))	
		VLR-DEG OF VAPORIZATION (ne R) 000-000 100.00000 900.00000 SITY 2 VISCOSITY 3 VISCUSITY 4 CT MM 18 PCT MM 27 PCT MM 000304 .000344 .000383 SITY 6 VISCOSITY 7 VISCOSITY 8 CT MM 55 PCT MM 64 PCT MM 00044 .000502 .00044 ITY 10 VISCOSITY 11 VISCOSITY 12 CT MM 91 PCT MM 100 PCT MM	
DENSITY	VAPOR CP.	LATENT HEAT	DECOMP. TEMP.
(Le/CUFT)	(HTU/LR-DEG)		
62.427961	1.000000	180.00000	
VISCOSITY 1	V15C051TY 2	viscosity 3	VISCOSITY 4
O PCT MMH	9 PCT MMH	18 PCT MMH	
.0002-5	.000304		
VISCOSITY 5	VISCOSITY 6	VISCOSITY 7	VISCOSITY A
36 PCT MMH	45 PCT MMH		
.000423	.000462		
VISCOSITY .	VISCOSITY 10	VISCOSITY 11	VI8COSITY 12
73 PCT MMH	82 PCT MMH		
.000580	.000020		
VISCOSITIES	AME GIVEN LE(MASS)	/FT-SET	

HINTURE OF AA = AMONIA AND BR = N2H4

Figure D-43. Derived Coefficients and Properties of Deposit Constituents (Page 4 of 5)

ESTIMATED MIXTURE VISCOSITIES

0 PCT **AA** 17 PCT **AA** 53 PCT **AA** 50 PCT **AA** .030*12

07 PCT **AA** 83 PCT **AA** 100 PCT **AA** .021050 .011200 .001344

VISCOSITIES ARE GIVEN LH(MASS)/FT-SEC

HICKORY-DICKORY BAYS 481.40 SECONDS FOR BUNNING OVERLAY(2.8) FINISMED AT CLOCK TIME 11-39-33.8

> CP TIME USED = 156.35 SECONDS PP TIME USED = 106.96 SECONDS SINCE JOB STARTED

Figure D-43. Derived Coefficients and Properties of Deposit Constituents (Page 5 of 5)

REVISED DATA INPUT WITH NAMELIST

SNAST3

TOTIMS = 0.2E-02.

SEND

SHAST4

P1 = 0.142E+00.
DELP1 = 0.1E-02.
NTR1 = 2.

HICKORY-DICKORY SAYS 6.00 SECONDS FOR RUNNING OVERLAY 2:1 FINISHED AT CLOCK TIME 11-39-39.0

> CP TIME USED = 156.64 SECONDS PP TIME USED = 107.95 SECONDS SINCE JOB STARTED

Figure D-44. Revised Data Input by NAMELIST

CYCLE CYCLE	EXPOSURE /OPERATION	DURATION (SEC.)
1	INITIAL	-1.0000
Ž	ASSIGNED	-1.0000
3	PLUME	70.000
4	SPACE	6.00000E+05
5	WRAP-UP	-1.0000

Figure D-45. Assignment of Calculation Cycles

Spacecraft Exterior Materials and Conditions, Initial State

The assignment of materials and their thickness, surface temperature, and surface finish at specified segments of the satellite is performed in Link 3,0 with data input from file BDATA. This input is listed for inspection in Figure D-46.

Initial Condition of Satellite Exterior

The data on the assignment of structural materials to the satellite and the properties of the materials are used by Link 11,0 and its subsidiary programs to calculate the condition of each segment of the surface. The results of the calculations include the solar absorptivity, thermal emissivity, diffuse and specular solar reflectivity, thermal reflectivity, thermal conductivity and transmissivity at several wavelengths. This information is listed for each segment (see Figure D-47 for a portion of this listing).

The condition is then summarized in a listing of the average solar absorptivity, thermal emissivity, and their ratio for the whole satellite. Then the percent transmission for a set of 8 different wavelengths of light through all transparent segments is listed. These outputs are presented in Figure D-48.

Assigned Satellite Conditions

The assigned post-exposure conditions for specific satellite locations, input from file TAPE9 (CDATA) by execution of Link (4,0), and used to test the functioning of the subsequent calculation routines in Links (7,0) and (11,0) are listed next. This input can be the satellite condition data generated in a previous run of the program, but that feature is not used in the Test Case run. The input is listed, as shown in Figure D-49. These data are then used to calculate the condition of the exterior, and then a segment-by-segment printout of the condition is again made. Figure D-50 shows the results of the assigned post-exposure condition at three of the segments, numbers 51, 52, and 53, identified in Figure D-49. The data in Figure D-50 consists first of the condition for the three segments in the pristine or original condition, then the condition for the same segments after the assigned exposure. The changes are those caused by the alteration of the surface condition.

The summary of current conditions, Figure D-51, includes not only the average α and ε values, and their ratio, but also the percent change in the ratio compared to the initial or baseline condition.

The execution then enters Link (10, 2) and checks to see whether the assigned conditions should be used as the base for further calculations or whether the initial conditions should be reestablished. The flag ISTST, read from file TAPE9, is zero, so the initial conditions are reestablished.

Plume Exposure Sequences

The total exposure duration and the calculation time increment are listed, see Figure D-52. Note that these are the true values to be used; in this case they are the specific corrected values supplied by NAMELIST/NAST3, not those input from ADATA.

SPACECRAPT EXTERIOR MATERIALS SEEK VALUES INPUT FOR INITIAL COMOITIONS

SEGMENT NO.	TEMPERAT	MATERIAL	(IN)	(MU-IN)	L DEPTH	(MI-1M)	3 - R D MATERIAL		inish (MU=IN)
1	520.00	SOLO	.0003	1.00	ALUMIN	.1000	2.00	NONE	-1.0	-1.0
į	520.00	SOLD	.0003	1.00	ALUMIN	.1000	2.00	NONE	-1.0	-1.0
;	520.00 520.00	COLD	,0003	1,00	ALUMIN	.1000	2.00	HONE	-1:0	•1.0 •1.0
•	520.00		.0003	1.00	ALUMIN	.1000	2.00	NONE	-111	-1.0 -1.0
:	520.00	#1K00#	1.5000	5.00	NONE	-1.0	-1.0	NONE	-1.0	-1.0
10	520.00	ALUMIN	.1000	10.00	HONE	-1.0	2.00 -1.0	HONE	-1.0	-1.0
11	520.00	ALUMIN ALUMIN	.1000	10.00	NONE	-1.0	-1.0	NONE	-1.6	•1.0 •1.0
13	520.00		.1000	10.00	NONE	-1.0	-1.0 -1.0	NONE	•1.0	•1.0
15	520.00	ALUMIN	.1000	10.00	HONE	-1.0	-1.0	NONE	-1.0	-1.0
17	520.00	ALUMIN	.1000	10.00	NONE	-1.0	-1.0	NONE	-1.0	-1.0 -1.0
10	520.00	ALUMIN	.1000	10.00	NONE	-1.0	-1.0	NONE	-1.0	-1.0 -1.0
20	520.00	ALUMIN	.1000	10.00	NONE	-1.0	-1.0	NONE	-1.0	-1.0
55	520.00	ALUMIN ALUMIN	.1000	14.00	NONE	-1.0	-1.0	NONE	-1.0	-1.0
23	520.00	IRPORT	,2500	2.50	NONE	-1.0	-1.0	NONE	-1.0	-1.0
25	520.00	ALUMIN	.1000	10.00	HOME	-1.0	-1.0	NONE	-1.0	-1.0 -1.0
26 27	520.00 520.00	ALUMIN	.1000	16.00	ndae Hone	-1.0	-1.0 -1.0	NONE	-1.0 -1.0	-1.0 -1.0
24 29	520.00	ALUMIN	.1000	10.00	NONE	-1.0	:1:	NONE	-1:	-1.0 -1.0
34	520.00	ALUMIN	.1000	10.00	NONE	-1.0	-1.0	NONE	-1.0	-1.0
35 31	520.00	ALUMIN	.1000	10.00	NONE	-1.0 -1.0	-1.0	HONE	-1.0	-1.0
33 34	520.00	ALUMIN ALUMIN	-1000	10.00	NONE	-1.0	-1.0	NONE	-1.0	-1.0
35	520.00	ALUMIN	.1000	10.00	HONE	-1.0	-1.0	NONE	-1.0	-1.0
36 37	520.00	ALUMIN ALUMIN	.1000	10.00	HOHE	-1.0	-1.0	NONE	-1.0	-1:0
38 39	520.00	ALUMIN	.1000	10.00	NONE	-1.0	-1.0	NONE	-1.0	-1.0
40	250.00	ALUMIN	.1000	10.00	NONE	-1.0	-1.0	NONE	-1.0	-1.0
41	526.00 520.00	BHTTE	.0050	32.00	MONE	-1.0	10.00	NONE	-1.0	-1.0
43	520.00	MUSTE	.0050	32.00	ALUMIN	.1000	14.00	NONE	-1.0	-1.0
45	520.00	ALUMIN	.1000	10.00	NONE	•1.0	-1.0	NONE	-1:	-1.0
46	520.00	ALUMIN	.0050	32.00	ALUMIN	-1.0	-1.0	NONE	-1.0 -1.0	-1.0
**	520.00	BLUMIN	.0050	32.00	ALUMIN	-1.0	14.80	NONE	-1.0	-1.0 -1.0
50	520.00	ALUMIN	.1000	10.00	NONE	-1.0	-1.0	NONE	-1.0	-1.0
25 21	520.00 520.00	ALUMIN	.0050	32.00	ALUMIN	-1.0	16.60	HONE	-1.0 -1.0	-1.0
53 54	520.00	BLUMIN	.0050	32.00	VONE	-1.0	16.00	NONE	-1.0	-1.0
55	250.00	ALUMIN	.1000	10.00	NONE	-1.0	-1.0	NONE	-1.0	-1.0
56 57	520.00	ALUMIN	.1000	32.00	MONE	1000	-1.0	NONE	-1:0	-1.0
58 59	520.00	MHITE	. 1000	32.00	ALUMIN	-1.0	16.00	NONE	-1.0 -1.0	-1.0 -1.0
• 0	520.00	ALUMIN	.1000	10.00	HONE	-1.0	-1.0	HONE	-1.0	-1.0
91	520.00	ALUMIN	.1000	32.00	ALUMIN	-1.0	-1.0	NONE	-1.0	-1.0
63 64	520.00	MHTTE	.1000	32.00	MOME VECHIN	-1.0	-1.0	NONE	-1.0	-1.0
65	520.00	ALUMIN	.1000	10.00	ALUMIN	.1000	-1.0	NONE	-1.0	-1.6
67	520.00	ALUMIN	.1000	10.00	HONE	-1.0	-1.0	NONE	-1.0	-1.0
**	320.00	ALUMIN	.1000	14:50	NONE	-1.0	-1.0	NONE	-1:	-1.0
70 71	520.00	BLACK	.0070	12.00	ALUMIN	.1000	16.00	NONE	-1.0	-1:
72	520.00	BLACK	.0070	48.00	ALUMIN	1.5000	10.00	NONE	-1.0	-1.0
73 74	520.00	DLACK	.0050	32.00	ALUMIN	1.5000	10.00	NONE	-1.0	-1.0 -1.0
75 76	520.00		.0074	48.00	ALUMIN	.1000	10.00	NONE	-1.0	-1.0
77	520.00	PLACK	.0070	48.60	ALUMIN	.1000	14.00	NONE	-1.0	-1.0
78 79	520.00	BLACK	.0050	32.00	ALUMIN	.1000	16.00	HONE	-1.0	-1.0 -1.0
**	\$20.00	BLACK	.0070	10.00	ALUMIN	:::::	1:::	NONE		:::
95	520.00	BLACK	.0070	48.00	VEANIN	.1000	16.00	NONE	-1.0	-1.0
**	520.00	BLACK	:0070	10.00	ALUMIN	1:000	14.00	NONE	:1::	:1::
85	520.00	BLACK	.0070	48.00	ALUMIN	.1000	16.00	NONE	-1.0	-1.0
87	520.00	BLACK	.0070	48.00	ALUMIN	.1000	14.00	NONE	-1.0	-1.6
••	250.00	MHITE	.0050	36.00	ALUMIN	.1000	16.00	HONE	-1.0	-1.0

Figure D-46. Initial Material Assignment to Spacecraft Exterior (Page 1 of 2)

CR99A ALUMIN ALUMIN ALUMIN ALUMIN 16.00 80 90 91 92 93 94 95 96 97 98 100 \$20.00 \$20.00 \$20.00 \$20.00 \$20.00 \$20.00 \$20.00 \$20.00 \$20.00 BLACK BLACK BLACK BLACK BLACK BLACK BLACK RLACK RLACK 48.00 48.00 48.00 48.00 48.00 48.00 48.00 48.00 32.00 32.00 32.00 32.00 32.00 32.00 32.00 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 .0070 .0070 .0070 .0070 .0070 .0070 .0070 .0050 .0050 .0050 .0050 .0050 .0050 .0050 .1000 .1000 .1000 .1000 .1000 .1000 .0075 .0075 ALUMIN ALUMIN GOLD ALUMIN -1.0 ALUMIN ALUMIN ALUMIN ALUMIN ALUMIN ALUMIN ALUMIN ALUMIN ALUMIN ALUMIN ALUMIN ALUMIN ALUMIN WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WAS A SHOULD WHITE WHITE WHITE WHITE WHITE WHITE WHITE WHITE WAS A SHOULD WHITE WHITE WAS A SHOULD WHITE WHITE WAS A SHOULD WHITE WAS A SHOULD WHITE WAS A SHOULD WHITE WAS A SHOULD WHITE WHITE WAS A SHOULD WHITE WAS A SHOULD WHITE WAS A SHOULD WHITE WAS A SHOULD WHITE WAS A SHOULD WHITE WAS A SHOULD WHITE WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WAS A SHOULD WA .1000 520.00 520.00 520.00 520.00 520.00 520.00 520.00 520.00 102 103 104 105 100 107 108 110 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 .1000 .1000 .1000 .1000 1.00 520.00 520.00 520.00 520.00 520.20 520.20 550.00 560.00 560.00 560.00 560.00 .0003 1.00 .1000 -1.0 BOLD GOLD GOLD GOLD GOLD 115 1.00 1.00 20.00 20.00 10.00 ALUMIN ALUMIN ALUMIN ALUMIN UNASNO UNASNO .1000 .1000 .0210 .0205 2.00 2.00 32.20 32.20 -1.0 -1.0 -1.0 -1.0 -1.0 .0003 -1.0 7301 7301 7302 7303 DLACK TEFLON SOLCEL SOLCEL SOLCEL SOLCEL .0000 .0500 .0500 .0500 .0500 .0500 .0500 .0500 .0500 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 10.00 -1.0 UNASND 7304 7305 7306 7307 7306 7309 7310 7311 7312 UNASNO UNASNO UNASNO UNASNO SOLCEL SOLCEL SOLCEL SOLCEL SOLCEL SOLCEL -1.0 -1.0 -1.0 -1.0 -1.0 -1.0 500.00 510.00 510.00 UNASND UNASHD UNASHD UNASHD 7313 7314 510.00 SOLCEL SOLCEL SOLCEL SOLCEL SOLCEL SOLCEL TEFLON .0500 10.00 .0500 10.00 .0500 10.00 .0500 10.00 .0500 10.00 .0500 10.00 UNASHD UNASHD UNASHD UNASHD UNASHD UNASHD UNASHD UNASND UNASND UNASND UNASND UNASND UNASND UNASND -1.0 -1.0 -1.0 -1.0 7315 7316 510.00 7317 7318 7319 7320 510.00 510.00 510.00 520.00 NONE NOTE VALUES OF -1.0 INDICATE NO DATA ENTERED.

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HICKORY-DICKORY SAYS 4.88 SECONDS FOR RUNNING OVERLAY(3.8) FINISHED AT CLOCK TIME 11-19-43.8

CP TIME USED = 158.10 SECONDS PP TIME USED = 109.90 SECONDS SINCE JOB STARTED MAXIMUM EXECUTION FIELD LENGTH REQUIRED = 202406

HICKORY-DICKORY SAYS 1028.66 SECONDS FOR RUNNING OVERLAY 11.1 FINISHED AT CLOCK TIME 12-10-12.6

> CP TIME USED = 171.99 SECONDS PP TIME USED = 157.44 SECONDS SINCE JOB STARTED

HICKORY-DICKORY SAYS 592.20 SECONDS FOR RUNNING OVERLAY 11.2 FINISHED AT CLOCK TIME 12-20- 4.8

> CP TIME USED = 174.05 SECONDS PP TIME USED = 166.46 SECONDS SINCE JOB STARTED

Figure D-46. Initial Material Assignment to Spacecraft Exterior (Page 2 of 2)

CROOA

CURRENT CONDITION OF BATELLITE EXTERIOR

INITIAL CONDITIONS

```
THETA AREA PLUME X
                           TEMP PRIN HET LAMBOA MU
  1 0.000 0.000 0.0000 1 -3.000 550.0 -1.0 -1.0 -1.0 -1.0 -1.0
       DEPTH FINISH ABBRY EMISY DIFRF SPECRF TRMRF TRMCHD RFIND 15 $ 1 .5
                                                  .2 .05 .01 TEMP
THETA AREA PLUME X
                          TEMP PRIM HET
                                    LAMBDA HU
  THETA AREA PLUME X TEMP PRIN HET
  3 ,305 ,590 3.14159 1 -3,295 520.0 -1,0 -1,0 -1,0 -1,0 -1,0
PEATIE! SMITTED
              THETA AREA PLUME Y
          .
                           TEMP PRIN HET
  6 .020 1.200 4.39016 1 -2,960 520.0 -1,0 -1,0 -1,0 -1,0
MATRL NAME DEPTH FINISH ABSRY EMISY DIFRE SPECIF TRHIFF TRHICHD RFIND IS $ 2 1 .S .2 .05 .01 TEMP
NO SECOND LAYER
              THETA AREA PLUME N
                           TEMP PRIN HET
  9 .020 1.200 5.05400 1 -2.900 520.0 -1.0 -1.0
                                         '-i.i ' -1.i.
      DEPTH FINISH ABBRY EMISY DIFRF SPECRF TRWRF TRHCHD RFIND 15 S 2 1 .5 .2 .05 .01 TEMP
      THIRD LAVER
              THETA AREA PLUME X
                          TEMP PRIM HET LAMBOA MU
  10 1,325 1,600 ,31410 1 -2,275 920,0 -1,0 -1,0 -1,0 -1,0 -1,0
MATRL NAME DEPTH FINISH ABBRY EMISY DIFFF SPECRF TRMRF TRMCND RFIND IS $ 1 .5 .2 .05 .01 TEMP
' î ÎLÛMÎN' "10000000" "10.00" 1230" 1040" 1329" 1764" 1910" 1930" "0.0" "0.0" "0.0" "0.0" "0.0" "0.0" "0.0" "520"
NO SECOND LAYER
NO TMIRD LAYER
```

Figure D-47. Satellite Exterior Conditions (Page 1 of 3)

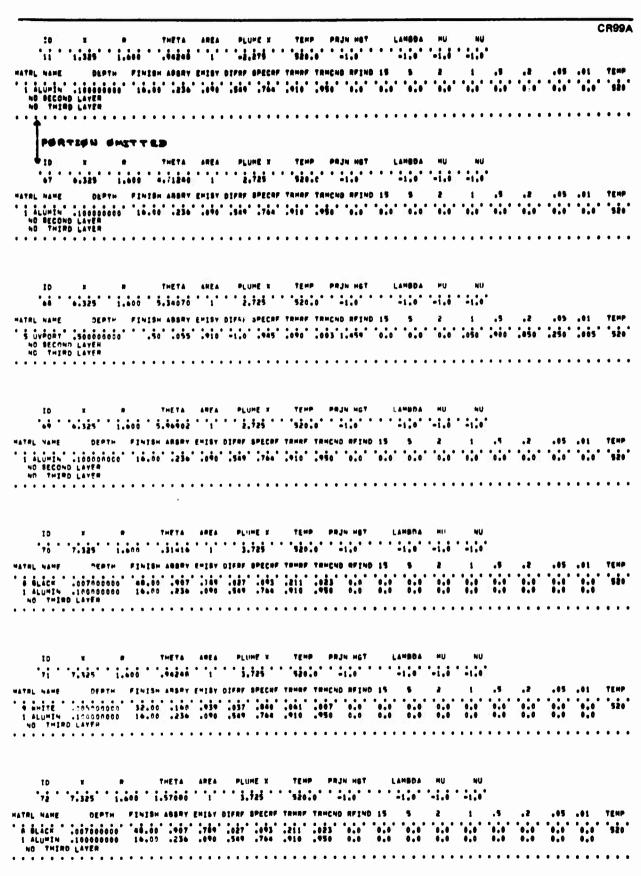


Figure D-47. Satellite Exterior Conditions (Page 2 of 3)

CR99A PORTED N SMITTED **→** 10 THETA AREA PLUME X TEMP PRJN HST LAMBOA HU .05 .01 TEMP 10 THETA AREA PLUME X TEMP PRIN HAT LAMBOA HU 7301 6.866 2.502 2.19911 1 5.266 550.0 -1.0 -1.0 -1.0 -1.0 -1.0 DEPTH FINISH ABORY EMISY DIFMF SPECAF TRHAF TRHCHD AFIND 15 5 1 ,5 ,2 ,05 ,01 TEMP 2 10 X L THETA AREA PLUME N TEMP PRJN HET LAMBOA MU 7302 ' 6.668 ' 4.161 ' 1.49900 ' '1' ' 3.668 ' ' 560.0' ' -1.0' ' -1.0' ' -1.0' ' -1.0 ' -1.0' MATRL NAME DEPTH FINISH ABBRY ENISY DIPRF SPECRF TRHRF TRHCHD RPIND 15 5 2 1 . 5 " \$ \$0(cel. '.osococo' '10.00' '100' '050' -1:0' '000' '150' '.os'' '0;0' '0;0' '250' '.oso' '0;0' '0;0' '0;0' '0;0' '540' NO THIRD LAYER * THETA AREA PLUME H TEMP PRIN HET LAMBDA MU 7303 ' '6:603' ' 4,056 ' 2,10011 ' '1' ' 3,003 ' ' '500;0' ' -1;0' ' -1;0' '-1;0' '-1;0' MATRL NAME DEPTH FIMISH ABBRY EMISY DIFFR SPECAF TANAF THUCHD AFIND 15 5 2 1 .5 .2 .05 .01 TEMP SECOND LAVER PERTION SMITTED NOTE VALUES OF -1.0... O INDICATE NO DATA ENTERED

Figure D-47. Satellite Exterior Conditions (Page 3 of 3)

EFFECTIVE HEAT TRANSPER CONSTANTS

SUM OF SESMENT AREAS = 125.00

EFFECTIVE BOLAR ABBORPTIVITY . .348 EFFECTIVE THERMAL EMISSIVITY . .423

AVERAGE VALUE-ALPHA/EPSILON RATIO FOR WHOLE SPACECRAFT = .8232

PERCENT TRANSMISSION THROUGH TRANSPARENT SESMENTS

BESMENT	HATERIAL	1540	540	240	140	.540	.240	.05MU	.014
	HOOMIN		••••••	1.418	20.70	49.87			
23	IRPORT	22.98	54.21	80.40	80.40	50.55			
23	UVPORT				19,53	89.25	86.74	47.04	6.652
7302	BOLCEL		•	76.14	79.86	92.00	•		450
7303	BOLCEL	•	•	70.14	79.86	92.00	Ī	•	
7304	SOLCEL	•	•	76.14	79.86	92.00	•	•	•
7305	BOLCEL	•	•	70.14	79.66	92.00	•	•	
		•	•				•	•	•
7306	BOLCEL	•	•	76.14	79.86	92.00	•	•	•
7307	BOLCEL	•	•	76.14	79.86	92.00	•	•	•
7308	BOLCEL	•	•	76.14	79.06	92.00	•	•	•
7309	BOLCEL	•	•	76.14	79.86	45.00	•	•	•
7310	BOLCEL	•	•	70.14	79.86	45.08	•	•	•
7311	BOLCEL	•		74.14	79.86	92.08	•	•	•
7312	BOLCEL			76.14	79.86	92.00	•		•
7313	SOLCEL			76.14	79.86	45.64			
7314	BOLCEL	- 1	Ĭ	76.14	79.86	92.00			
7315	SOLCEL	•	•	76.14	79.86	92.0A	•	•	•
7316	SOLCEL	•	•	70.14	79.86	92.00	•		•
		•	•		79.86	92.00	•	•	•
7317	BOLCEL	•	•	76.14			•	•	•
7318	BOLCEL	•	•	76.14	79.86	92.00	•	•	•
7319	BOLCEL	•	•	76.14	79.86	45.00	•	•	•
WE	AN SOLCEL	•	•	76.14	79.86	45.08	•	•	•

MICKORY-DICKORY SAYS 328.80 SECONDS FOR RUNNING OVERLAY 11.8 (PRIMARY OVERLAY ONLY)

HICKORY-DICKORY SAYS 2749.80 SECONDS FOR RUMNING OVERLAY 11.0 INCLUDING ALL SECONDARY OVERLAYS FINISHED AT CLOCK TIME 12-25-33.0

> CP TIME USED = 193.14 SECONOS PP TIME USED = 198.85 SECONOS SINCE JOS STARTED

Figure D-48. Summary of Initial Satellite Condition

SPACECRAFT EXTERIOR MATERIALS

ASSIGNED CONDITIONS

BEGMENT NO.	TEMPERATURE (DES 9)	TOP LA MATERIAL DEPTH (IN)		Y E R FINISH (MU-IN)	MATERI	LAYER FINISH (MU=IN)	
42 51 52 53 61 62 63 72 62	000.0 610.0 650.0 610.0 700.0 700.0 430.0 610.0	ALUMIN ALUMIN ALUMIN ALUMIN ALUMIN ALUMIN ALUMIN ALUMIN ALUMIN ALUMIN ALUMIN ALUMIN ALUMIN ALUMIN ALUMIN	2.0006-03	80.00 250.00 -1.00 50.00	NONE ALUMIN NONE NONE NONE NONE NONE NONE NONE NO	-1.0 .100 -1.0 .100 -1.0 -1.0 -1.0	-1.0 39.00 -1.0 39.00 -1.0 -1.0 -1.0 -1.0

NOTE VALUES OF -1.0 INDICATE NO DATA ENTERED.

HICKORY-DICKORY SAYS 7.80 SECONDS FOR RUNNING OVERLAY 7.0 FINISHED AT CLOCK TIME 12-25-48.0

> CP TIME USED = 195.07 SECONDS PP TIME USED = 204.19 SECONDS SINCE JOB STARTED .

HICKORY-DICKORY SAYS 47,40 SECONDS FOR RUNNING OVERLAY 11.1 FINISMED AT CLOCK TIME 12-26-36.0

> CP TIME USED = 202.00 SECONDS PP TIME USED = 222.06 SECONDS SINCE JOB STARTED

HICKORY-DICKORY SAYS 3.00 SECONDS FOR RUNNING OVERLAY 11.2 FINISMED AT CLOCK TIME 12-26-39.0

> CP TIME USED = 202.86 SECONDS PP TIME USED = 223.71 SECONDS SINCE JOB STARTED

Figure D-49. Assignment of Post-Exposure Conditions

```
PRISTINE CONDITION
                       PLUME Y
                             TEMP PRIM HET
                                       LAMBDA HU
    '91 '5.325' 1.600' '.74246' 1' '1.725' '520.0' '-1.0' '-1.0' '-1.0' '-1.0'
        DEPTH FINISH ABBRY EMISY DIFFF SPECAF TOWAR TRUCHO RFIND 15 5
                                          2
                                              1 .5 .2 .05 .01 TEMP
     THETA AREA
                       PLUMF Y
                             TEMP PRIN HET
                                       LAMBDA HII
    52 5.325 1.000 1.57000 1 1.725 520.0 -1.0 -1.0 -1.0 -1.0
 MATRL NAME DEPTH FINISH ABBRY EMISM DIPRF SPECRF TRHRF TRHCND RFIND 15 5 1 .5 .2 .05 .01 TEMP
       ALUMIN ,180000
NO SECOND LAYER
NO THIRD LAYER
               THETA AREA
                       PLIME I
                             TEMP PRIM HGT
                                       LAMBOA MU
    53 5,325 1,000 2,19911 1 1,725 520.0 11.0 -1.0 -1.0 -1.0
       DEPTH FINISH ABBRY EMISY DIFRE SPECEF TRANF TRACAD AFIND 15 9
                                          2
                                              1 .5
                                                   .2 .05 .01 TEMP
 RESULT OF ASSIGNED POST EXPOSURE CONDITION
               THETA AREA
                       PLUME Y
                             TON MERG THET
      DEPTH FINISH ABBRY EMISY DIFFF SPECAF TAWAF TANCHO AFIND 15 5 2 1 .5 .6 .61 TEMP
 THIRD LAVER
                             TON MERG 943T
                                      LAPBOA MU
   52 5.325 1.606 1.57686 1 1.725 650.0 -1.6 -1.6 -1.6 -1.6 -1.6
       DEPTH FINISH ARRHY EMISY DIFFF SPECRF TRHAF TRHAF TS 2 1 .5 .5 .01 TEMP
 MATRL NAME
 1 ALUMEN .09500
NO SECOND LATER
NO THIRD LATER
       PLUME X TEMP PRIN HET
    DEPTH FINISH ABBRY EMISY DIFRE SPECAF TAMAF TANCHO RFIND 15 5 2
                                              1
                                                .5
                                                      .05 .01 TEMP
```

Figure D-50. Comparison of Selected Segments Before and After Assignment of Post-Exposed Conditions

EFFECTIVE MEAT TRANSFER CONSTANTS

SUM OF SEGMENT AREAS = 125.00

EFFECTIVE SOLAR ASSORPTIVITY = .357 EFFECTIVE THERMAL EMISSIVITY = .401

AVERAGE VALUE-ALPHA/EPSILON MATIN FOR WHOLE SPACECRAFT # .8888

PERCENT CHANGE IN ALPHA/EPSILON MATID PHOM BASELINE VALUE # 7.470

PERCENT TRANSMISSION THROUGH TRANSPARENT SEGMENTS

SEGMENT	MATERIAL	15MU	5mu	\$#U	1#0	.540	.240	.05MU	.01#0
•	#1400#	•	•	1.418	24.90	49.67		,	•
53	IRPORT	44.55	54.21	80.40	80.40	50.55			•
	UVPORT				19.53	89.25	84.74	47.84	6.652
7362	BOLCEL	•		76.14	79.86	42.00	•		
7303	BOLCEL			76.14	79.66	92.08			·
7304	SOLCEL	•	1	70.14	79.86	92.08		· ·	
7305	BOLCEL			70.14	79.86	92.00		Ĭ	•
7300	BOLCEL		Ĭ	70.14	79.46	92.00		•	•
7307	SOLCEL	:	•	76.14	79.06	92.00	•	•	
7300	SCLCEL	•	•	70.14	79.86	40.50	•	•	•
7309	BOLCEL	•	•	70.14	79.86		•	•	•
		•	•			92.04	•	•	•
7310	BOLCEL	•	•	76.14	79.86	45.00	•	•	•
7311	BUTCEL	•	•	70.14	79.06	45.00	•	•	•
7312	BOLCEL	•	•	70.14	79.86	92.00	•	•	•
7313	SOLCEL	•	•	76.14	79.86	45.00	•	•	•
7314	BOLCEL	•	•	76.14	79.86	45.66	•	•	•
7315	SOLCEL	•	•	76.14	79.86	45.66	•	•	
7316	BOLCEL	•	•	74.14	79.84	42.00			
7317	BOLCEL	•		70.14	79.86	92.00			
7316	BOLCEL			76.14	79.00	92.00	I		i i
7319	BOLCEL			76.14	79.00	92.00			i
	AN BOLCEL	-	-	76.14	79.86	40.50	-		•
		•	•		,,,,,,	45.44	•	•	•

WICKORY-DICKORY SAYS 99,48 SECONDS FOR RUNWING GVERLAY 11.8 (PRIMARY GVERLAY ONLY)

MICKORY-DICKORY SAYS 150.00 SECONDS FOR RUNNING OVERLAY 11.0 INCLUDING ALL SECONDARY OVERLAYS FINISHED AT CLOCK TIME 12-20-10.0

> CP TIME USED = 221.49 SECONDS PP TIME USED = 254.91 SECONDS SINCE JOB STARTED

MICKORY-DICKORY BAYS 5.40 SECONDS FOR RUNNING OVERLAY 10.2 FINISMED AT CLOCK TIME 12-20-24.0

> CP TIME USED = 222.46 SECONDS PP TIME USED = 257.27 SECONDS SINCE JOB STARTED

Figure D-51. Summary of Effects Caused by Assigned Post-Exposure Conditions

The satellite segments impinged by the plume for various combinations of thrustor operation and plume geometry input from file DDATA are then calculated and printed. This facility can be used to identify the segments of the satellite most frequently exposed to plume impingement. The test case has two such demonstrations, and the results are shown in Figure D-53.

The specified test case thrustor i. d. and plume configuration, input by NAMELIST/NAST4, are then used for calculation the i. d. 's of the segments impinged by the plume, to be used for calculating the plume exposure effects. These data are printed and identified as the base for further calculations, see Figure D-54.

The capability of coordinate transformation from thrustor based coordinates is next demonstrated in the output, based on data input by file DDATA. This is an ancillary routine which requires specific inputs of the thrustor coordinates; it does not form part of the calculation of plume impingement effects. The listing from this routine is shown in Figure D-55.

The next listing (Figure D-56) is that of the coordinates of the satellite segments which were identified as being impinged by the plume (Figure D-54). The coordinates are given in the thrustor based coordinate system for use in selecting appropriate plume characteristics from KINCON AND MULTRAN.

This output terminates the run for the case where STPFLG = 1. The program is then rerun with STPFLG = 2 and with the plume characteristics supplied as a new version of file EDATA

RUN WITH STPF6G = 2 FØR CALCULATION OF EXPOSURE EFFECTS

The plume characteristics data used as EDATA input for this run are shown in Figure D-57.

D.6 REFERENCES

- D-1. K. B. Wiberg, "Subroutine LØCATN," Computer Programming for Chemists, W. A. Benjamin, Inc., New York (1965).
- D-2. J.H. Neilson and A. Gilchrist, Wear, Vol. 11, pp. 111-122 (1968).
- D-3. R. E. Trebal, Illustration 3.6 Mass Transfer Operations, McGraw Hill, 2nd Edition (1968).

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71 72 73 A?	
90 91 92 93	
04 170 101 102	
Projection segments liplinged by thiluston Mo. 2	ины: Р в 1,250
7304 7402 7868 7865 7306 7408	
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FROM THRUSTOR AG, 1 MMEN PO .2. IC AGS,	
PPOLECTION SECHENTS IMPINGED BY THRUSTOR NG. 1	WFET. F = .200
	FICKCRY-CICKORY SAVS 0.00 SECONDS FOM PLP. I'M OVERLAY(3.1) FINISHED AT CLOCK TOPE 0-36-58,8
	CP TIPE USED = 177.40 SECONDS PP TIPE USED = 249.29 SECONDS SINCL JOS STARTED

Figure D-53. Segments Impinged for Selected Thrustors and Plume Configurations

FROM THRUSTOR NO. 2 WHEN PO .34	
PROJECTION SEMBATS IMPINED BY THRUSTOR NO. 2 WHEN F = ,142	
7302	
	HICKORY-DICKORY SAYS 1.20 SECONDS
	FINISHED AT CLOCK TAPE 0-37- 0.0
	CP TIME USED = 177.40 SECONDS PP TIME USED = 245.44 SECONDS
ABRIATOR TUBERTO PAR SIREPALENC ALIGNATURA DA PEPPER	61486 700 -0144455

Figure D-54. Segments Impinged by Plume

COORDINATE TRANSFORMATION

THRUSTOR SYSTEM TO SATELLITE SYSTEM

THRUSTOR NUMBER 3

AXIAL CO	. STANION	RADIAL	CORRDINATE		ANGULAR	COORDINATE
22	и .	RR	R		THE	THETA
(FT)	(FT) .	(FT)	(FT)	•	(RAD)	(RAD)
*********				•		
4.0700	7.6700 .	1.500	0 3.5895		2.9870	1225
5.1000	8.7000 .	1.000	0 1.1023		.0500	0240
1.7300	5.3300 .	1.000	3,1000		3.1416	.0004
DE	CITAGTEMO	M CALCULA	TION			

HICKORY-DICKORY BAYS 1.20 SECONDS FOR RUNNING OVERLAY(5.2) FINISMED AT CLOCK TIME 12-28-40.8

> CP TIME USED = 224.49 SECONDS PP TIME USED = 263.65 SECONDS SINCE JOS STARTED

MICKORY-DICKORY SAYS 0.00 SECONDS FOR RUNNING OVERLAY(5.3) FINISMED AT CLOCK TIME 12-28-40.8

> CP TIME USED = 224.53 SECONDS PP TIME USED = 264.17 SECONDS SINCE JOB STARTED

Figure D-55. Transformation of Coordinates, Thrustor Based to Satellite Based

THRUSTOR-BASED COORDINATES
OF FLUME-IMPIRGED SEGMENTS

"" from "thmustor ito" 2 "When the 114

SEG AXIAL RADIAL ANGULAR
ID DIST DIST COCRD
2 (F1) R (FT) THE(RAD)
7302 3.0022 .71581 1.13957

* ANGLE FROM LINE JOINING AXES OF SATELLITE ALC THPUSTON

USE COOFDINATES Z AND R IN DETERMINATION OF PLUME CHARACTERISTICS FOR INPUT TO REPLACE DEFAULT VALUES CO EDATA = TAPE11

Appendix E

HYDRAZINE MONOPROPELLANT ENGINE ANALYSIS

N2H4

A VAPORIZED MONOPROPELLANT AND INITIAL PULSE MODEL COMPUTER PROGRAM

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Appendix E

HYDRAZINE MONOPROPELLANT ENGINE ANALYSIS

A Vaporized Monopropellant and Initial Pulse Model Computer Program

E. 1 INTRODUCTION

The current program N2H4 is used to model the contaminant production, i.e., the hydrazine droplets, hydrazine vapor, and ammonia from an N₂H₄ catalytic reactor.

The program N2H4 is a modification of the code written by Kesten, Smith, and Smith (Reference E-1) at the United Aircraft Research Laboratories (UARL). The UARL program was based on a model of the vapor phase region only. The modeling of a monopropellant engine considering only the vapor phase places its utility in the same state as models of a bipropellant system starting with premixed gases. That is, a very substantial amount of the phenomena critical to the process are excluded from the model.

MDAC has modified the UARL program with the addition of the following elements.

- a. Time-dependent inertial-resistive feed line modeling
- b. Time-dependent injection stream velocities
- c. The estimation of cold start spikes from empirical UARL data
 - 1. Time to pressure rise
 - 2. Peak Pressure
 - 3. Propellant accumulation
 - 4. Contaminant discharge (associated with initial cold start transient)

The program with the current additions can be utilized for either cold or hot starts. The initial conditions which are given as input permits the program to determine whether the start will be smooth (hot) or rough (cold). If a smooth start is indicated, the program calculates the start and steady state operation in the manner programmed by Kesten. If a rough start is indicated, the program provides working information about the initial pressure surge, but it is incapable of following this with the subsequent

steady operation; it shuts down and prints out the characteristics of the rough start.

a. Estimating cold start transients

The following parameters are estimated by correlations of UARL data.

- 1. Time to pressure rise
- 2. Peak pressure
- 3. Contaminant discharges associated with cold start transient.

b. Hot starts

Hot starts and the subsequent steady operation may be simulated with the modified program to the same level of accuracy as in the original Kesten program. The program still contains the UARL assumption of "instantaneous evaporation" and hence overestimates the speed of hydrazine decomposition. The program cannot handle system instabilities.

The difficulty of application of the UARL model to real transient situations is illustrated in Figures E-la and E-lb taken from References E-2 and E-3, respectively. In Figure E-la attention is directed to the experimental curve of chamber pressure. Note (1) the time delay from opening of the propellant valve to the rise in pressure and (2) the shape of the pressure pulse. Compare this curve to the theoretical pressure pulse calculated via the UARL code as shown in Figure E-lb. The program does not properly simulate the experimental data for this case.

E. 2 ANALYSIS

a. Scope

The monopropellant subprogram at its current state must be considered quasi-steady-state. It is necessary to include a substantial subprogram to compute steady state conditions as inputs to the time dependent 'main' program. This program uses numerical integration techniques centered around the catalyst bed reaction kinetics and associated mass and heat fluxes.

The initial flow of propellant is calculated beginning with the valve opening. The propellant is assumed to accelerate through inertial, resistive feed lines. The monopropellant is fed into the chamber through orifices which are direct extensions of the feed tubes. In the absence of experimental data, the injected streams are assumed to impact into the catalyst bed, with stream breakup parameters falling between a splash plate and a shower head model.

The injection velocity is such that in an unpacked chamber a droplet of liquid propellant would traverse a 2-inch-long chamber in the order of 1 millisecond.

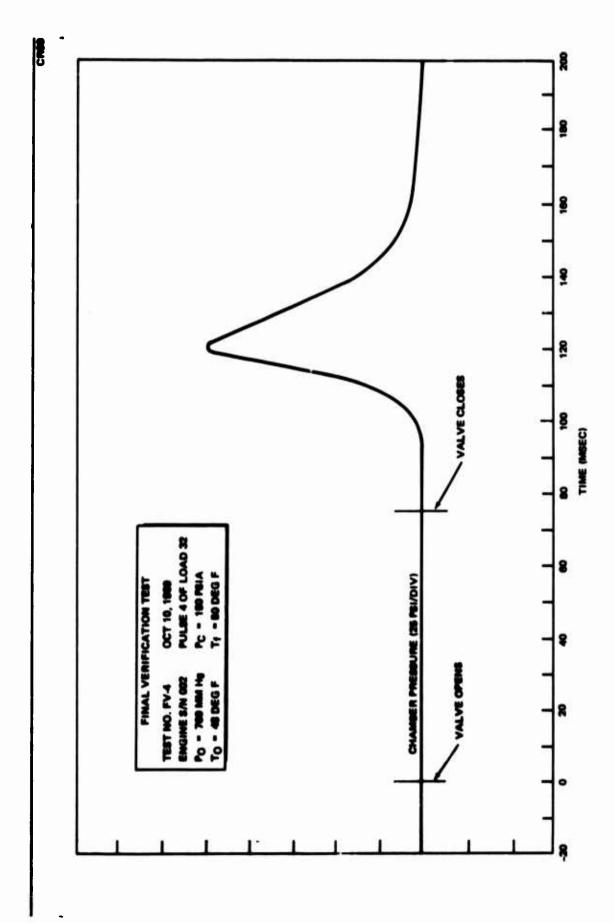


Figure E-1a. Final Verification Test

CREE

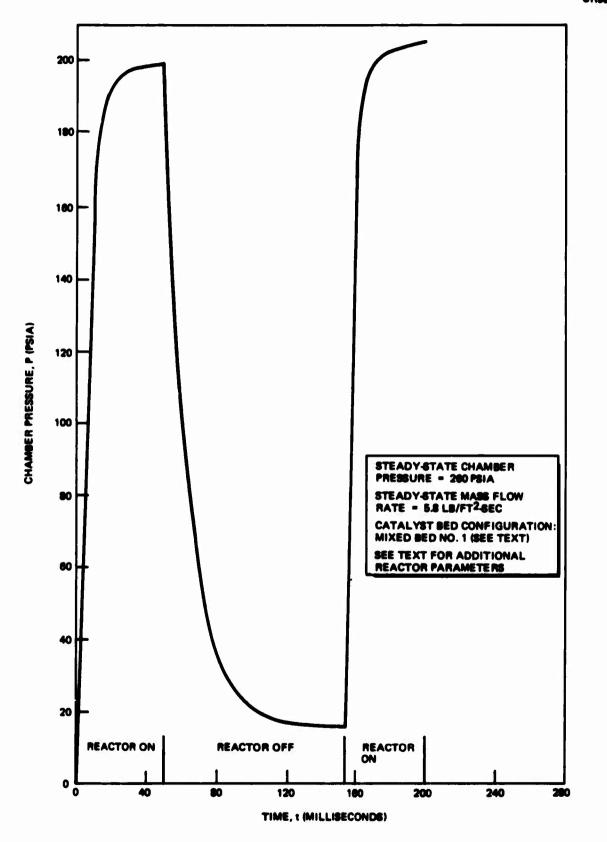


Figure E-1b. Variation of Chamber Pressure With Time

In the estimation of startups constrained by evaporation, assumptions most favorable to rapid evaporation were used, and evaporation limitations were used as the test for cold starts. The processes of distribution and evaporation of liquid hydrazine are considered crucial to a model capable of accurately predicting dynamic behavior in the cold start of a catalyst-pack hydrazine motor.

The hot vapor catalyst bed reactions used included both heterogenous and homogenous decomposition of hydrazine. Consideration was also given to the decomposition of ammonia and the heat transfer between catalyst particles and the surrounding gases.

The heat flux to the wall and atmosphere was treated in a lumped manner, with the chamber wall being considered an isothermal heat sink.

The mass flow of gas was calculated as a function of time and axial distance with the changes in composition and temperature based upon the catalyst bed processes controlling rates. There is no gas acceleration treatment in either the chamber or the throat, nor calculation of the pressure drop across the catalyst bed.

Gas composition profiles are calculated for each interval of model time. The gas composition at the throat section gives values for undecomposed hydrazine and ammonia expelled as contaminant. An infinite evaporation rate of hydrazine was assumed and this produced a tendency to underestimate the residual hydrazine and ammonia. In the case of cold starts, the MDAC modifications yield estimates of the liquid contamination attributable to the initial pressure spike.

The crucial limitations in the model are (1) lack of liquid phase dispersion and breakup, (2) lack of competition between wetted catalyst reactions and vapor-catalyst reactions, (3) lack of detailed evaporation processes, and (4) the chamber gas dynamics are treated as slug-flow i.e., gas velocity and composition differences only in the radial dimension. These deficiencies grossly 'imit the ability of this program to model start transients (particularly cold starts), pulse transients, or low frequency instabilities.

There is no capability in this program to treat situations in which the initial ambient pressure is less than the propellant vapor pressure. Low-temperature stops have been added by MDAC to avoid meaningless computer output in these regimes.

Performance estimation requires that the downstream end thermochemical data be manually combined with assumed motor parameters

b. Modeling the System Processes

1. Feed System

The feed systems are approximated with single lumped parameters representing the inertial and resistive aspects of the feed system, i.e.,

$$\frac{dq}{dt} = \frac{P_t - P_c - 1/2 \rho Rq|q|}{\rho \Sigma L/A}$$
 (E-1)

where

q = volumetric flow rate through line

P₄ = tank pressure

P = chamber pressure

R = a coefficient representing steady flow pressure drop in the line

ΣL/A = the summation of the segment length divided by crosssectional area for the series of segments which make up the feed system.

A common design style of monopropellant feed system consists of a cluster of long small bore tubes which directly feed the injector orifices. The feed valves are treated as simple "off-on" devices, and at the present level of sophistication of the program it is felt that no further accuracy would be gained by use of a valve ramp function. The resistive aspects of the feed system was determined from experimental flow rates versus pressure drops for test engines. The current feed system will compute the line fill time for any degree of line void which is input to the program. The feed will follow the time history of the chamber and compute the flow rate as a function of time and chamber pressure, i.e.,

$$q = \phi(t, Pc). \qquad (E-2)$$

This inertial/resistive feed system is the basis for computing inputs in the INLETM subroutine. It provides means of estimating the accumulation of propellants in the chamber in a cold start situation.

c. Physical Properties

In those segments of the program developed by UARL, physical properties of propellants and products were introduced in tabular form. The sections added at MDAC have approximated the physical properties of hydrazine and the associated chemical products as algebraic functions of temperature, using the relations developed by Priem (Reference E-4).

d. Propellant Stream Distribution and Breakup

The injection velocity of the hydrazine feed is computed. At present, there is no propellant spray distribution system in this model. However, propellant evaporation rates are used for testing the accumulation of propellants on cold start. The assumption that droplet sizes lie between shower head and splash ring values leads to the approximation:

$$D = 0.25 D_0,$$
 (E-3)

where

D = median droplet diameter, and

D₀ = orifice diameter of the injector

e. Monopropellant Decomposition

The monopropellant decomposition packed bed model was developed by Kesten (Reference E-1). The calculation is time and distance dependent but as stated earlier is based solely on the vapor phase condition and the mass, enthalpy, and species balances are treated in slug flow. The rates of mass and heat flux between the interstitial gases and the particle surfaces are treated as transport-process-controlled for the hydrazine catalysis. The much slower catalytic decomposition of ammonia is treated as controlled by chemical kinetics.

The UARL model assumes that the hydrazine vaporizes so rapidly that this process can be assumed to occur in zero axial distance. Subsequently, hydrazine decomposes exothermically either heterogenously at the catalyst surface or homogeneously in the gas phase. The ammonia breaks up endothermically on the catalyst surfaces.

f. Testing for Cold Start Spikes

In the MDAC extension of the program, start spikes are assumed to be due to the accumulation of liquid hydrazine in the chamber, and are estimated by calculating the amount of evaporation of liquid hydrazine occurring in the time needed for it to pass through the fine particle stage of the catalyst bed. Arbitrarily it has been assumed that sufficient hydrazine must evaporate to yield the heat required to vaporize 50 percent of the propellant fed to the chamber in order to avoid a hard start. In practice, about 1/6 of the propellant must evaporate in the transit time through the catalyst fine zone to avoid triggering the cold start test calculation. The drop velocity is assumed to be equal to injection velocity.

The "flight" distance or "effective" bed path length is assumed to be twice the physical thickness of the fine particle zone, due to the tortuous path.

The evaporation of drops with forced convective heating has been investigated by Cramer (Reference E-5) and Webber (Reference E-6), and the mass rate of change is given by

$$\dot{M} = \left(\frac{\pi K}{cp}\right) D \left[2.0 + 0.5N_{R_e}^{0.5}\right] ln \left[1 + \left(\frac{cp}{\Delta H}\right) \Delta T\right]$$
 (E-4)

where

M = mass rate of change with time,

k = thermal conductivity of the boundary layer about the drop,

c = specific heat at constant pressure,

D = drop diameter,

NR = Reynolds number,

 ΔH = heat of vaporization,

ΔT = difference between medium and droplet surface temperature.

The evaporation rates existing in a hydrazine catalyst bed at or near steady-state conditions will be slow compared to bipropellant engines. Under cold start conditions, evaporation rates are orders of magnitude slower. In the test for cold start flooding, the thermal conductivity, K, of the boundary layer around the hydrazine drop is taken to be that of air. This assumption is justified because normally cold starts occur when the temperature is less than 560°R, and in this case the hydrazine vapor never exceeds 5 percent of the mole fraction of the hydrazine-air mixture.

At the present time the program is limited to (1) a single hard-start transient with a cold catalyst bed or (2) hot start calculations assuming instantaneous vaporization of hydrazine. The temperature value of 560°R is an experimentally determined figure below which the engine will normally experience a hard start. Should the model later be expanded to cover multiple pulses or transient calculations leading to the equilibrium state, the following information should be noted:

When the temperature of the hydrazine is of the order of the boiling point, the correct value of K would be that given by Priem (Reference E-1) derived from the properties of hydrazine vapor. As the engine continues to heat, i.e., when the catalyst temperature reaches 1,700 to 1,900°R, this value will be too low, and the effect of decomposition products must be considered. The thermal decomposition products NH_3 , N_2 , and H_2 have a substantially higher value of K than N_2H_4 alone.

g. Simulating Cold Start Transients and the Associated Contamination

When the cold start test routine senses a condition indicating the accumulation of liquid hydrazine in the catalyst bed i.e., the hydrazine temperature is less than 560°R, the program computes the accumulation of liquid hydrazine by integrating the propellant feed rate from propellant entrance time to pulse cut off or pulse spike time, whichever comes first. Referring above to the feed system model, the volumetric feed rate, q, at time, t, is given by integrating Equation (E-1).

The accumulation of propellants in the cold chamber in preparation for a chug and spike is obtained by the summation

$$M_{\text{puddle}} = \rho \int_{1}^{t_2} q dt$$
 (E-5)

where:

M_{puddle} = accumulating mass of liquid N₂H₄,

t = time of first arrival of propellant to chamber after filling feed lines,

t₂ = time to propellant flow cutoff (either due to pulse cutoff or chamber pressure spike whichever is first).

The times to spiking and the peak pressure were fitted to empirical data gathered on a 5-lb and a 100-lb engine reported by UARL (Reference E-3). These data were collected over a temperature range from 495° to 560°R. The time to pressure rise was curve fitted as a function of temperature, and the peak pressure was fitted as a function of time to pressure rise.

The contaminants discharged from the chamber from a cold start surge are estimated by calculating the fraction of the puddle mass (M_{puddle}) which must react to generate enough gas and temperature to produce the pressure spike, assuming the reaction,

$$3N_2H_4 \longrightarrow 4NH_3 + 3N_2 + \Delta H$$

Gaseous ammonia will be blown out at the cold start spike. It is assumed that the upstream catalyst particles will have had the longest time to heat up. Much of the pressure surge will, therefore, be upstream of the bulk of the accumulating puddle mass, and consequently, an estimated one half of the residual unreacted puddle mass will be blown out as liquid N₂H₄. That portion of the residual puddle mass which is not blown out of the chamber with the initial chug is assumed to evaporate and react (on a longer time scale) to produce a slower discharge of gaseous NH₃ and N₂. Experimental monitoring of liquid hydrazine spray under these conditions has been reported at a recent Plume Interference Conference (Reference E-7).

If the chamber or propellants are warm enough to pass the cold start test routine, the calculation will proceed immediately to the UARL time-dependent hydrazine reactor program. This program has now been coupled with inertial/resistive feed system so that it more realistically reflects feed fluid dynamics. The program is, however, still without an integrated liquid-phase distribution and evaporation scheme.

h. Chamber Calculations

The UARL time-dependent program generates a time-axial distance calculating matrix for particle temperature, interstitial gas temperature, interstitial composition, and mass flow rates. An average chamber pressure is calculated for each time point. Gas velocity effects in the chamber are not estimated and the contraction region or throat are not treated.

E. 3 CATALYST BED BREAKUP LITERATURE SEARCH

The effective-use life of the catalyst bed is the most frequent limiting factor for service life of a hydrazine monopropellant reactor. The bulk of

the literature on catalyst bed breakup does not mention or consider catalyst breakup debris in the rocket exhaust or as a plume effect. The recent conference on plume effects (Reference E-7) contains the only specific mention of catalyst particles in the rocket exhaust. The rest of the literature in this area deals only with loss of effectiveness of the catalyst bed.

a. Types of Catalyst Losses

A survey of the current literature indicates that catalyst loss is due to mechanical abrasion, systems vibration, and spalling due to thermal cycling. References E-8, E-9, E-10, and E-11 suggest a source of these phenomena are low-frequency instabilities; Reference E-15 quotes system support vibration; and references E-13 and E-15 also trace sources of the loss phenomena to high-velocity stream erosion and thermal shock.

The losses are correlated with the particular system design and mode of operation. References E-9 through E-12 and E-16 through E-18 cite various losses of the catalyst particle masses ranging from a low of 0, 1 percent mass loss/min to values as high as 50 percent of the total mass during unstable runs.

References E-13 and E-19 indicate that there can be substantial function loss or drop in performance for monopropellant engines. The figures are put in the range of 50 to 67 percent over several minutes of run time.

The accumulation of damage from mechanical and thermal sources associated with transient phenomena appears far more important than the total run time. There is insufficient information presently available to weight particular conditions on an a priori basis.

The current state of the literature on catalyst breakup is relatively sparse in detailed information on catalyst particle breakup mechanics and loss exhaust rates. Consequently, the best approach at this time would be to develop a system by system statistical model built upon the available empirical data.

E. 4 MODEL DEVELOPMENT

Table E-1 illustrates the various phenomena that were modeled in the original UARL program; those which have been added by MDAC; and those which are recommended for future inclusion.

E. 5 CELL LOADER STRUCTURE

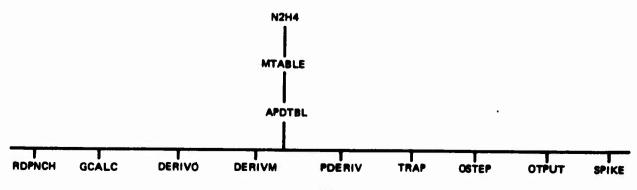


Table E-1

PROCESSES NEEDED FOR A TRANSIENT
MONOPROPELLANT CONTAMINATION MODEL

Process	UARL	Added Current	Recommended Addition
Feed System			
Inertial/Resistive Flow	No	Yes	Yes
Dribble Start	No		Yes
Dribble Cutoff	No		Yes
Injection			
Stream Breakup Atomization	No	Yes	Yes
Low-Pressure Flashing	No		Yes
Chamber Gas Dynamics	No	No	Yes
Drop Mechanics			
Breakup	No		Yes
Acceleration	No		Yes
Evaporation	No	Yes	
Drop Size Distribution	No		Yes
Vacuum Start	No		Yes
Wall Effects			
Heat Transfer	Yes		Real
	(isothermal)		Transients
Wetting			
Liquid Drag Breakoff	No		Yes
Evaporation	No		Yes
Decomposition of N ₂ H ₄			
Liquid N2H4	Yes		
Vapor Heterogenous	Yes		
Homogeneous	Yes		
Decomposition of NH ₃			
Vapor Heterogenous	Yes		
Homogenous	Yes		
Heat Exchange - Gas/Catalysts	Yes		
Calculation of Cold Start			
Conditions	No	Yes	
Contaminant	No	Yes	
Run Contaminants	No		Yes
Transient Operations	Nol		Yes
Performance Calculations	No		Yes
Affects of Chamber			
Dynamics on Catalyst Bed	No	Yes^2	Yes
Dimensional Effects	No		Yes

The program is based on steady state inputs and will not calculate a "true" transient operation.

²Only most primitive calculations have currently been made.

E. 6 PROGRAM SUBROUTINES

a. N2H4

This is the main routine and contains all MDAC modifications. It controls program decision making for both continuous and pulse operation cases; it also calculates all times, temperatures, concentrations, pressures, etc., using values generated in the various subroutines. Input/output is controlled by this routine.

b. INLETM

This routine calculates the inlet conditions of the bed at the start of each time increment.

c. GCALC

This routine calculates the mass flow rate used in particle and interstitial equations.

d. DERIVM

This routine calculates certain functions used in the particle temperature and concentration equations.

e. PDERIV

This routine calculates partial derivatives of interstitial temperature and concentration with respect to axial position.

f. DERIVO

This routine is not used in the present version of N2H4, but is retained for possible future use with multiple pulse applications.

g. OSTEP

This routine interpolates to obtain temperatures, concentrations, etc., when the accumulated time has "overstepped" the duty cycle interrupt time. The routine is used only for pulse operation.

h. SPIKE

This routine is not used in the present version of N2H4, but is retained for possible future use with multiple-pulse applications.

i. TRAP

This routine is used to integrate certain functions using the trapezoidal rule.

j. RDPNCH

This routine is not used in the present version of N2H4, but is retained for possible future use with multiple-pulse applications.

k. OTPUT

This routine retrieves and prints information stored on drum during the course of program calculations.

1. UNBAR

This is an interpolation routine used to obtain values from a table.

m. Block Data

Many of the arrays and parameters contained in the labeled common's are input via block data. In particular these data cover tables of:

- 1. Temperature vs Viscosity
- 2. Utilization Factor (log)
- 3. Temperature vs N2H4 Heat of Reaction
- 4. Temperature vs NH3 Heat of Reaction
- 5. Temperature vs N2H4 Specific Heat
- 6. Temperature vs NH₃ Specific Heat
- 7. Temperature vs N2 Specific Heat
- 8. Temperature vs H2 Specific Heat
- 9. Vapor Pressure vs Temperature

E. 7 PROGRAM USER'S MANUAL

a. Inputs

Input to N2H4 is carried out via the NAMELIST mode. However, it should be mentioned that certain of the original control parameters of the UARL code have been arbitrarily set within the MDAC program. This was done to make the former compatible with the modifications made in the latter. Certain of the parameter values are obtained from the steady state program, STDST, explained further on. The NAMELIST's and their corresponding parameter lists are as follows:

1. NAMELIST/ENGINE

Parameter	Identity
zo	Axial distance to injector end (ft)
GOSS	Mass flow rate, obtained from STDST (lb/ft ²)
MBSS	Steady state average molecular weight obtained from STDST
PSS	Pressure chamber, steady state obtained from STDST (Psia)
TSS	Steady state downstream temperature obtained from STDST (*R)
TVAP	Vaporization temperature obtained from STDST (*R)

HIV Enthalpy at liquid-vapor/vapor interface (Btu/lbm) Distributed feed rate (lbm/ft3-sec) F HF Enthalpy of the feed (Btu/lbm) TF not used in present calculation ENMX1, ENMX2. ENMX3 not used in current calculation PF pressure of propellant feed (psia) ZEND not used in present calculation VC volume of chamber (free volume) AC Cross-sectional area, of reactor (ft²) MW chamber wall thermal mass (lbm) CW Chamber wall specific heat (Btu/lbm-°R) ΑW Chamber wall total surface area (ft²) TA Ambient temperature in chamber (°R) Forced convection heat transfer coefficient (Btu/ft²-sec-*R) HA HA1 Natural convection heat transfer coefficient (Btu/ft2-sec-·R1.25) HA2 Radiative heat transfer coefficient (Btu/ft²-sec R⁴) TPI Initial catalyst temperature (°R) PCI Initial chamber pressure (psia) Initial hydrazine concentration in the catalyst (lbm/ft³) CPHYI **CPAMI** Initial ammonia concentration in catalyst (lbm/ft³) CIAMI Initial interstitial ammonia concentration (lbm/ft³) Initial nitrogen concentration in the catalyst (lbm/ft³) CPN2I Initial interstitial nitrogen concentration (lbm/ft³) CIN2I Initial hydrogen concentration in catalyst (lbm/ft3) CPH2I CIH2I Initial interstitial hydrogen concentration (lbm/ft³) AG Acceleration due to gravity (ft/sec2) DORIF Diameter of orifice (ft) Number of orifices (No. /ft²) ORINO **FDLINL** Feedline length (ft) FULFR Fraction of feed line full at start TLIQ Temperature of liquid hdyrazine (°R) **TPULS** Time of pulse signal (sec) FINZON Thickness of fine particle bed (ft) CIHYI Initial Interstitial hydrazine concentration (lbm/ft³)

2. NAMELIST/CATALYST

Parameter	Identity
M	Time mesh dimension
N	Axial distance mesh dimension
NOFZ	Number of axial stations used in input tables
Z(75)	Axial stations in total solution mesh
Z1(37)	Axial stations used in table inputs
RADT(37)	Catalyst particle radii (ft)
SURFT(37)	Total external catalyst particle surface areas/unit Volume of catalyst bed (ft ² /ft ³)
VOIDT(37)	Interparticle void fractions

3. NAMELIST/CONTROL

Parameter	Identity	
DRUM MPRNT	Fortran logical unit designator Print indicator (default value = MPRNT = 1 gives much more	0 gives summary printout,
NPASS DELT(20)	total number of passes through Time increments between each	mesh defined by M and N

b. Steady-State Data

The program STDST supplies the needed steady state input values required by the Monopropellant Engine program N2H4. STDST has not been incorporated into CONTAM (i.e., driven by the EXEC program) because there is only one set of equilibrium conditions for a given engine design, and once these values are determined, any transient problem may be solved using these steady state conditions.

STDST, except for the conversion to NAMELIST input, is the UARL Steady State one-dimensional program (Reference E-20), and the reader is directed to the reference for the details of the program.

It should be noted that this program does not lend itself to overlaying in that many subroutines continually call upon other subroutines during a run. Consequently, STDST has been supplied in a stand-alone non-cell loader program, and requires a memory capacity of 150,000 words on a CDC 6500 computer.

The input to the program is given via the NAMELIST/STEADY/. The variables in this list are as follows:

NCASE	Number of data cases with each run, use NCASE = 1
NOFZ	Number of axial stations used in the table inputs
ZO	Axial distance to the end of buried injector (ft)
GO	Inlet mass flow rate (lbm/ft ² -sec)
FC	Rate of feed of hydrazine from buried injectors (lbm/ft ³ -sec)
HF	Enthalpy of liquid hydrazine entering catalyst bed (btu/lbm) set to zero for cold flow
TF	Temperature of liquid hydrazine entering bed (°R)
ENMX1	Constant used to determine the size of axial station incre- ments in the liquid region. It equals 200. Increasing this number would result in a decrease in size of axial station increments (and an increase in computer run time).
ENMX2	Constant used to determine the size of axial station increments in the liquid-vapor region. It equals 40. Increasing this number would result in a decrease in size of axial station increments (and an increase in computer run time).
ENMX3	Constant used to determine the size of axial station incre- ments in the vapor region. It equals 80. Increasing this number would result in a decrease in size of axial station increments (and an increase in computer run time).

PRES Inlet chamber pressure (psia)
ZEND Catalytic bed length (ft)

ZPOS Number of independent variables in tables

RADII Catalyst particle radius (ft)

SURFAR Catalyst particle surface area per unit of volume of bed

 (ft^2/ft^3)

VOIDFR Interparticle void fraction

E.8 CONTAM INTERFACING

The monopropellant engine model is not nearly to the same level of development as that of the bipropellant. Consequently, in determining the inputs for the MULTRAN portion of CONTAM certain assumptions and approximations must be made. The following list is a suggested NAMELIST input for the MULTRAN portion of CONTAM when the MDAC modifications are put to use in the program N2H4.

\$DATA CAPN= 0.6, CPG=18, 150, CPL=S.A.B., CPS=S.A.B., GAMMA=

1.206, GMGO=4.24E-5, HPL=S.A.B., HPS= S.A.B., PR=0.7, RCAP=

3102 *Same as bipropellant

E. 9 SAMPLE CASES

The sample cases shown are those used by Kesten, et al (References E-1 and E-20). The steady state program, STDST, is illustrated first, and then the program N2H4. Two examples of N2H4 are shown; a "hot" catalyst bed run, the original Kesten program; and the "cold" bed or the program with the MDAC modifications.

a. Steady State Program

The following list identifies the required STDST outputs and their corresponding name for the N2H4 program input.

N2H4 Input STDST Output

GOSS GO in the INPUT CONSTANTS

MBSS MBAR printed out at end of output

PSS last PRES printed out TSS last TEMP printed out.

b. Monopropellant Engine, Hot catalyst Bed

The term "hot catalyst bed" refers to the cases when the parameters TF (feed temperature), TA (ambient temperature), TPI (bed particle temperature), and TLIQ (liquid fuel temperature) are all greater than 561°R.

The output has been edited and during a run considerably more printout will be obtained. With the hot bed there are no droplets produced. If the user wishes to make a contamination run for the hydrazine, ammonia,

nitrogen, and hydrogen vapors, the last value on each of the interstitial lists should be used. In the case shown these values appear just before the "completion of pass 1" message.

The figures given are in units of lbm/ft³. The droplet sizes should be set to vanishingly small, and the rest of CONTAM will calculate for the vapor phase only.

c. Monopropellant Engine, Cold Catalyst Bed

When temperature parameters listed above fall below 560°R, the Mono-propellant Engine program will use the MDAC modifications. In this case the output is much simplified. The time required to reach the pressure peak of the start transient is shown, the pressure obtained, the amounts of liquid hydrazine and ammonia produced, and the amount of ammonia vapor produced.

At the present time there is some evidence (Reference E-7) to suggest that very small droplets are formed during a cold start transient. Since the Monopropellant program has no capability to calculate droplet sizes, it is recommended that the diameter of 20 microns be assumed for the droplets.

E. 10 REFERENCES

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