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A COMPREHENSIVE CYCLE ANALYSIS AND DIGITAL
COMPUTER SIMULATION FOR SPARK-IGNITED ENGINES

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TABLE OF CONTENTS

	<u>Page</u>
ACKNOWLEDGMENTS.....	ii
LIST OF TABLES.....	v
LIST OF FIGURES.....	vi
LIST OF APPENDICES.....	viii
NOMENCLATURE.....	x
I INTRODUCTION.....	1
A. Purpose.....	1
B. Previous Work.....	1
C. Scope of This Paper.....	10
II THE THERMODYNAMIC PROPERTIES AND COMPOSITION OF THE WORKING FLUID IN A SPARK-IGNITED ENGINE.....	11
A. General.....	11
B. Calculation of Ideal Gas Thermodynamic Properties.....	13
1. Empirical Heat Capacity Equations.....	13
2. Computation of Thermodynamic Functions and Equilibrium Constants from Empirical Heat Capacity Equations.....	15
C. The Composition of the Products of Combustion.....	24
1. Assumption of Chemical Equilibrium.....	24
2. The Chemical Composition at Equilibrium.....	31
a. General.....	31
b. Computation of Homogeneous Gas Equilibrium....	33
c. Computation of Frozen Equilibrium	43
3. Computer Solution of the Equations of Chemical Equilibrium.....	46
III SIMULATION OF THE FUEL-AIR ENGINE CYCLE ON A DIGITAL COMPUTER.....	47
A. General.....	47
B. Results and Discussion.....	50
IV COMBUSTION IN A SPARK-IGNITED ENGINE.....	53
A. General.....	53
B. Flame Propagation.....	54
C. Thermal Theory of Flame Propagation for A Bimolecular Second-Order Reaction.....	59

TABLE OF CONTENTS (CONT'D)

	<u>Page</u>
D. Use of the Thermal Theory Equations to Predict Turbulent Flame Propagation Velocity.....	66
V HEAT TRANSFER IN A SPARK-IGNITED ENGINE.....	69
A. General.....	69
B. Convective Heat Transfer.....	72
1. Temperature Distribution and Heat Flow in a Finite Plate in Contact with a Uniform Medium whose Temperature Varies Periodically.....	72
2. The Convective Heat Transfer Coefficient.....	80
C. Radiant Heat Transfer.....	82
VI THE ANALYTICAL MODEL FOR SPARK-IGNITED ENGINE CYCLE CALCULATIONS.....	85
A. General.....	85
B. The Cycle Processes.....	86
C. The Cycle Characteristics.....	102
D. Computational Method.....	103
VII CALCULATED RESULTS FROM THE ANALYTICAL MODEL.....	104
A. General.....	104
B. The "Normal Cycle".....	105
C. Comparison of the Analytical and Experimental "Normal Cycles".....	106
D. Effect of Exhaust Temperature.....	111
E. Influence of Heat Transfer and Flame Propagation Rates.....	112
F. Availability and Irreversibility.....	115
G. The Analytical Model and the Fuel-Air Cycle.....	117
H. Concluding Remarks.....	118
APPENDICES.....	136
BIBLIOGRAPHY.....	283

LIST OF TABLES

<u>Table</u>		<u>Page</u>
I	Ideal Gas Empirical Heat Capacity Equations.....	16
II	Fundamental Thermodynamic Properties.....	17
III	Calculated and Tabular Values of Ideal Gas Heat Capacity, C_p°	18
IV	Calculated and Tabular Values of Ideal Gas Enthalpy, $H^{\circ} - H_0^{\circ}$	20
V	Calculated and Tabular Values of Ideal Gas Entropy, S°	21
VI	Calculated Values of \log_{10} of the Equilibrium Constant, K_p	23
VII	Pertinent Dimensions for Engine of Figure 11.....	121
VIII	Volume-Area-Length Relationships for the Combustion Chamber of Figure 11.....	122
IX	Normal Engine Cycle Operating Conditions.....	123
X	Amount of Heat Transferred.....	124
XI	Digit Changed in the Partial Pressure of Each Constituent for Selected Values of Relative Error.....	158

LIST OF FIGURES

<u>Figure</u>		<u>Page</u>
1	The Otto Cycle.....	3
2a	Mole Fraction Vs Temperature, O ₂	38
2b	Mole Fraction Vs Temperature, N ₂	38
2c	Mole Fraction Vs Temperature, CO ₂	39
2d	Mole Fraction Vs Temperature, H ₂ O.....	39
2e	Mole Fraction Vs Temperature, CO.....	40
2f	Mole Fraction Vs Temperature, H ₂	40
2g	Mole Fraction Vs Temperature, OH.....	41
2h	Mole Fraction Vs Temperature, NO.....	41
2i	Mole Fraction Vs Temperature, O.....	42
2j	Mole Fraction Vs Temperature, H.....	42
3	Indicator Diagram for the Fuel-Air Cycle Engine at Part Throttle.....	48
4	Efficiency and Mean Effective Pressure Vs Compression Ratio for Lean Mixtures.....	51
5	Efficiency and Mean Effective Pressure Vs Compression for a Mixture of Octene and 110 Percent Theoretical Air.....	52
6	Wrinkled Flame Front Model of Scurlock and Grover....	57
7	Schematic Diagram and Temperature Profile in the Region of the Reaction Zone.....	60
8	Heat Transfer Model for the Internal Combustion Engine.....	74
9	Indicator Diagram for the Engine of the "Analytical Model" at Part Throttle.....	87
10	Mathematical Model for Combustion and Heat Transfer..	96-97

LIST OF FIGURES (CONT'D)

<u>Figure</u>		<u>Page</u>
11	View of Engine Combustion Chamber Showing Spherical Flame Propagation Pattern.....	120
12	Cylinder Pressure Vs Crank Angle for a 10:1 Compression Ratio Engine With Normal Rate of Flame Propagation.....	125
13	Pressure-Volume Diagram for a 10:1 Compression Ratio Engine With Normal Rate of Flame Propagation..	126
14	Pressure-Volume Diagram for a 10:1 Compression Ratio Engine With Rapid Rate of Flame Propagation..	127
15	Pressure-Volume Diagram for a 10:1 Compression Ratio Engine With Reduced Rate of Flame Propagation	128
16	Cylinder Pressure Vs Crank Angle for a 10:1 Compression Ratio Engine With Normal Heat Transfer	129
17	Calculated Thermal Efficiency and Mean Effective Pressure Vs Flame Propagation Rate for a 10:1 Compression Ratio Engine.....	130
18	Entropy-Volume Diagram for a 10:1 Compression Ratio Engine with Normal Rate of Flame Propagation.....	131
19	Availability Vs Crank Angle for a 10:1 Compression Ratio Engine With Normal Rate of Flame Propagation and Heat Transfer.....	132
20	Availability Vs Crank Angle for a 10:1 Compression Ratio Engine With No Heat Transfer and Normal Flame Propagation Rate.....	133
21	Flame Speed and Effective Flame Temperature Vs Crank Angle for a 10:1 Compression Ratio Engine With Normal Flame Propagation and Heat Transfer Rates.....	134
22	Relationship Between Mass and Volume of Burned Charge for a 10:1 Compression Ratio Engine With Normal Rate of Flame Propagation.....	135
23	Flow Diagram for the Subroutine Computing the Composition of the Products of Combustion.....	159

LIST OF APPENDICES

<u>Appendix</u>	<u>Page</u>
A.	POLYNOMIAL APPROXIMATION BY THE METHOD OF LEAST SQUARES..... 136
B.	EQUILIBRIUM CONSTANT AS A FUNCTION OF TEMPERATURE. 140
C.	CALCULATION OF THERMODYNAMIC PROPERTIES FROM EMPIRICAL HEAT CAPACITY EQUATIONS..... 144
A.	General..... 144
B.	Method of Calculation..... 146
1.	Computation of the Enthalpy..... 146
2.	Computation of the Entropy..... 147
3.	Computation of $(\partial H/\partial T)_P$ 149
4.	Computation of $(\partial S/\partial T)_P$ 149
5.	Computation of the Equilibrium Constant... 150
C.	Computer Programs..... 151
D.	THE TIME RATE OF TEMPERATURE CHANGE DURING EXPANSION..... 152
E.	METHOD OF SOLUTION OF THE EQUATIONS OF SIMULTANEOUS GAS EQUILIBRIUM..... 154
F.	MATHEMATICAL ANALYSIS, FLOW DIAGRAM, AND COMPUTER PROGRAM FOR THE FUEL-AIR CYCLE..... 160
A.	General..... 160
B.	Mathematical Analysis..... 161
1.	Exhaust Residual - Initial Approximation... 161
2.	Mixing of the Fresh Charge and Exhaust Residual..... 162
3.	Isentropic Compression..... 163
4.	Constant Volume Adiabatic Combustion..... 164
5.	Isentropic Expansion..... 165
6.	Expansion to Exhaust Pressure..... 165
7.	Exhaust Residual..... 166
8.	Net Work..... 166
9.	Thermal Efficiency..... 166
10.	Mean Effective Pressure..... 167
C.	Flow Diagram..... 168
D.	Computer Program..... 172

LIST OF APPENDICES (CONT'D)

<u>Appendix</u>		<u>Page</u>
G.	ITERATIVE METHODS FOR SOLUTION OF IMPLICIT FUNCTIONS OF A SINGLE VARIABLE.....	178
	A. General.....	178
	B. Methods of Solution.....	178
	1. Newton's Method.....	178
	2. Half-Interval Method.....	179
H.	CALCULATION OF THE RATE OF RADIANT HEAT TRANSFER....	180
I.	FLOW DIAGRAM AND PROGRAM LISTING FOR THE ANALYTICAL MODEL.....	184
	A. Flow Diagram.....	184
	B. Explanation of Symbols Used in Flow Diagram and Computer Program.....	198
J.	SPECIAL THERMODYNAMIC CONSIDERATION ASSOCIATED WITH INTER-RELATED STATE CHANGES OF TWO DISTINCT GASEOUS REGIONS.....	231
	A. Isentropic Change of Volume of Each Region.....	231
	B. Change of Volume With Known Values of Internal Energy and Entropy.....	232
K.	CALCULATED DATA.....	235

NOMENCLATURE

A	Area
ATC	After top center
a	Atoms of carbon per molecule of fuel
a	Activity
BDC	Bottom center piston position
BTC	Before top center
Bore	Engine bore
b	Atoms of hydrogen per molecule of fuel
C_p	Constant pressure heat capacity per mole
C_v	Constant volume heat capacity per mole
c	Mole ratio of nitrogen to oxygen in atmospheric air
c_p	Constant pressure heat capacity per unit mass
E	Activation energy
F	Exhaust residual mass fraction
F	A constant, $F = RT/V$
F_0	Moles of fuel inducted per engine power cycle
f	Fugacity
G	Gibbs free energy
G	Mass rate of flow per unit area
H	Enthalpy
h	Convective heat transfer coefficient
J	Conversion factor from heat to work units
K_c	Equilibrium constant based on concentrations
K_p	Equilibrium constant based on partial pressures

NOMENCLATURE (CONT'D)

K_L	Flame speed factor (Equation (4-24))
k	Heat capacity ratio
k	Thermal conductivity
L	Hydraulic radius
M	Molecular weight
mep	Indicated mean effective pressure
n	Polytropic exponent
n_i	Number of moles of component 'i'
P	Pressure
P_i	Partial pressure of component 'i'
Q	Amount of heat transferred
q	Rate of heat transfer
R	Universal gas constant
Rod	Engine connecting rod length
r	Compression ratio
rpm	Engine speed
S	Entropy
S_ℓ	Laminar flame speed
S_t	Turbulent flame speed
Stroke	Piston stroke
T	Temperature
TDC	Top center piston position
t	Time
U	Internal energy

NOMENCLATURE (CONT'D)

V	Volume
V_{bot}	Clearance volume plus swept volume
V_{top}	Clearance volume
v	Specific volume
W	Work
w	Average piston velocity
X	Square root of partial pressure of O_2
x	Air-fuel ratio, fraction chemically correct air
Y	Square root of partial pressure of H_2
Z	Square root of partial pressure of N_2
α	Thermal diffusivity, $\alpha = \rho c_p/k$
ΔH_c	Enthalpy of combustion
ΔH_f	Enthalpy of formation
η	Thermal efficiency
η_v	Volumetric efficiency
θ	Crank angle degrees, $0^\circ = \text{BDC piston position}$
θ	$\theta = T/T_0$
μ	Viscosity
ρ	Density
ϕ	Fuel-air ratio, fraction chemically correct fuel, $\phi = 1/x$

SUPERSCRIPTS

'	Designates a trial value
o	Ideal gas state

NOMENCLATURE (CONT'D)

SUBSCRIPTS

b	Burned
e	Exhaust manifold
f	Flame element
f	Fuel
i	Inlet manifold
P	Constant pressure
p	Piston
t	Total
u	Unburned
w	Wall
0	Reference state
298	298.16

I. INTRODUCTION

A. Purpose

The purpose of this investigation was to study the four stroke spark ignited internal combustion engine cycle from a fundamental point of view. The effects of mixture composition, combustion chamber geometry, flame propagation, and heat transfer were incorporated into an analytical model, suitable for programming on a digital computer, which approximates more closely than heretofore possible the actual performance of an engine. This work is primarily concerned with the development of such a model and the subsequent evaluation of the individual effects of the heat transfer and flame propagation phenomena on the thermal efficiency and indicated mean effective pressure of the hypothetical engine. Experimentally determined performance data are compared with values calculated using the analytical model.

B. Previous Work

In 1876 Nicolas Otto constructed the first successful gas engine which employed compression prior to the combustion of the charge. This innovation, although suggested earlier by Beau de Rochas in 1862, produced an engine which was more efficient than any heretofore and marked the beginning of the modern four stroke cycle spark-ignited engine^(1, 2). Although it was the introduction of the compression process which accounted for the improved efficiency, Otto himself did not seem to realize this and attributed the improvement in efficiency to a stratification of the charge. Only when thermodynamic analyses of the various gas engine cycles of the day were made were the relative merits of Otto's engine understood.

The earliest attempts to analytically study engines largely followed the fundamental work of Sadi Carnot. As in any mathematical simulation of a real physical process, an analytical model was developed which incorporated the salient features of the physical process which it approximated. In the case of Otto's engine, the processes which the working fluid underwent could be closely approximated by the constant volume cycle, now commonly called the Otto cycle. The Otto cycle is composed of four highly idealized processes (see Figure 1).

Process 1-2. Compression of the working fluid from volume V_1 to V_2 . This occurs reversibly and adiabatically, therefore, isentropically.

Process 2-3. Constant volume heat addition. An amount of heat Q_1 is added to the working fluid thereby raising its temperature to T_3 and its pressure to P_3 . This occurs when the piston is at the top center position.

Process 3-4. Expansion of the working fluid from V_2 to V_1 . This expansion occurs isentropically.

Process 4-1. Constant volume heat rejection. An amount of heat Q_2 is transferred from the working fluid, lowering its temperature to T_1 and its pressure to P_1 , thus completing the cycle.

The isentropic compression and expansion follow the path $PV^k = a$ constant, where 'k' is the ratio of heat capacity of the working fluid at constant pressure to that at constant volume. The work of the cycle is $Q_1 - Q_2$ and its thermal efficiency is

$$\eta = \frac{Q_1 - Q_2}{Q_1} \quad (1-1)$$

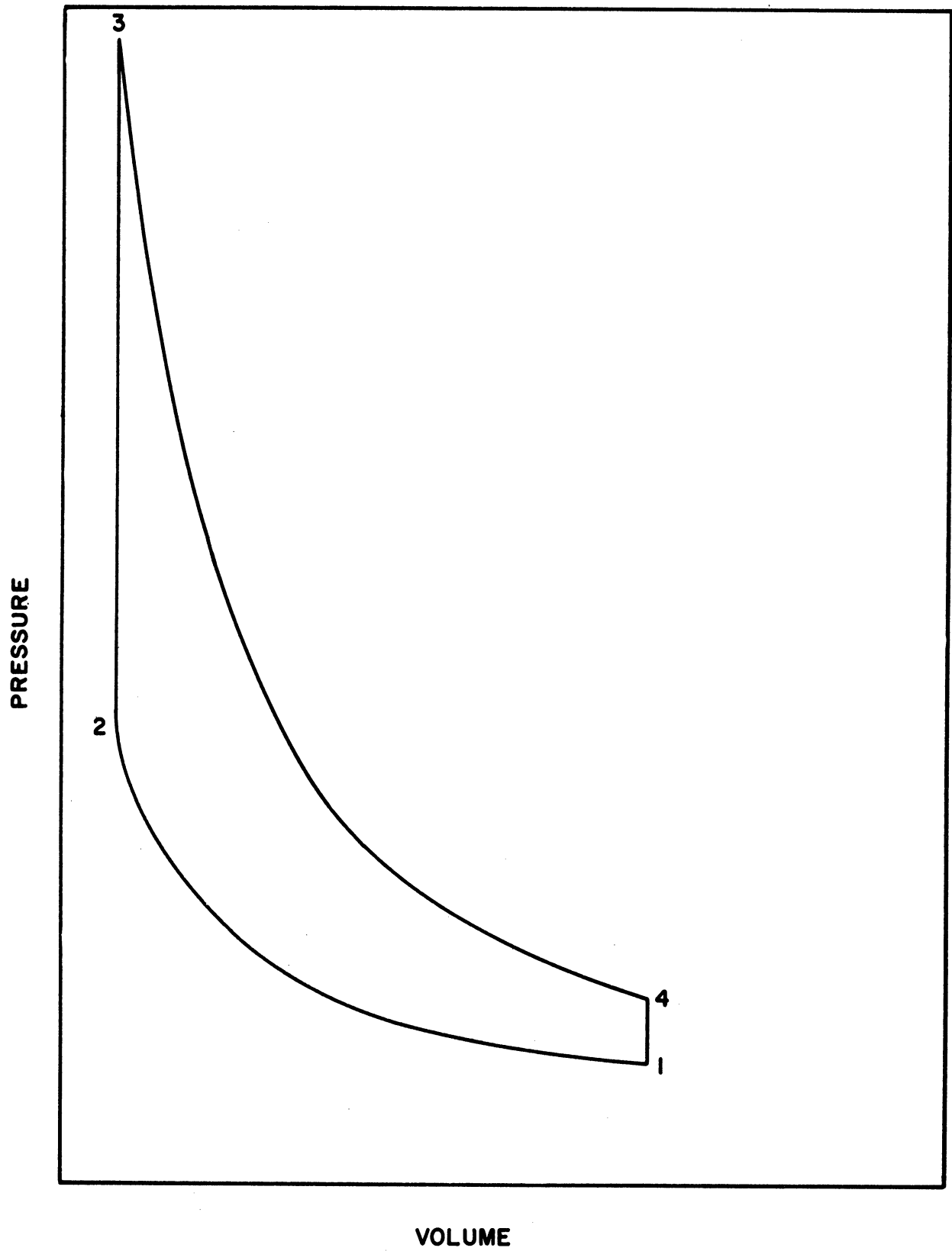


Figure 1. The Otto cycle.

For an Otto cycle engine of compression ratio 'r' this can be expressed as

$$\eta = 1 - \left(\frac{1}{r}\right)^k \quad (1-2)$$

The mean effective pressure, defined as the quotient of the cycle work and the displacement is

$$mep = \frac{\eta \cdot Q_1}{V_1 - V_2} \cdot J \quad (1-3)$$

Thus, as a consequence of such an analysis, convenient mathematical expressions were developed for the efficiency and mean effective pressure of an engine operating on this idealized Otto cycle.

Because at this time knowledge about the properties of the working fluid was meager, it was decided to adopt as an analytical model for the Otto cycle, the constant volume cycle operating on an idealized fluid having the properties of air at standard conditions. In 1905 a committee of the Institution of Civil Engineers⁽³⁾ recommended that the air cycle efficiency be officially adopted as the standard of comparison against which measured values from internal combustion engines might be compared. This decision was based on the result of tests which consistently showed that at a given compression ratio measured efficiencies were from 50% to 70% of air cycle efficiencies. This report left little doubt concerning the value of compression in the internal combustion engine.

Shortly thereafter, Clerk⁽⁴⁾, using heat capacities deduced by experimental methods⁽⁵⁾, showed that in considering the actual heat capacity of the working fluid, the attainable thermal efficiencies were considerably lower than those predicted by the air cycle analysis. In 1908, Hopkinson⁽⁶⁾ analyzed the Otto cycle engine using the heat capacity

of the mixture calculated from the known values for the individual constituents. By this, he was able to quantitatively predict the influence of the fuel-air ratio on the thermal efficiency and mean effective pressure and thus confirm experimental evidence that leaner mixtures tended to give higher thermal efficiencies. His analysis took account of the exhaust residual.

Not until 1921, was an analysis of the constant volume cycle carried out which considered the actual composition of the working fluid including the effects of dissociation. At this time this became possible because accurate heat capacity data and sufficient information on the extent of dissociation of carbon, hydrogen, oxygen, and nitrogen systems at elevated temperatures were available. This analysis, by Tizard and Pye⁽⁷⁾, assumed that during compression the working fluid was primarily composed of air, but that after combustion and during expansion, the working fluid consisted of O_2 , N_2 , CO_2 , H_2O , CO , and H_2 in a state of chemical equilibrium. Because of the complexity of the analysis, no analytical expressions for thermal efficiency or mean effective pressure could be derived.

Whereas the effect of compression ratio on the performance of the Otto cycle engine was shown by the air cycle analysis, that of mixture composition was shown by Clerk's analysis, and that of fuel-air ratio was shown by Hopkinson's analysis, the effects of dissociation were introduced by Tizard and Pye. In doing so they answered three pressing problems of the day. First, because of dissociation effects, the highest combustion temperatures and mean effective pressures occur with mixtures 10 to 15 percent richer than stoichiometric. Second, for increasingly rich mixtures the mean effective pressure, which is a measure of the power output, is rela-

tively constant and only decreases slowly as the fuel-air ratio is increased. Third, the thermal efficiency of the engine increases as the mixture is leaned out. This last effect is due to the fact that both heat capacity and dissociation increase with an increase in temperature.

Whereas the best measured thermal efficiencies are as low as 50 to 70 percent of the air cycle efficiency, they are 80 to 85 percent of those calculated by Tizard and Pye's method. Many people had been openly critical of the validity of the air cycle analysis, but now it was realized that at a given compression ratio, the efficiency calculated from the air cycle represented an optimum value which could be approached in practice only with very lean mixtures. This study was the first of many which analyzed the Otto cycle engine in this manner.

In 1924, Goodenough and Felbeck⁽⁸⁾ presented a comprehensive discussion of the theory of chemical equilibrium and the effects of dissociation on the maximum temperatures and pressures attainable in combustion processes. Using the results of this analysis Goodenough and Baker⁽⁹⁾, in 1927, presented a complete thermodynamic analysis of the Otto cycle together with the results of numerous calculations. Their analysis followed along the same lines as Tizard and Pye's, but in addition included the effects of the mixing of the exhaust residual with the fresh charge.

In 1935, Hershey, Eberhardt, and Hottel⁽¹⁰⁾ presented thermodynamic charts from which the properties of the working fluid in the Otto cycle engine could be read. These eliminated the tedious calculations of Goodenough and Baker. The charts, based on spectroscopic heat capacity data, allowed for complex dissociation at high temperatures. The species included O_2 , N_2 , CO_2 , H_2O , CO , H_2 , OH , NO , O , and H in the products of combustion.

The charts have been widely used from the time of their introduction for the analysis of the Otto cycle engine and have been revised a number of times to allow for improvements in fundamental data (see bibliography of Reference 11). While the charts allow a considerable reduction in the mathematical difficulties of the Otto cycle analysis, they are not general, having been developed for a particular fuel and a selected number of fuel-air ratios. The analysis of the Otto cycle proposed by Hershey, Eberhardt, and Hottel was very similar to that proposed by Goodenough and Baker but considered the induction and exhaust processes in detail.

The Otto cycle is termed the fuel-air cycle when the actual properties of the working fluid are considered just as it was termed the air cycle when air was considered to be the working fluid. The fuel-air cycle is described in more detail in Chapter III.

In spite of the refinements of the fuel-air cycle analysis, measured thermal efficiencies were still between 80 and 90 percent of those calculated. This is because many important phenomena present in actual engines are disregarded in the simplicity of the Otto cycle engine. Two significant items neglected are the flame propagation and the heat transfer. However incorporating these effects into an analytical model poses extremely complex mathematical problems. Nevertheless some investigators have attempted to include one or another of these phenomena in their analysis.

Attempts to consider the effects of flame propagation (progressive burning of the charge), were first made in connection with thermodynamic studies of combustion in constant volume bombs. Among the earliest was an analysis by Nagel⁽¹²⁾ in 1907. A similar analysis was made by

Rosecrans⁽¹³⁾ in 1926 which made use of the equilibrium studies of Goodenough and Felbeck. Endres⁽¹⁴⁾ was among the first to consider the effects of piston motion. Hershey, Eberhardt and Hottel⁽¹⁰⁾ suggested a method by which the temperature gradients in a constant volume bomb could be calculated considering mixture composition, variable specific heats, and dissociation. In 1937, Hottel and Eberhardt⁽¹⁵⁾ extended the work of Reference 10 and proposed a method for calculating the temperature gradients in an engine considering both piston motion and spark advance. Pressure volume diagrams were calculated for the combustion portion of the cycle assuming no heat was transferred to the cylinder walls.

Rassweiler and Withrow⁽¹⁶⁾ and Rassweiler, Withrow, and Cornelius⁽¹⁷⁾ suggested a method for treating the combined effects of flame propagation and piston motion in internal combustion engines in order to analyze combustion photographs. In 1940 Withrow and Cornelius⁽¹⁸⁾ proposed a method for calculating the pressure rise in an internal combustion engine. They combined the analysis of the previous two references with the thermodynamic charts of Reference 10. Rabezanna, Kalmar, and Candelise⁽¹⁹⁾ in 1939, proposed a method for the analysis of flame propagation and expansion during combustion which took into account combustion chamber geometry, spark advance, and piston motion. Although this analysis considered the thermodynamics of the working fluid in an uncomplicated manner, pressure-time diagrams could be calculated from measured flame propagation rates by their method.

These analyses of flame propagation in internal combustion engines were hindered in two respects. First, fundamental information about the nature of the flame propagation in an internal combustion engine

was not available. Second, the sheer magnitude of the calculations involved, assuming that flame propagation information were available, would have precluded any extensive analysis. All of these studies assumed that the engine operated under adiabatic conditions.

In 1940, David and Leah⁽²⁰⁾ presented charts of "attainable thermal efficiency" and fuel consumption for Otto cycle engines. These charts were computed from an analysis similar to that of Hershey, Eberhardt, and Hottel, but included a prescribed amount of heat transfer during the constant volume combustion and subsequent expansion.

A concise presentation of References 10 and 15 was published in 1948⁽¹¹⁾, the thermodynamic charts having been revised in light of more accurate thermodynamic data. However no new technique for internal combustion engine analysis was introduced. In 1950, Leah⁽²¹⁾ illustrated a technique for approximating the thermal efficiencies the Otto cycle engine for any hydrocarbon fuel from charts. Edson⁽²²⁾ in 1961 presented the results of calculations on the fuel-air cycle model made on a digital computer. These covered a wide range of variables. A portion of the work done in the investigation being reported involves an analysis similar to that made by Edson and is reported in Chapter III.

The problem of considering the combined effects of mixture composition, variable specific heats, dissociation, flame propagation, and heat transfer simultaneously presented such a formidable mathematical obstacle that not until the widespread use of automatic computing machines was such an analysis contemplated. In 1960, Edson⁽²³⁾ proposed an analytical model for combustion in the internal combustion engine which was suitable for programming on a digital computer. His analysis included the effects of

flame propagation and piston motion but did not include heat transfer. To date no computed results have been reported.

C. Scope of This Paper

The remainder of this paper describes the development and evaluation of a comprehensive cycle analysis and digital computer simulation for spark-ignited internal combustion engines. Chapter II describes a method which is used for the determination of the thermodynamic properties and composition of the working fluid. Chapter III presents the results of computations using a well known internal combustion engine cycle analysis but employing the methods of Chapter II to describe the working fluid.

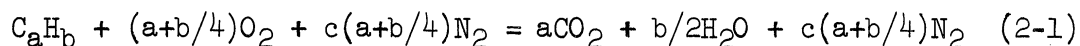
Chapters IV and V discuss the nature of the flame propagation and heat transfer in the spark-ignited engine. Chapter VI describes an analytical model for engine cycle calculation which includes both heat transfer and flame propagation effects. Chapter VII presents results of computer calculation made using this analytical model. These are compared with experimental engine data.

II. THE THERMODYNAMIC PROPERTIES AND COMPOSITION OF THE WORKING FLUID IN A SPARK-IGNITED ENGINE

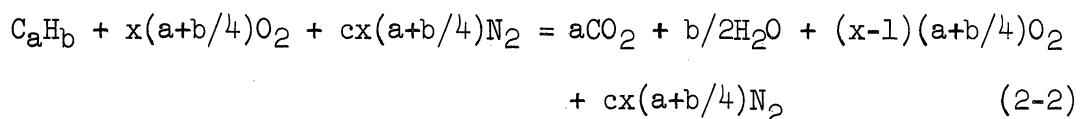
A. General

This chapter describes a method by which the thermodynamic properties and composition of the working fluid of an internal combustion engine can be calculated. The method presented is ultimately employed to describe the working fluid of the "analytical model" engine proposed in Chapter VI. It is also employed in the fuel-air engine cycle analysis discussed in Chapter III.

For the reaction between a hydrocarbon fuel and oxygen and nitrogen, the following equation can be written when stoichiometric proportions of oxygen and nitrogen, subsequently referred to as air, are present:



where 'c' is the mole ratio of nitrogen to oxygen in air and 'a' and 'b' are the number of carbon and hydrogen atoms in a molecule of fuel. When an excess of air is present, the following equation can be written:



where 'x' is the fraction of chemically correct air entering the reaction. When 'x' is less than one, that is for rich mixtures, no oxygen is present in the products but some CO and H₂ will be. These equations tend to oversimplify the combustion process and many more chemical species will be present in the products due to dissociation.

While it is true that atmospheric air contains substances other than oxygen and nitrogen, only water vapor exists to any appreciable extent. In order to establish a reference base for all the calculations, dry air is assumed. Dry air at sea level has the following composition⁽²⁴⁾:

Basis: 100 moles dry air

O ₂	20.99	
N ₂	78.03	
A	0.94	
CO ₂	0.03	
H ₂	0.01	<u>79.01</u>
		100.00

For the purposes of this work, the argon, carbon dioxide, and hydrogen normally found in air are lumped together with the nitrogen and are assumed to have the same thermodynamic properties as nitrogen. Thus air is assumed to be composed of 79.01 parts of nitrogen by volume to 20.99 of oxygen, their ratio 'c' being 3.764 to 1. The error in the value of the thermodynamic properties calculated on this basis is about 0.4% or less.

Furthermore, all substances including the fuel are treated as ideal gases and mixtures are assumed to follow Dalton's law of partial pressures. The error in the thermodynamic properties introduced by this assumption varies as it depends on the temperature and pressure. Because of the nature of the Otto cycle, the highest pressures always coincide with the highest temperatures and thus the deviation from ideal gas behavior tends to be small. Such deviations as may exist are greatest during the induction and compression processes. Reference 11 presents charts for

corrections to ideal gas thermal functions calculated by the Beattie-Bridgman equation of state for octene-air mixtures. For such a hydrocarbon-air mixture which is compressed isentropically from room temperature and pressure, the ideal gas thermodynamic properties will be no more than about 0.5 percent greater than those of the real gas for the states encountered along this path.

B. Calculation of Ideal Gas Thermodynamic Properties

1. Empirical Heat Capacity Equations

In a carbon, hydrogen, oxygen, nitrogen system at high temperatures, dissociation produces many chemical species whose individual thermodynamic properties must be known in order to compute the gross properties of the working fluid which they comprise. These thermodynamic properties are normally presented in tabular form or may be calculated from equations approximating the tabular data. Since the analytical model for the internal combustion engine presented in this study is designed to be simulated mathematically on a digital computer, it was decided to calculate the thermodynamic properties from ideal gas empirical heat capacity equations. This obviates the use of a large amount of computer storage capacity for the tables.

Hougen, Watson, and Ragatz have tabulated a number of heat capacity equations in the appendix of their recent text⁽²⁵⁾ which includes most of the equations developed by Kobe and his associates⁽²⁶⁾. Either a single equation was used for a rather narrow temperature range (such as 273-1500°K) giving fairly accurate results, but not covering the higher temperatures, or a single equation for a larger temperature range (such as 273-3800°K) with a significant decrease in accuracy. Spencer⁽²⁷⁾ presents

heat capacity equations calculated by the method of least squares for a number of hydrocarbons. Williams⁽²⁸⁾ presents heat capacity equations for a large number of substances. Two sets of constants are given for each substance, one set for the temperature range 500-3000°K, the other for the range 3000-6000°K. In these references the constants for the empirical heat capacity equations are presented as five or six digit numbers.

In order to achieve greater accuracy than the above equations afford in the desired temperature range, it was decided to develop highly accurate empirical heat capacity equations for the required substances. Two equations were developed for each compound, the first covering the range 298.16-1500°K and the second the range 1500-3500°K. Moreover, since a high speed computing machine capable of handling eight digit numbers was to be employed, the constants were calculated and used as eight digit numbers. The basic data used to develop the equations came from tables of ideal gas heat capacity at one atmosphere as a function of temperature. For the substances employed here, two sources were used, the American Petroleum Institute Report 44⁽²⁹⁾ and the National Bureau of Standards Circular 564⁽³⁰⁾. Using these data, empirical equations for the heat capacity as a function of temperature were developed by the method of least squares, the details of which are given in Appendix A. The equations for each substance are of the form:

$$C_p = a_0 + a_1T + a_2T^2 + a_3T^3 \quad (2-3)$$

298.16-1500°K

$$C_p = b_0 + b_1T + b_2T^2 + b_3T^3 \quad (2-4)$$

1500-3500°K

Table I gives the data source, the a's and b's, the maximum percent deviation, and the root mean square deviation for those substances which comprise the working fluid of an internal combustion engine. For these substances Table II lists the enthalpies of formation and absolute entropies at 298.16°K, the enthalpies and entropies at 1500°K, and the enthalpy and free energy of reaction at 1500°K, for selected reactions (all quantities at 1 atmosphere pressure). Table III gives selected values of heat capacity as a function of temperature, listing both the tabular and calculated value at each point. The heat capacity representation in the range 1500-3500°K proved to be very accurate. The maximum deviation for any of the substances considered was less than 0.1 percent of the tabular value. In this range, the use of such approximating equations will represent the tabular function within the stated uncertainty of the table. In the range from 298.16-1500°K, where the heat capacity function changes rapidly with temperature, the accuracy is less. The maximum percentage deviation is about 0.6 percent of the tabular value for some substances, less than 0.3 percent for most.

2. Computation of Thermodynamic Functions and Equilibrium Constants From Empirical Heat Capacity Equations

The computation of the ideal gas thermodynamic functions and equilibrium constants from the empirical heat capacity equations developed in the previous section is most conveniently made by arranging the constants for each substance (the a's and b's) and the necessary fundamental data in matrix form. The thermodynamic functions then can be calculated, as required, by a series of matrix multiplications. This procedure is outlined in Appendix C.

TABLE I
IDEAL GAS EMPIRICAL HEAT CAPACITY EQUATIONS

$$C_p^0 = a_0 + a_1T + a_2T^2 + a_3T^3 \text{ cal/mole-}^\circ\text{K}$$

298.16-1500 °K

Substance	Data Source	*a0	a1	a2	a3	Max. % Dev.	RMS Dev.
O ₂	NBS-564	0.61907849E+01	0.29234242E-02	-0.70522172E-06	-0.80285969E-10	0.64	0.022
N ₂	API-44	0.70507626E+01	-0.11515000E-02	0.30557188E-05	-0.11525971E-08	0.35	0.015
CO ₂	NBS-564	0.51113583E+01	0.15430851E-01	-0.99580134E-05	0.24073171E-08	0.36	0.020
H ₂ O	NBS-564	0.77613959E+01	0.19662306E-04	0.31476535E-05	-0.10812777E-08	0.22	0.0092
CO	API-44	0.69101229E+01	-0.56810040E-03	0.26467764E-05	-0.10721376E-08	0.47	0.020
H ₂	NBS-564	0.68529104E+01	0.16183303E-03	0.61791131E-07	0.14864846E-09	0.53	0.018
OH	API-44	0.76220243E+01	-0.24470010E-02	0.29835421E-05	-0.82731476E-09	0.30	0.0063
NO	API-44	0.70200658E+01	-0.36766843E-03	0.25451986E-05	-0.10864604E-09	0.68	0.029
O	API-44	0.55969158E+01	-0.16383585E-02	0.14788035E-05	-0.44240119E-09	0.31	0.0080
H	API-44	0.49680000E+01	0.00000000E+00	0.00000000E+00	0.00000000E+00	0.0	0.0
n-Heptane	API-44	-0.11854254E+01	0.16126768E+00	-0.86976233E-04	0.18208351E-07	0.30	0.084
n-Octane	API-44	-0.14126484E+01	0.18397817E+00	-0.99985303E-04	0.21072621E-07	0.30	0.092
1-Octene	API-44	-0.59674715E+00	0.17107204E+00	-0.94379942E-04	0.20222447E-07	0.21	0.086

$$C_p^0 = b_0 + b_1T + b_2T^2 + b_3T^3 \text{ cal/mole-}^\circ\text{K}$$

1500-3500 °K

Substance	Data Source	*b0	b1	b2	b3	Max. % Dev.	RMS Dev.
O ₂	NBS-564	0.78906694E-01	0.51781745E-03	0.53216719E-07	-0.13820401E-10	0.028	0.0009
N ₂	API-44	0.64837196E+01	0.19298323E-02	-0.54849168E-06	0.56397533E-10	0.056	0.0021
CO ₂	NBS-564	0.10728501E+02	0.33912981E-02	-0.97734787E-06	0.10230199E-09	0.070	0.0044
H ₂ O	NBS-564	0.53120816E+01	0.58710071E-02	-0.14930128E-05	0.14120276E-09	0.026	0.0019
CO	API-44	0.67392679E+01	0.17599157E-02	-0.50255810E-06	0.51864546E-10	0.052	0.0021
H ₂	NBS-564	0.52684432E+01	0.22852214E-02	-0.50854453E-06	0.48668136E-10	0.050	0.0019
OH	API-44	0.54319835E+01	0.23640016E-02	-0.57035617E-06	0.53582743E-10	0.018	0.0009
NO	API-44	0.71280200E+01	0.15047494E-02	-0.43395095E-06	0.46032279E-10	0.048	0.0019
O	API-44	0.50290719E+01	-0.34659544E-04	-0.42474464E-08	0.43342867E-11	0.001	0.0002
H	API-44	0.49680000E+01	0.00000000E+00	0.00000000E+00	0.00000000E+00	0.0	0.0

*Note O.xxx E+xx = (0.xxx ...) · (10^{+xx})

TABLE II

*FUNDAMENTAL THERMODYNAMIC PROPERTIES

Substance	ΔH_{f298}° cal/mole	S_{298}° cal/mole-°K	$(H_{1500}^{\circ} - H_{298}^{\circ})$ $+\Delta H_{f298}^{\circ}$ cal/mole	S_{1500}° cal/mole-°K	ΔH_{c298}° kcal/mole
O ₂	0	49.014 ^b	9705.3 ^b	61.6639 ^b	--
N ₂	0	45.767 ^a	9181.3 ^a	57.786 ^a	--
CO ₂	-94051.8 ^a	51.070 ^b	-79303.6 ^b	69.812 ^b	--
H ₂ O	-57797.9 ^a	45.106 ^b	-46303.6 ^b	59.8567 ^b	--
CO	-26415.7 ^a	47.301 ^a	-17130.3 ^a	59.436 ^a	--
H ₂	0	31.208 ^b	8673.6 ^b	42.7206 ^b	--
OH	10060.0 ^a	43.888 ^a	18863.3 ^a	55.568 ^a	--
NO	21600.0 ^a	50.339 ^a	31099.8 ^a	62.760 ^a	--
O	59159.0 ^a	38.469 ^a	65204.9 ^a	46.641 ^a	--
H	52089.0 ^a	27.327 ^a	58059.7 ^a	35.419 ^a	--
n-Heptane	44890.0 ^a	102.24 ^a	--	--	1075.85 ^a
n-Octane	49820.0 ^a	111.55 ^a	--	--	1222.77 ^a
1-Octene	19820.0 ^a	110.55 ^a	--	--	1194.97 ^a

Reaction	ΔH_{1500}° cal/mole	$(\Delta G^{\circ}/RT)_{1500}$
CO ₂ = CO + $\frac{1}{2}$ O ₂	67026.0	12.192
H ₂ O = H ₂ + $\frac{1}{2}$ O ₂	59830.0	13.180
NO = $\frac{1}{2}$ N ₂ + $\frac{1}{2}$ O ₂	-21656.0	-5.7379
OH = $\frac{1}{2}$ H ₂ + $\frac{1}{2}$ O ₂	-9673.8	-1.5465
O = $\frac{1}{2}$ O ₂	-60352.0	-12.292
H = $\frac{1}{2}$ H ₂	-53723.0	-10.948

* All quantities at 1 atmosphere pressure.

^a Reference 29.

^b Reference 30.

TABLE III

CALCULATED AND TABULAR VALUES OF IDEAL GAS HEAT CAPACITY, C_p^0

Temp ($^{\circ}$ K)	Units: cal/mole- $^{\circ}$ K										
	n-Octane	O ₂	N ₂	CO ₂	H ₂ O	CO	H ₂	OH	NO	O	H
298.16	45.11 <u>45.14</u>	6.998 -----	6.949 <u>6.960</u>	8.891 -----	8.018 -----	6.948 <u>6.965</u>	6.911 -----	7.136 <u>7.141</u>	7.108 <u>7.137</u>	5.228 <u>5.236</u>	4.968 <u>4.968</u>
300.00	45.35 <u>45.35</u>	7.002 <u>7.024</u>	6.949 <u>6.961</u>	8.909 <u>8.895</u>	8.021 <u>8.027</u>	6.949 <u>6.965</u>	6.911 <u>6.895</u>	7.134 <u>7.139</u>	7.109 <u>7.134</u>	5.227 <u>5.234</u>	4.968 <u>4.968</u>
500.00	68.21 <u>68.32</u>	7.466 <u>7.431</u>	7.095 <u>7.070</u>	10.638 <u>10.665</u>	8.423 <u>8.415</u>	7.154 <u>7.120</u>	6.968 <u>7.000</u>	7.041 <u>7.048</u>	7.337 <u>7.289</u>	5.092 <u>5.080</u>	4.968 <u>4.968</u>
700.00	85.61 <u>85.66</u>	7.864 <u>7.884</u>	7.347 <u>7.351</u>	11.859 <u>11.845</u>	8.947 <u>8.954</u>	7.442 <u>7.451</u>	7.047 <u>7.037</u>	7.087 <u>7.087</u>	7.637 <u>7.657</u>	5.023 <u>5.028</u>	4.968 <u>4.968</u>
900.00	98.54 <u>98.43</u>	8.192 <u>8.212</u>	7.649 <u>7.671</u>	12.688 <u>12.667</u>	9.540 <u>9.546</u>	7.761 <u>7.787</u>	7.157 <u>7.142</u>	7.233 <u>7.234</u>	7.959 <u>7.990</u>	4.998 <u>5.005</u>	4.968 <u>4.968</u>
1100.00	108.0 <u>108.0</u>	8.446 <u>8.439</u>	7.947 <u>7.947</u>	13.240 <u>13.243</u>	10.153 <u>10.151</u>	8.061 <u>8.058</u>	7.304 <u>7.305</u>	7.439 <u>7.440</u>	8.249 <u>8.243</u>	4.995 <u>4.994</u>	4.968 <u>4.968</u>
1300.00	115.1 <u>115.2</u>	8.623 <u>8.605</u>	8.186 <u>8.165</u>	13.631 <u>13.656</u>	10.731 <u>10.723</u>	8.289 <u>8.265</u>	7.494 <u>7.516</u>	7.665 <u>7.663</u>	8.457 <u>8.426</u>	4.994 <u>4.986</u>	4.968 <u>4.968</u>
1500.00	120.7 <u>120.6</u>	8.718 <u>8.739</u>	8.309 <u>8.330</u>	13.977 <u>13.952</u>	11.224 <u>11.233</u>	8.395 <u>8.419</u>	7.736 <u>7.720</u>	7.872 <u>7.875</u>	8.528 <u>8.560</u>	4.974 <u>4.982</u>	4.968 <u>4.968</u>
1500.00	-----	8.740 <u>8.739</u>	8.335 <u>8.330</u>	13.962 <u>13.952</u>	11.236 <u>11.233</u>	8.423 <u>8.419</u>	7.716 <u>7.720</u>	7.876 <u>7.875</u>	8.564 <u>8.560</u>	4.982 <u>4.982</u>	4.968 <u>4.968</u>
2000.00	-----	9.029 <u>9.029</u>	8.601 <u>8.602</u>	14.420 <u>14.423</u>	12.212 <u>12.213</u>	8.664 <u>8.665</u>	8.194 <u>8.195</u>	8.307 <u>8.308</u>	8.770 <u>8.771</u>	4.977 <u>4.977</u>	4.968 <u>4.968</u>
2500.00	-----	9.302 <u>9.301</u>	8.761 <u>8.759</u>	14.697 <u>14.691</u>	12.865 <u>12.862</u>	8.808 <u>8.806</u>	8.564 <u>8.565</u>	8.614 <u>8.614</u>	8.897 <u>8.895</u>	4.984 <u>4.983</u>	4.968 <u>4.968</u>
3000.00	-----	9.550 <u>9.551</u>	8.860 <u>8.861</u>	14.868 <u>14.872</u>	13.300 <u>13.303</u>	8.896 <u>8.898</u>	8.861 <u>8.859</u>	8.838 <u>8.838</u>	8.980 <u>8.981</u>	5.004 <u>5.004</u>	4.968 <u>4.968</u>
3500.00	-----	9.762 <u>9.762</u>	8.937 <u>8.934</u>	15.012 <u>15.005</u>	13.625 -----	8.966 <u>8.963</u>	9.124 <u>9.123</u>	9.016 <u>9.015</u>	9.052 <u>9.049</u>	5.042 <u>5.041</u>	4.968 <u>4.968</u>

Underlined entry is tabular value.

Tables IV and V list the calculated values of enthalpy and entropy alongside their corresponding tabular values as a function of temperature. The values of enthalpy are calculated from the following expression:

$$H_T^{\circ} - H_0^{\circ} = H_T^{\circ} - H_{T_0}^{\circ} + (H_{T_0}^{\circ} - H_0^{\circ})$$

where

$$H_T^{\circ} - H_{T_0}^{\circ} = \int_{T_0}^T C_p^{\circ} dT$$

The entropy is calculated from

$$S_T^{\circ} = S_T^{\circ} - S_{T_0}^{\circ} + (S_{T_0}^{\circ})$$

where

$$S_T^{\circ} - S_{T_0}^{\circ} = \int_{T_0}^T \frac{C_p^{\circ}}{T} dT$$

The quantities $(H_{T_0}^{\circ} - H_0^{\circ})$ and $(S_{T_0}^{\circ})$ are the tabular values at the reference temperature ' T_0 '. For the substances reported here they are from the data sources indicated in Table I and are doubly underlined in Tables IV and V. The low temperature range reference temperature is 298.16°K for the substances from API 44 and 300°K for those from NBS 564. The high temperature range reference temperature is 1500°K in every case. In general, the calculated values of enthalpy and entropy deviate less, percentagewise, from their tabulated values than do the calculated values of the heat capacity from tabular values. Such calculations for mixtures are always more accurate than the least accurate component.

TABLE IV
CALCULATED AND TABULAR VALUES OF IDEAL GAS ENTHALPY, $H^\circ - H^\circ_0$

Temp (°K)	n-Octane	Units: cal/mole										H
		O ₂	N ₂	CO ₂	H ₂ O	CO	H ₂	OH	NO	O	H	
298.16	8608.0	2074.8	2072.3	2238.3	2366.8	2072.6	2023.8	2106.2	2194.2	1607.4	1481.2	
	<u>8608.0</u>	-----	<u>2072.3</u>	-----	-----	<u>2072.6</u>	-----	<u>2106.2</u>	<u>2194.2</u>	<u>1607.4</u>	<u>1481.2</u>	
300.00	8691.0	2087.6	2085.1	2254.7	2381.5	2085.4	2036.5	2119.3	2207.3	1617.0	1490.3	
	<u>8691.0</u>	<u>2087.6</u>	<u>2085.1</u>	<u>2254.7</u>	<u>2381.5</u>	<u>2085.6</u>	<u>2036.5</u>	<u>2119.2</u>	<u>2206.8</u>	<u>1617.0</u>	<u>1490.4</u>	
500.00	20147.0	3535.5	3487.2	4218.9	4023.5	3493.8	3424.2	3534.2	3650.2	2647.6	2483.9	
	<u>20130.0</u>	<u>3528.9</u>	<u>3485.0</u>	<u>4223.6</u>	<u>4021.0</u>	<u>3490.0</u>	<u>3429.5</u>	<u>3535.0</u>	<u>3644.0</u>	<u>2645.4</u>	<u>2484.0</u>	
700.00	35612.0	5069.7	4930.1	6476.1	5758.8	4952.4	4825.3	4945.0	5146.9	3658.2	3477.5	
	<u>35581.0</u>	<u>5062.1</u>	<u>4925.3</u>	<u>6482.0</u>	<u>5756.7</u>	<u>4945.5</u>	<u>4831.8</u>	<u>4946.9</u>	<u>5136.6</u>	<u>3655.5</u>	<u>3477.6</u>	
900.00	54093.0	6676.5	6429.3	8936.4	7606.8	6472.6	6245.1	6375.7	6706.5	4659.7	4471.1	
	<u>54081.0</u>	<u>6673.8</u>	<u>6428.0</u>	<u>8938.2</u>	<u>7606.1</u>	<u>6471.0</u>	<u>6248.4</u>	<u>6377.4</u>	<u>6700.5</u>	<u>4658.7</u>	<u>4471.2</u>	
1100.00	74799.0	8244.6	7989.5	11532.9	9576.2	8055.6	7690.5	7842.3	8328.2	5658.8	5464.7	
	<u>74767.0</u>	<u>8340.3</u>	<u>7991.5</u>	<u>11532.2</u>	<u>9576.1</u>	<u>8056.4</u>	<u>7692.9</u>	<u>7844.1</u>	<u>8323.7</u>	<u>5658.4</u>	<u>5464.7</u>	
1300.00	97142.0	10049.9	9604.3	14221.8	11665.6	9692.2	9169.5	9352.8	10000.7	6657.9	6458.3	
	<u>97162.0</u>	<u>10045.4</u>	<u>9604.7</u>	<u>14224.6</u>	<u>11664.4</u>	<u>9690.2</u>	<u>9174.2</u>	<u>9354.8</u>	<u>9991.8</u>	<u>6656.4</u>	<u>6458.3</u>	
1500.00	120736.0	11785.4	11256.1	16982.4	13862.9	11363.0	10691.7	10907.2	11701.9	7655.2	7451.9	
	<u>120750.0</u>	<u>11780.0</u>	<u>11253.6</u>	<u>16986.5</u>	<u>13861.1</u>	<u>11358.0</u>	<u>10697.4</u>	<u>10909.5</u>	<u>11694.0</u>	<u>7653.3</u>	<u>7451.9</u>	
1500.00	-----	11780.0	11253.6	16986.5	13861.1	11358.0	10697.4	10909.5	11694.0	7653.3	7451.9	
	-----	<u>11780.0</u>	<u>11253.6</u>	<u>16986.5</u>	<u>13861.1</u>	<u>11358.0</u>	<u>10697.4</u>	<u>10909.5</u>	<u>11694.0</u>	<u>7653.3</u>	<u>7451.9</u>	
2000.00	-----	16222.7	15492.7	24091.1	19738.8	15635.0	14680.3	14961.0	16032.0	10142.8	9935.9	
	-----	<u>16223.2</u>	<u>15494.8</u>	<u>24093.1</u>	<u>19739.2</u>	<u>15636.0</u>	<u>14795.0</u>	<u>14960.0</u>	<u>16030.0</u>	<u>10142.8</u>	<u>9936.0</u>	
2500.00	-----	20806.2	19836.7	31376.3	26019.1	20006.0	18873.3	19196.0	20451.0	12632.6	12419.9	
	-----	<u>20806.4</u>	<u>19839.0</u>	<u>31377.2</u>	<u>26019.1</u>	<u>20008.0</u>	<u>18873.3</u>	<u>19193.0</u>	<u>20450.0</u>	<u>12632.5</u>	<u>12420.0</u>	
3000.00	-----	25520.4	24243.7	38770.4	32567.1	24434.0	23231.8	23562.0	24921.0	15128.8	14903.9	
	-----	<u>25520.8</u>	<u>24245.4</u>	<u>38770.4</u>	<u>32567.2</u>	<u>24435.0</u>	<u>23231.7</u>	<u>23559.0</u>	<u>24924.0</u>	<u>15128.7</u>	<u>14904.0</u>	
3500.00	-----	30350.1	28693.0	46240.0	39301.0	28899.0	27728.8	28026.0	29429.0	17639.3	17387.9	
	-----	<u>30350.8</u>	<u>28697.0</u>	<u>46241.3</u>	-----	<u>28900.0</u>	<u>27728.9</u>	<u>28025.0</u>	<u>29428.0</u>	<u>17639.3</u>	<u>17388.0</u>	

Underlined entry is tabular value.

TABLE V
 CALCULATED AND TABULAR VALUES OF IDEAL GAS ENTROPY, S° , AT 1 ATMOSPHERE PRESSURE

Units: cal./mole-°K

Temp (°K)	n-Octane	O ₂	N ₂	CO ₂	H ₂ O	CO	H ₂	OH	NO	O	H
298.16	<u>111.55</u> <u>111.55</u>	49.014 ----- <u>45.767</u>	45.767 <u>45.767</u>	51.070 ----- <u>47.301</u>	45.106 ----- <u>47.301</u>	47.301 <u>47.301</u>	31.208 ----- <u>43.888</u>	43.888 <u>43.888</u>	50.339 <u>50.339</u>	38.469 <u>38.469</u>	27.393 <u>27.393</u>
300.00	<u>111.83</u> <u>111.84</u>	49.057 <u>49.057</u>	45.810 <u>45.809</u>	51.125 <u>51.125</u>	45.150 <u>45.154</u>	47.344 <u>47.343</u>	31.251 <u>31.251</u>	43.932 <u>43.932</u>	50.383 <u>50.384</u>	38.501 <u>38.501</u>	27.423 <u>27.423</u>
500.00	140.59 <u>140.56</u>	52.745 <u>52.729</u>	49.388 <u>49.385</u>	56.104 <u>56.116</u>	49.340 <u>49.333</u>	50.937 <u>50.927</u>	34.793 <u>34.808</u>	47.548 <u>47.553</u>	54.063 <u>54.048</u>	41.136 <u>41.131</u>	29.961 <u>29.961</u>
700.00	166.45 <u>166.37</u>	55.322 <u>55.304</u>	51.813 <u>51.805</u>	59.890 <u>59.904</u>	52.255 <u>52.248</u>	53.388 <u>53.373</u>	37.150 <u>37.166</u>	49.921 <u>49.927</u>	56.578 <u>56.557</u>	42.837 <u>42.831</u>	31.633 <u>31.633</u>
900.00	189.60 <u>189.53</u>	57.340 <u>57.328</u>	53.695 <u>53.692</u>	62.977 <u>62.986</u>	54.574 <u>54.568</u>	55.296 <u>55.287</u>	38.933 <u>38.945</u>	51.718 <u>51.723</u>	58.537 <u>58.520</u>	44.096 <u>44.091</u>	32.881 <u>32.881</u>
1100.00	210.34 <u>210.28</u>	59.010 <u>58.999</u>	55.260 <u>55.259</u>	65.581 <u>65.587</u>	56.548 <u>56.543</u>	56.884 <u>56.878</u>	40.383 <u>40.394</u>	53.189 <u>53.194</u>	60.163 <u>60.150</u>	45.098 <u>45.094</u>	33.878 <u>33.878</u>
1300.00	228.99 <u>228.98</u>	60.436 <u>60.423</u>	56.608 <u>56.606</u>	67.826 <u>67.835</u>	58.292 <u>58.286</u>	58.250 <u>58.243</u>	41.618 <u>41.630</u>	54.450 <u>54.457</u>	61.559 <u>61.544</u>	45.933 <u>45.928</u>	34.708 <u>34.708</u>
1500.00	245.86 <u>245.84</u>	61.678 <u>61.664</u>	57.790 <u>57.786</u>	69.801 <u>69.812</u>	59.863 <u>59.857</u>	59.445 <u>59.436</u>	42.707 <u>42.720</u>	55.562 <u>55.568</u>	62.776 <u>62.760</u>	46.646 <u>46.641</u>	35.419 <u>35.419</u>
1500.00	----- -----	61.664 <u>61.664</u>	57.786 <u>57.786</u>	69.812 <u>69.812</u>	59.857 <u>59.857</u>	59.436 <u>59.436</u>	42.720 <u>42.720</u>	55.568 <u>55.568</u>	62.760 <u>62.760</u>	46.641 <u>46.641</u>	35.419 <u>35.419</u>
2000.00	----- -----	64.218 <u>64.218</u>	60.223 <u>60.225</u>	73.897 <u>73.896</u>	63.232 <u>63.232</u>	61.895 <u>61.896</u>	45.008 <u>45.008</u>	57.896 <u>57.894</u>	65.254 <u>65.252</u>	48.074 <u>48.074</u>	36.848 <u>36.848</u>
2500.00	----- -----	66.262 <u>66.263</u>	62.161 <u>62.164</u>	77.147 <u>77.147</u>	66.032 <u>66.032</u>	63.845 <u>63.845</u>	46.878 <u>46.878</u>	59.785 <u>59.781</u>	67.226 <u>67.225</u>	49.185 <u>49.185</u>	37.957 <u>37.957</u>
3000.00	----- -----	67.981 <u>67.981</u>	63.768 <u>63.769</u>	79.842 <u>79.841</u>	68.418 <u>68.419</u>	65.459 <u>65.459</u>	48.467 <u>48.466</u>	61.376 <u>61.374</u>	68.856 <u>68.857</u>	50.095 <u>50.095</u>	38.862 <u>38.863</u>
3500.00	----- -----	69.469 <u>69.470</u>	65.139 <u>65.140</u>	82.145 <u>82.144</u>	70.494 -----	66.836 <u>66.835</u>	49.852 <u>49.852</u>	62.752 <u>62.751</u>	70.245 <u>70.247</u>	50.869 <u>50.869</u>	39.628 <u>39.628</u>

Underlined entry is tabular value.

Table VI lists values of the equilibrium constants, calculated from Equation (C-5) (see Appendix C) for the principle dissociation reactions in the high temperature products of a hydrocarbon-air reaction. A derivation of the expression for the equilibrium constant as a function of temperature is presented in Appendix B. The calculated values of the equilibrium constants usually deviate less, percentagewise, from their tabulated values than does the maximum percentage deviation exhibited by the heat capacity function of any of the reaction constituents. However, since the calculation for the equilibrium constant involves differences of thermodynamic functions, the percent deviation tends to get larger as the absolute magnitude of the equilibrium constant diminishes. Because the heat capacity data were taken from several references no strictly comparable tabular values are available.

In Tables III, IV, and V, a double entry is made at 1500°K. This is the break point between the two temperature ranges. The first entry is calculated from the heat capacity equations for the lower range whereas the second is calculated from those for the higher range. Any discontinuity at 1500°K could be removed by adding appropriate constants to the fundamental data at the beginning of each temperature range. However, as no particular advantage could be seen in doing this, it was not done.

Appendix C also indicates the procedure used to calculate

$$\left(\frac{\partial H}{\partial T}\right)_P \quad \text{and} \quad \left(\frac{\partial S}{\partial T}\right)_P .$$

TABLE VI

CALCULATED VALUES OF \log_{10} OF THE EQUILIBRIUM CONSTANT, K_p

Temp ($^{\circ}$ K)	Reaction					
	$\text{CO}_2 = \text{CO} + 1/2\text{O}_2$	$\text{H}_2\text{O} = \text{H}_2 + 1/2\text{O}_2$	$\text{OH} = 1/2\text{O}_2 + 1/2\text{H}_2$	$\text{NO} = 1/2\text{N}_2 + 1/2\text{O}_2$	$\text{H} = 1/2\text{H}_2$	$\text{O} = 1/2\text{O}_2$
1500.00	-5.294	-5.721	0.6711	2.491	4.755	5.338
2000.00	-2.863	-3.535	0.3214	1.703	2.790	3.135
2500.00	-1.418	-2.218	0.1150	1.230	1.601	1.808
3000.00	-0.4642	-1.337	-0.0203	0.9155	0.8029	0.9205
3500.00	0.2112	-0.7061	-0.1145	0.6921	0.2302	0.2856

C. The Composition of the Products of Combustion

1. Assumption of Chemical Equilibrium

Having developed a scheme by which the thermodynamic properties of a system can be calculated when the chemical composition of that system is known, it is now necessary to investigate the nature of high temperature combustion in order to establish not only what chemical species exist, but also the amount of each. The conditions under which combustion occurs are always idealized for the purpose of calculation and assumptions of homogeneity, steady state, and absence of wall effects are made which then allow the subject to be treated by the methods of steady state thermodynamics. Throughout this work such assumptions were made in determining the composition of the products of combustion.

Combustion in an internal combustion engine occurs under conditions which are often far from ideal. Consequently results based on such calculations generally tend to over estimate the performance of the engine. Thus it is necessary to investigate the above mentioned assumptions in order to predict, at least qualitatively, in so far as possible, what effects deviations from these idealizations might have upon the chemical composition of the working fluid and ultimately upon the performance of the engine.

Homogeneity

It is a long established fact that the mixture in an internal combustion engine is not entirely homogeneous⁽³⁴⁾. Aside from wall effects, which will be discussed later, the inhomogeneity arises during the induction of the fresh charge. During this process a mixture of air and both gaseous and liquid fuel in the form of droplets is inducted and mixed

with residual exhaust gas. Whether or not efficient mixing occurs depends largely on the design of the carburetor, manifold, and inlet valve as well as the engine speed. In any event the charge during compression is composed more or less of pockets rich in exhaust gas interspersed within regions of charge, some of which are richer and other leaner than the measured fuel-air ratio. When combustion occurs, the products are heterogeneous, reflecting the fact that the initial mixture was not homogeneous. Thus, for example, if a stoichiometric mixture of fuel and air were burned, one might expect to find evidence of products resulting from both rich and lean combustion. That is, carbon dioxide and hydrogen as well as oxygen would be present in concentrations in excess of expected equilibrium amounts. David and Leah⁽²⁰⁾ have an interesting discussion of inhomogeneities caused by improper mixing. The effect of these inhomogeneities, assuming the difficulties of ignition and flame propagation can be overcome, is to cause incomplete combustion of the charge unless the mixture is quite lean. Pockets of unburned or partially burned gas may exist in the combustion chamber. Thus one would expect to find somewhat lower thermal efficiencies and mean effective pressures whenever mixing is not complete.

Additional inhomogeneity resulting from the flame propagation process is manifest as a continuous stratification of the burned products. For a discussion of this phenomenon see Brown⁽³⁵⁾ or Hershey, Eberhardt, and Hottel⁽¹⁰⁾. This subject will be treated more extensively in the discussion of the theoretical model for the internal combustion engine in Chapter VI.

Steady State

The assumption of steady state is tantamount to the assumption of chemical equilibrium. The implication is that the system, comprising the products of combustion has been allowed to remain in a given state of temperature and volume long enough to reach a homogeneous condition. That is all chemical reactions have proceeded to such an extent that the rate of the forward reaction is just equalled by the rate of the reverse reaction. At this point, the free energy of the system is at its minimum value and no spontaneous change can occur within an isolated system. Thus, because this is an equilibrium state, the composition can be completely determined by the methods of classical thermodynamics. However, if the time available for combustion is short, true equilibrium may not be attained.

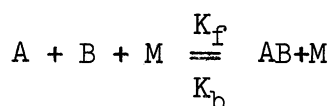
The time during which the system is maintained at constant conditions is termed the residence time. In an internal combustion engine the residence time, strictly speaking, is zero. The system volume and temperature are constantly changing. Fortunately this change is usually relatively slow compared to the time required to achieve equilibrium. This is especially true during combustion as compared to expansion.

To estimate the order of magnitude of the residence time in an internal combustion engine during combustion, one might consider the time required for one degree of crank revolution. During this period, changes in temperature and pressure are not large. For an engine operating at 2000 rpm this time is about 10^{-4} seconds. While it is difficult, if not impossible at this time to quantitatively predict the time interval necessary to reach near equilibrium conditions in a system as complex as the

working fluid of an internal combustion engine, it is known that at the temperatures and pressures existant during combustion the relaxation times for many of the chemical species present are from 10^{-6} to 10^{-8} seconds or less (36, 37). Thus one might expect that during combustion the chemical composition of the products could be calculated from equilibrium thermodynamics with very little error.

However during expansion, when temperatures and pressures are lower and many recombination reactions are taking place, equilibrium is not always maintained. This condition is termed equilibrium lag. The reactions lag and give rise to compositions which would be considered in equilibrium if the temperature were higher. Not all reactions lag the same amount and for a complex system as in the internal combustion engine, an exact analysis on this basis would be difficult to make and apply. Nevertheless a simple analysis is illustrative of the more complex system behavior.

Reference 37 describes a method whereby a pseudo temperature can be found for simple reactions which is suitable for the calculation of non-equilibrium compositions. In the case of recombination reactions requiring a three body collision between molecules A, B, and a third body M the following reaction can be written,



where K_f and K_b denote the forward and reverse reaction rates respectively.

The equilibrium constant K_c for this reaction is,

$$K_c = \frac{(AB)}{(A)(B)} = \frac{K_f}{K_b}$$

where the parentheses denote concentrations in moles per unit volume. The resultant expression for the pseudo temperature is (37),

$$T' = T + \frac{[1 + (C_v T / \Delta U)] [-\partial T / \partial t]}{K_f(M) [(A) + (B) + 1/K_c']}$$

where

T' = the pseudo temperature

C_v = the heat capacity at constant volume for the system

$-\Delta U$ = the heat of reaction at constant volume

$\frac{\partial T}{\partial t}$ = the reciprocal of the residence time per degree

K_c' = the equilibrium constant at T'

During the expansion stroke of an internal combustion engine, $-\partial T / \partial t$ is of the order of 10^5 to 10^6 °K per second (see Appendix D).

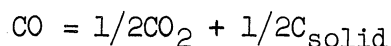
Reference 37 suggests the following values for common recombination reactions: $-\Delta U$ from 50 to 150 kcal/mole and K_f about 10^{11} (moles/liter)⁻²/sec. The quantity $(1 + C_v T / \Delta U)$ will be between 0 and 1 and consequently T' will be nearly equal to T unless the concentrations of A and B are very small in which case further recombinations will have only small effects.

It must be remembered that the above numerical values will change as the state of the system changes and in particular the rate of recombination will become lower as the pressure and temperature of the system are reduced. Moreover other types of reactions occur which under certain circumstances may be quite slow. The decomposition of NO into N_2

and O_2 is an example of such a reaction. Concentration of NO considerably in excess of equilibrium amounts have been observed in the exhaust of both rocket and internal combustion engines. In the analysis of the expansion process in rocket engines where $-\partial T/\partial t$ is 10^7 or 10^8 (46), chemical equilibrium is normally assumed. Thus the assumption of chemical equilibrium during expansion for internal combustion engine calculations is at least as good as it is for rocket engine calculations.

The effect of assuming chemical equilibrium during expansion is to increase the thermal efficiency and indicated mean effective pressure of the engine cycle. This is because the recombination of the radicals releases energy which can be used to do work.

There is a temperature, however, below which the rate of reaction is so slow that for all practical purposes no further reaction occurs and the equilibrium is effectively frozen. This temperature varies with the particular reaction. Gaydon⁽³⁶⁾ recommends $1500^\circ K$ for the reaction



and Hottel⁽¹¹⁾ suggests $1600^\circ K$ for the water gas reaction. Some early studies of Lovell and Boyd⁽³⁸⁾ suggest $1600-1700^\circ K$ as the temperature at which the equilibrium freezes in the exhaust of an internal combustion engine. However, $1500^\circ K$ is used throughout this work as the temperature below which no further changes occur in composition.

Wall Effects

The presence of a wall can effectively quench a combustion reaction. This phenomena occurs because the walls absorb heat or chain carriers, one or both of which must be transferred ahead of the flame to

the unburned gas in sufficient quantity to continue the propagation of the flame. Thus in the region near a wall, unreacted or partially reacted quantities of mixture may be found. In an engine this region appears as a layer a few thousands of an inch thick around the inside of the combustion chamber. Daniel⁽³⁹⁾ presents photographs which illustrate the presents of this layer. It is commonly thought that quenching is a major cause of partially reacted products, such as CH_4 , in the exhaust.

Because this dead space is a region of partially reacted or unreacted mixture, its effect is to reduce the thermal efficiency and mean effective pressure of an internal combustion engine.

Summary

In summary, inhomogeneities in the original mixture and quenching at the wall result in incomplete combustion of the charge. Even if the charge were completely homogeneous and were all burned, the rate of reaction may be so slow for some reactions that a state of chemical equilibrium is not attained in the products, especially during the expansion part of the cycle. Nevertheless to circumvent the inherent difficulties of treating these deviations from ideality, complete homogeneity and chemical equilibrium will be assumed to exist for the purposes of calculating the composition of the working fluid above 1500°K in the analytical model of the internal combustion engine. Below 1500°K , the composition will be assumed frozen at that composition which existed at 1500°K . These assumptions will always produce calculated thermal efficiencies and mean effective pressures which are higher than those found if the exact composition of the working fluid were used.

2. The Chemical Composition at Equilibrium

a. General

The products of a hydrocarbon-air reaction consist of a number of atoms and molecules whose individual amounts vary with changes in pressure and temperature according to the laws of chemical kinetics, subject to the condition that the mass of each basic atomic specie remains constant. When the condition of chemical equilibrium is imposed on the system, the amount of each constituent can be found by purely thermodynamic consideration. For the range of pressures and temperatures encountered in the internal combustion engine, 10 gaseous species must be considered in the products:

Reactants	Products	Specie No.
Fuel-C _a H _b	--	1
Air{ -O ₂ -N ₂	O ₂	2
	N ₂	3
	CO ₂	4
	H ₂ O	5
	CO	6
	H ₂	7
	OH	8
	NO	9
	O	10
	H	11

Additional species may be considered such as N, NO₂, C, or CH₄. However their amounts, calculated from equilibrium considerations, will be extremely small and their contribution to the overall thermodynamic properties of the system will be negligible.

In order to establish the amount of each of the 10 constituents 10 equations are required. Since some of these equations are non-linear,

an explicit solution can not be obtained and resort must be made to a trial and error solution. Many solution techniques have been developed for such equations. Some are specifically directed at the solution of equations for the combustion of a hydrocarbon fuel with air and some are quite general in scope and are capable of treating systems of arbitrary reactants with solid, liquid or gaseous products. Hershey, Eberhardt, and Hottel⁽¹⁰⁾ as well as Goodenough and Felbeck⁽⁸⁾ employed the former, or specific type of solution, to their combustion equations. Vickland⁽⁴⁰⁾ analyzed a number of methods, both general and specific, finally adopting a combination of them which was suitable for a computer solution. Huff and Co-workers⁽⁴¹⁾ proposed a method for treating arbitrary reactants and solid, liquid, or gaseous products which has been widely used for combustion calculations. Brinkly⁽³⁷⁾ proposed still a different method for treating a general system. This method was used by Edson⁽²²⁾. A summary and comparison of the most widely used general methods is presented in Zeleznik⁽⁴²⁾. In comparing the rate of convergence for the general methods considered, it was found in Reference 42 that no significant advantage was obtained using any one general method in preference to another.

In spite of the availability of numerous methods for computing equilibrium compositions, many of which provide for the simultaneous computation of the adiabatic flame temperature, it was felt that a much more efficient solution might be obtained if a computational method were used which was not general but was explicitly developed to solve only those equations which arise in the combustion of a hydrocarbon fuel with air. Such a system would contain carbon, hydrogen, oxygen, and nitrogen, and be comprised of the 10 listed species.

The computation scheme which is illustrated in the following section is based on the work of Ritter von Stein⁽⁴³⁾. An adaptation of this method for the combustion of a hydrocarbon fuel and air is presented by Schmidt⁽⁴⁴⁾.

b. Computation of Homogeneous Gas Equilibrium

A number of assumptions have been made or implied so far concerning the nature of the products of combustion. The following assumptions are essential to the computation of the gas composition:

- 1) The products are in a state of chemical equilibrium.
- 2) All the products are gaseous and can be treated as perfect gases.
- 3) The system comprising the products of combustion is homogeneous and is maintained at constant temperature and volume.
- 4) The fuel is a pure hydrocarbon of the form C_aH_b .
- 5) Air is a mixture of O_2 and N_2 .
- 6) Only 10 species exist in the products.
- 7) The mass of each basic atomic specie is known and remains constant.

The 10 equations necessary to determine the amount of each of the 10 constituents are determined from a consideration of the following:

- 1) The conservation of mass.
- 2) The law of mass action, governing chemical equilibrium.
- 3) The temperature and volume of the system.

The Conservation of Mass

The reactants are supposed to consist entirely of fuel, C_aH_b , and air, O_2 and N_2 , and contain N_c mole atoms of carbon, N_h mole atoms of

hydrogen, N_O mole atoms of oxygen, and N_n mole atoms of nitrogen. Four mass balance equations can then be written between the number of mole atoms of each basic specie in the products and that in the reactants.

To determine the number of mole atoms of carbon in the products the following equation can be written:

$$\text{mole atoms of carbon in products} = n_{CO_2} + n_{CO}$$

where n is the number of moles of a particular specie. Re-arranging this equation in terms of partial pressures yields:

$$\text{mole atoms of carbon in products} = V/RT(p_{CO_2} + p_{CO})$$

where T and V are the temperature and volume of the system. Since the mole atoms of carbon in the products must be equal those in the reactants

$$N_c = V/RT(p_{CO_2} + p_{CO})$$

Derived in a similar manner the four mass balance equations are:

$$F \cdot N_c = CO_2 + CO \tag{2-5}$$

$$F \cdot N_h = 2H_2O + 2H_2 + OH + H \tag{2-6}$$

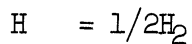
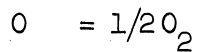
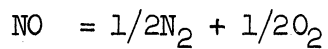
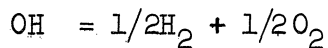
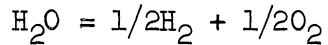
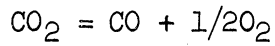
$$F \cdot N_O = 2CO_2 + H_2O + 2O_2 + OH + NO + CO + O \tag{2-7}$$

$$F \cdot N_n = 2N_2 + NO \tag{2-8}$$

where $F = RT/V$ and the formula names have been substituted for the partial pressures. The quantities on the right side of the equations are the "atom pressures" referred to by Ritter von Stein.

The Law of Mass Action

The following reactions are considered to occur among the products:



By assuming the products to be perfect gases, the expressions for the equilibrium constants of these reactions can be written in terms of the partial pressures of the constituents. Letting $\text{O}_2 = X^2$, $\text{H}_2 = Y^2$, and $\text{N}_2 = Z^2$, they are

$$K_1 = \frac{\text{CO} \cdot X}{\text{CO}_2} \quad (2-9)$$

$$K_2 = \frac{Y^2 \cdot X}{\text{H}_2\text{O}} \quad (2-10)$$

$$K_3 = \frac{Y \cdot X}{\text{OH}} \quad (2-11)$$

$$K_4 = \frac{Z \cdot X}{\text{NO}} \quad (2-12)$$

$$K_5 = \frac{X}{\text{O}} \quad (2-13)$$

$$K_6 = \frac{Y}{\text{H}} \quad (2-14)$$

These six equations together with the 4 mass balance equations must then be solved together for the 10 unknown partial pressures. The equilibrium constants can be calculated by the method of Appendix C, Section 5.

Solution

By re-arranging and substituting equations 2-9 through 2-14 into equations 2-5 through 2-8, the number of equations to be solved is reduced to four.

$$F \cdot N_c = CO(1 + \frac{X}{K_1}) \quad (2-15)$$

$$F \cdot N_h = 2(Y^2 + \frac{Y^2X}{K_2}) + \frac{YX}{K_3} + \frac{Y}{K_6} \quad (2-16)$$

$$F \cdot N_o = 2(\frac{CO \cdot X}{K_1}) + \frac{Y^2X}{K_2} + 2X^2 + \frac{YX}{K_3} + \frac{ZX}{K_4} + CO + \frac{X}{K_5} \quad (2-17)$$

$$F \cdot N_n = 2Z^2 + \frac{ZX}{K_4} \quad (2-18)$$

Solving Equation (2-18) for the positive value of Z,

$$Z = -\frac{X}{4K_4} + \sqrt{(\frac{X}{4K_4})^2 + \frac{F \cdot N_n}{2}} \quad (2-19)$$

and Equation (2-16) for the positive value of Y,

$$Y = \frac{-\left(\frac{X}{K_3} + \frac{1}{K_6}\right) + \sqrt{\left(\frac{X}{K_3} + \frac{1}{K_6}\right)^2 + 8 \left[1 + \frac{X}{K_2}\right] \cdot F \cdot N_h}}{4\left(1 + \frac{X}{K_2}\right)} \quad (2-20)$$

If Equation (2-15) is then substituted into Equation (2-17),

$$F \cdot N_o = \frac{\frac{X \cdot F \cdot N_c}{K_1}}{1 + \frac{X}{K_1}} + \frac{Y^2X}{K_2} + 2X^2 + \frac{YX}{K_3} + \frac{ZX}{K_4} + F \cdot N_c + \frac{X}{K_5} \quad (2-21)$$

In light of Equations (2-19) and (2-20), Y and Z are functions of X only. Then Equation (2-21) is only a function of X, since the equilibrium constants are only a function of temperature. It then remains to obtain the positive value of X which satisfies Equation (2-21). A method for doing this is presented in Appendix E along with an estimate of the rate of convergence of the method.

The quantity X having been determined, Y and Z can be calculated from Equations (2-19) and (2-20). The partial pressures can be calculated from Equations (2-9) through (2-15) re-arranged as follows:

$$O_2 = X^2 \quad (2-22)$$

$$N_2 = Y^2 \quad (2-23)$$

$$CO_2 = \frac{F \cdot N_c}{K_1 / X + 1} \quad (2-24)$$

$$H_2O = \frac{XY^2}{K_2} \quad (2-25)$$

$$CO = \frac{CO_2 \cdot K_1}{X} \quad (2-26)$$

$$H_2 = Y^2 \quad (2-27)$$

$$OH = \frac{XY}{K_3} \quad (2-28)$$

$$NO = \frac{XZ}{K_4} \quad (2-29)$$

$$O = \frac{X}{K_5} \quad (2-30)$$

$$H = \frac{Y}{K_6} \quad (2-31)$$

The number of moles of each constituent is the partial pressure divided by the factor 'F'.

Figures 2a through 2j show calculated mole fractions of the various constituents in the products as a function of the temperature and pressure of the system for a stoichiometric mixture of octane and air.

These curves are in good agreement with those of Reference 40.

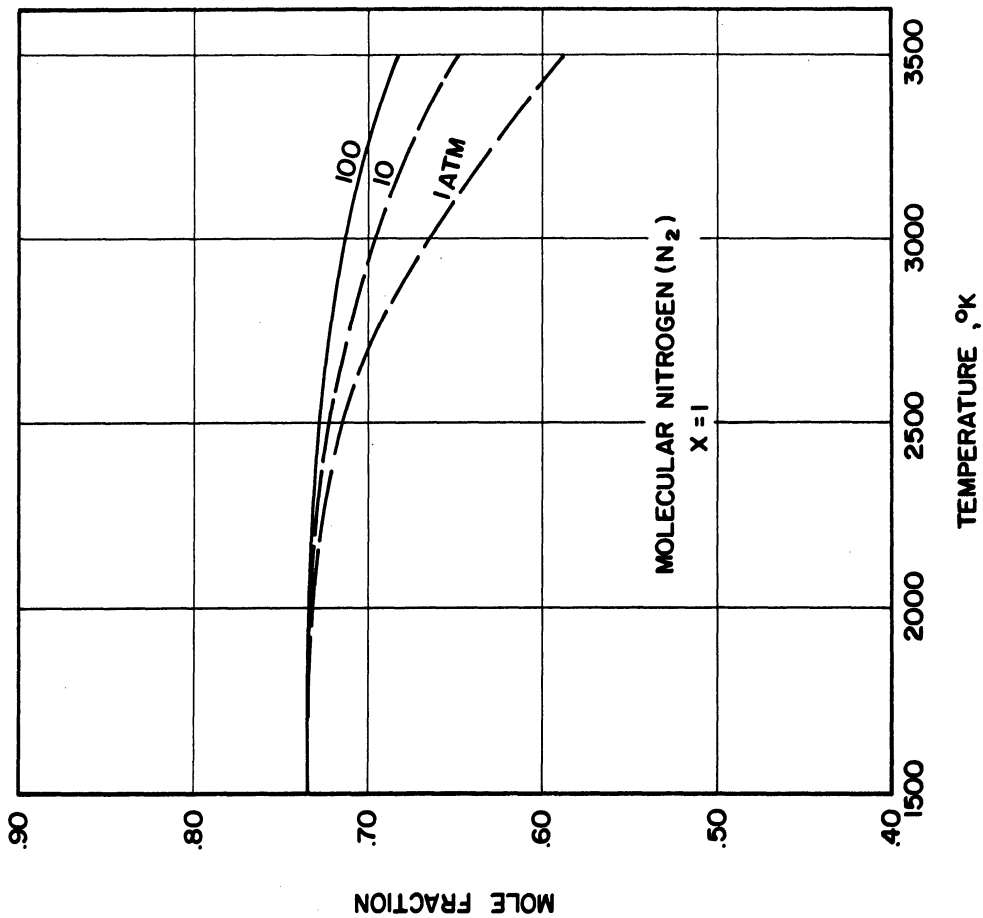


Figure 2b. Mole fraction vs temperature, N_2 .

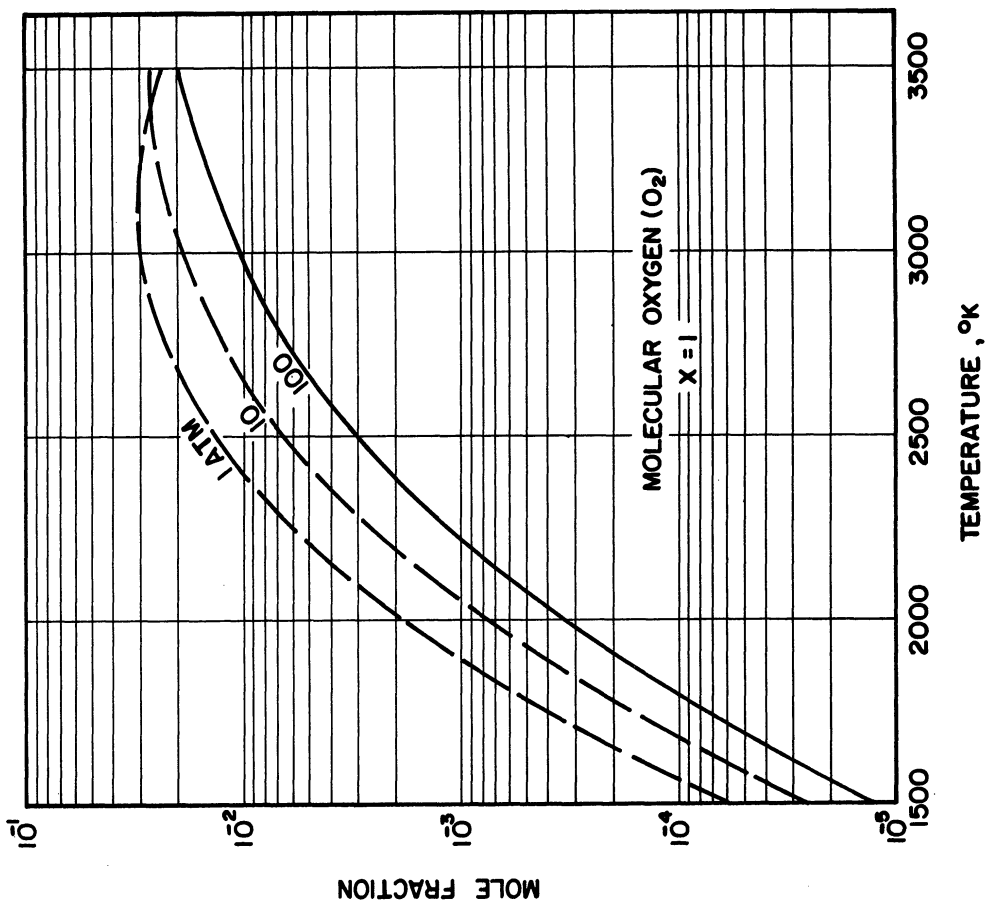


Figure 2a. Mole fraction vs temperature, O_2 .

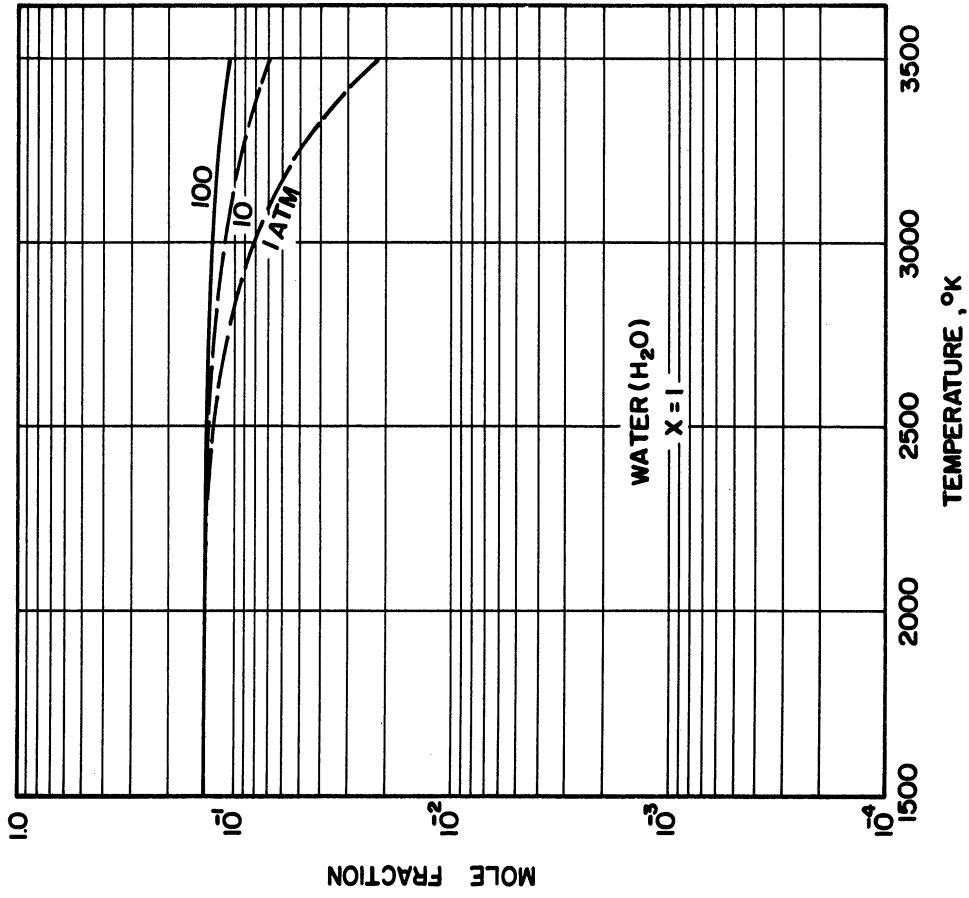


Figure 2d. Mole fraction vs temperature, H₂O.

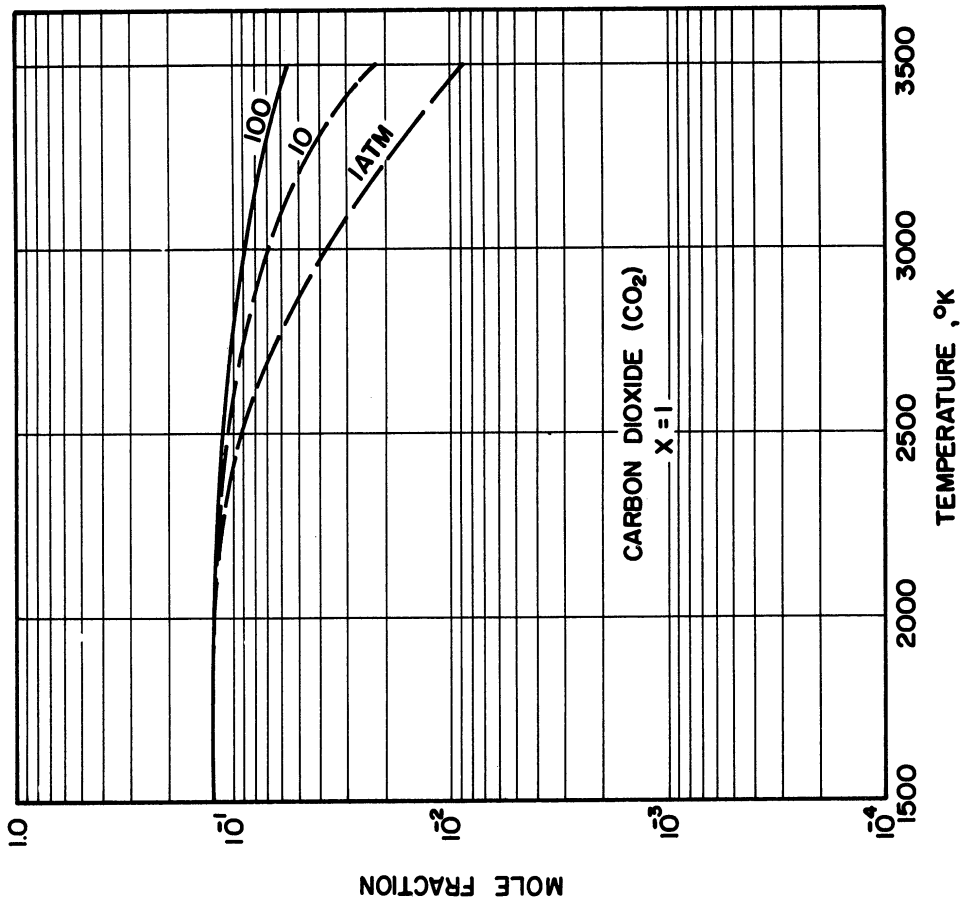


Figure 2c. Mole fraction vs temperature, CO₂.

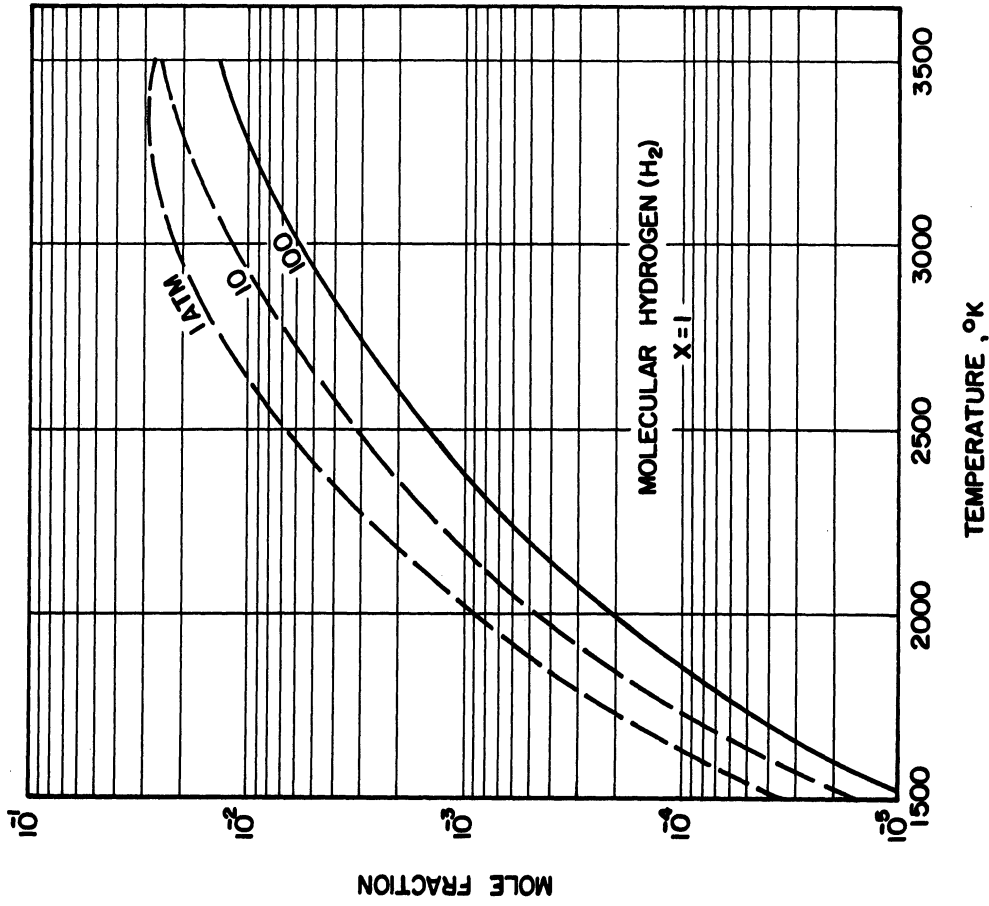


Figure 2f. Mole fraction vs temperature, H₂.

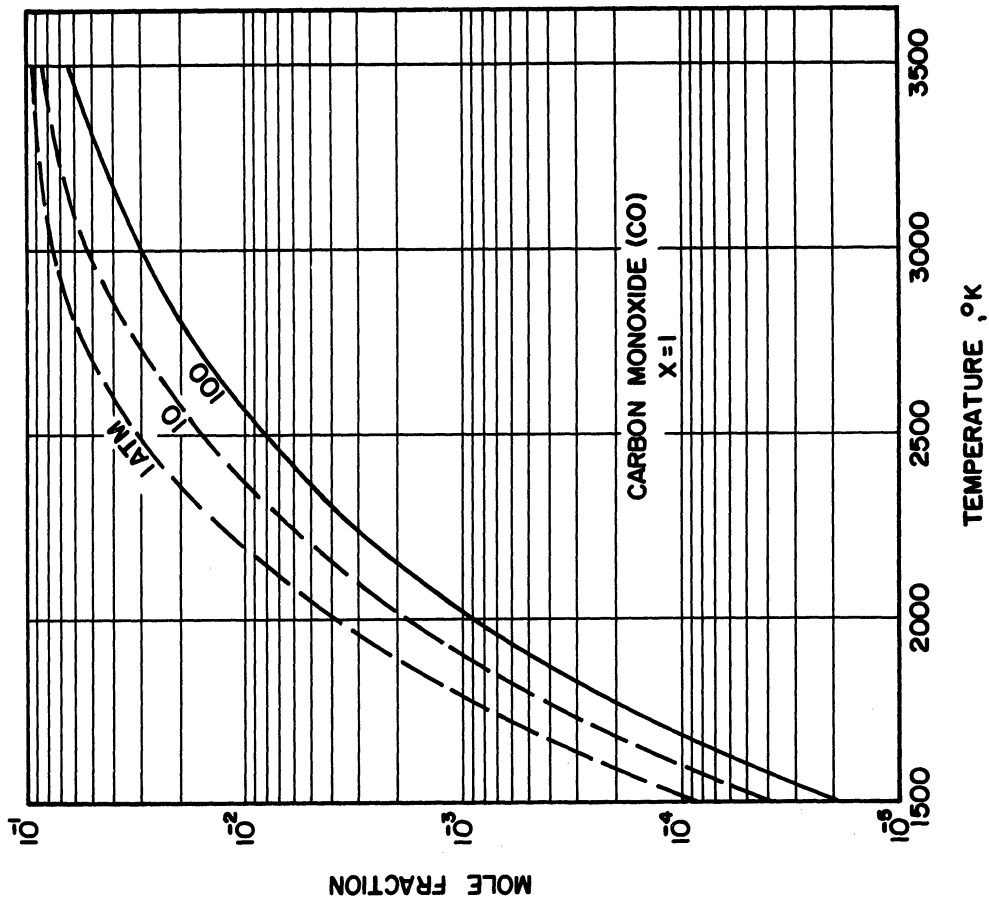


Figure 2e. Mole fraction vs temperature, CO.

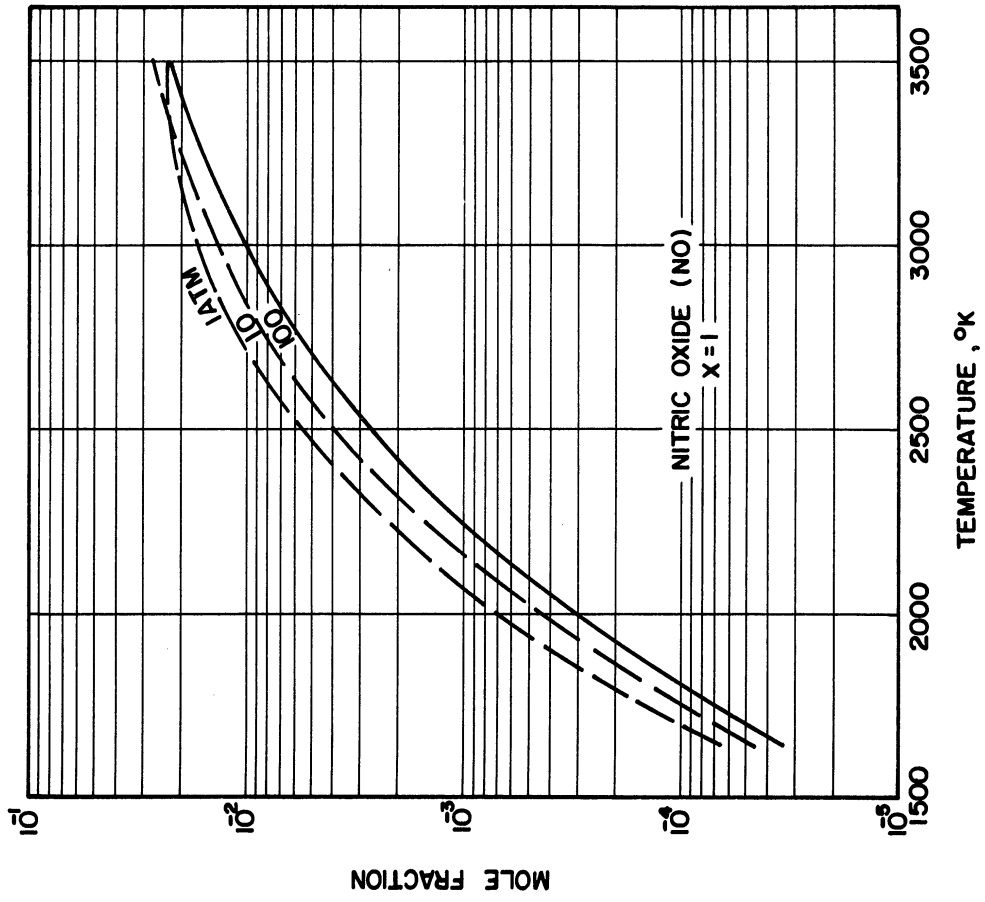


Figure 2h. Mole fraction vs temperature, NO.

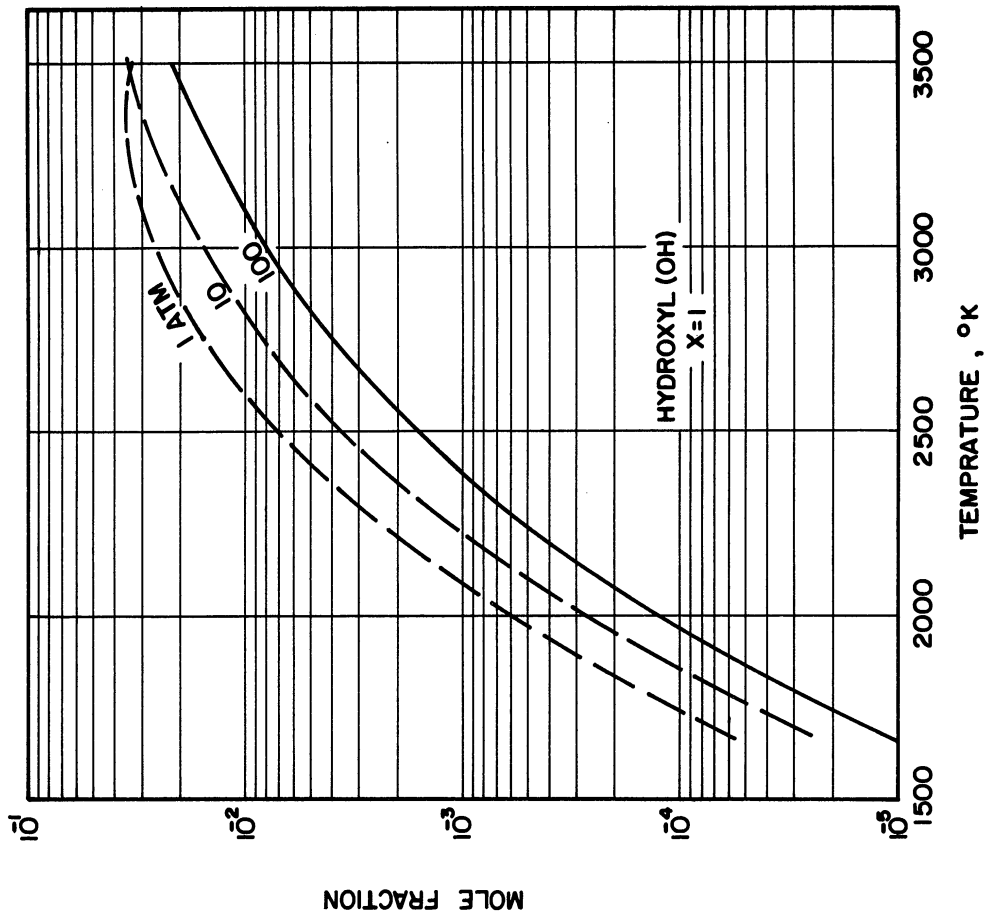


Figure 2g. Mole fraction vs temperature, OH.

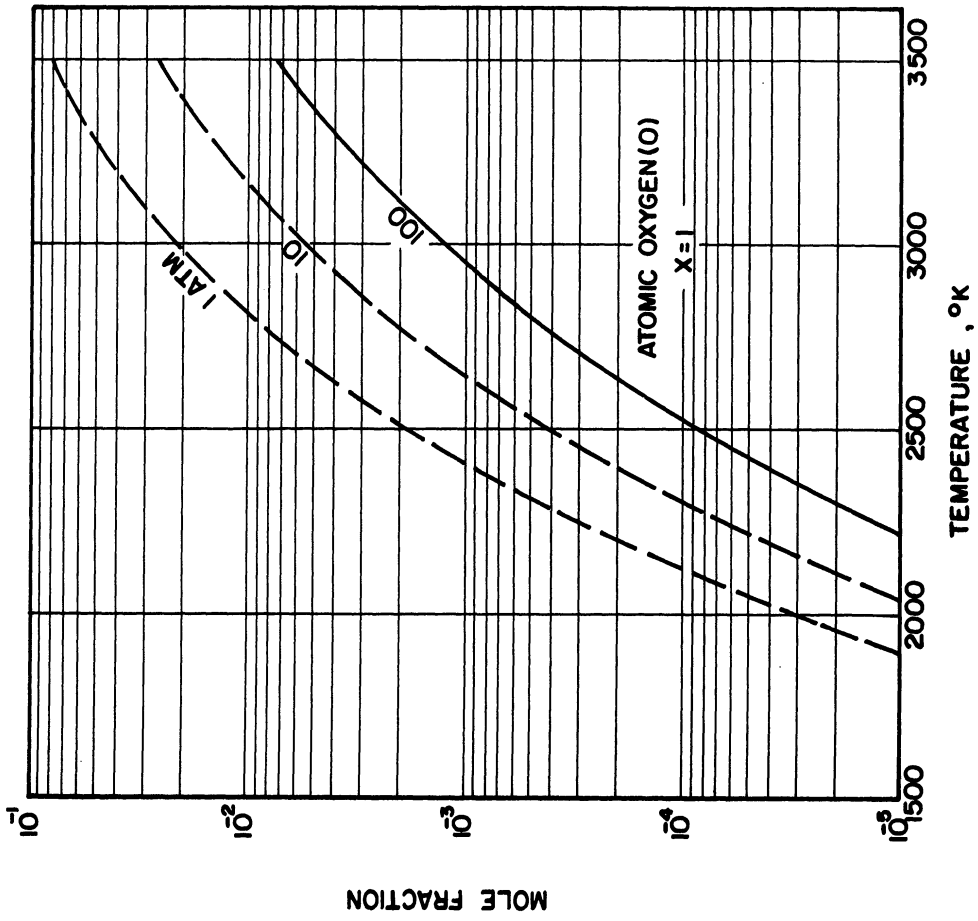


Figure 2j. Mole fraction vs temperature, O.

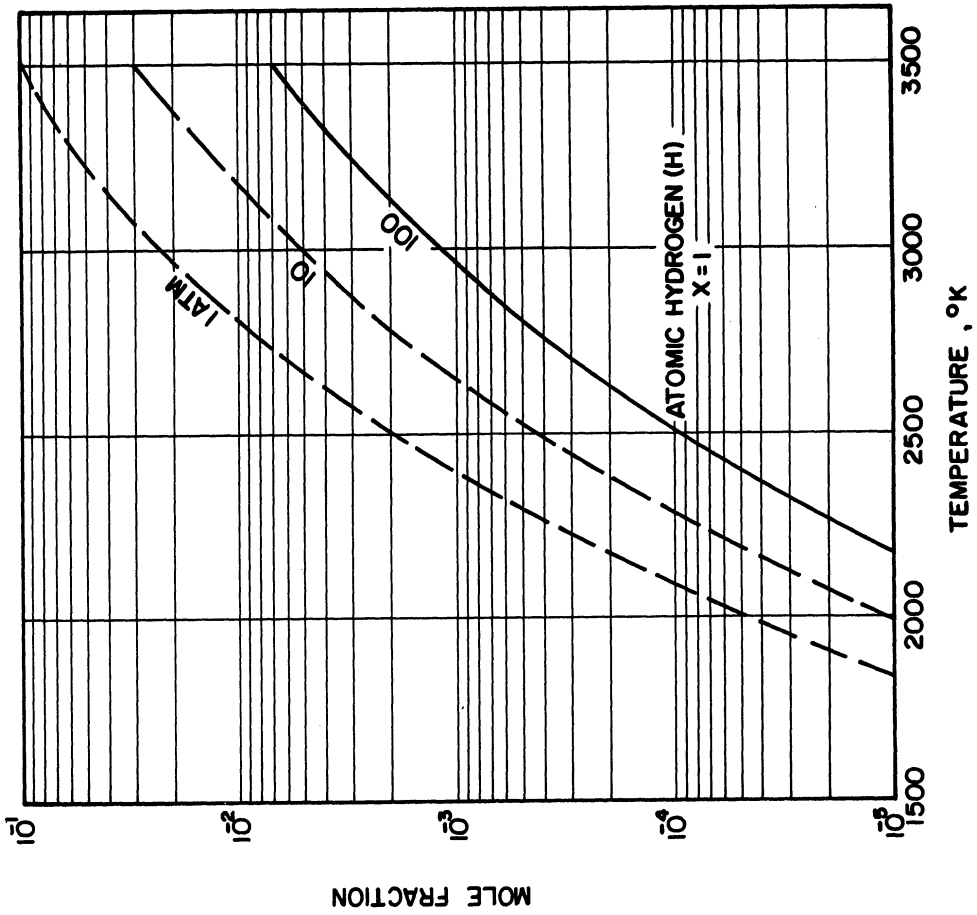


Figure 2i. Mole fraction vs temperature, H.

In order to calculate the composition at a given temperature and pressure, rather than volume, a second trial and error solution must be made. If an initial estimate is made of what the volume might be at the given temperature and pressure and the calculations made as above, the total number of moles can be computed. From the number of moles, the pressure, and the temperature, a new estimate of the volume can be made and the calculations repeated once more, leading to yet a third estimate of the volume. The entire process can then be repeated a third time and so on until the volumes calculated for any two successive iterations are within a specified amount of each other. For a reasonable good estimate of the volume initially, only about two additional iterations are required to achieve agreement in the fifth or sixth digit between two successive values of the volume.

c. Computation of Frozen Equilibrium

The products of combustion below 1500°K are assumed frozen. In addition to the assumptions made in connection with homogeneous gas equilibrium in Part 2 of this section the following additional assumptions are made for frozen equilibrium:

- 1) The composition of the products is frozen at the equilibrium composition which would exist at 1500°K.
- 2) For lean and chemically correct mixtures, only O_2 , N_2 , CO_2 , and H_2O are present.
- 3) For rich mixtures, only N_2 , CO_2 , H_2O , CO , and H_2 are present.

Lean and Stoichiometric Mixtures

For the calculation of the composition of lean and stoichiometric mixtures under the assumption of frozen equilibrium the mass balance equations are:

$$F \cdot N_c = CO_2$$

$$F \cdot N_h = 2H_2O$$

$$F \cdot N_o = 2CO_2 + H_2O + 2O_2$$

$$F \cdot N_n = 2N_2$$

Finally, the partial pressures of the constituents are:

$$CO = F \cdot N_c \quad (2-32)$$

$$H_2O = F \cdot N_h/2 \quad (2-33)$$

$$O_2 = \left| F \cdot N_o/2 - F \cdot N_h/4 - F \cdot N_c \right| \quad (2-34)$$

$$N_2 = F \cdot N_n/2 \quad (2-35)$$

where the absolute value of the partial pressure of the oxygen is taken to avoid the possibility of an extremely small negative value of this quantity occurring near the stoichiometric point due to round-off errors. This consideration applies to computer solutions only. The number of moles of each constituent is its partial pressure divided by 'F'.

Rich Mixtures

For the calculation of the composition of rich mixtures under the assumption of frozen equilibrium the mass balance equations are:

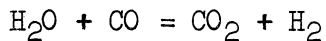
$$F \cdot N_c = CO_2 + CO$$

$$F \cdot N_h = 2H_2O + 2H_2$$

$$F \cdot N_0 = 2CO_2 + CO + H_2O$$

$$F \cdot N_n = 2 \cdot N_2$$

The following reaction is assumed to occur among the products of rich mixtures at 1500°K



The expression for the equilibrium constant of this reaction is

$$K = \frac{CO_2 \cdot H_2}{H_2O \cdot CO} \quad (2-36)$$

As this reaction involves no change in the total number of moles, the computation at a given temperature and pressure is equivalent to that at a given temperature and volume. And in fact, the reaction is independent of the pressure of the system. Therefore the composition of the products depends only on the temperature. As a result, fixing the value of the equilibrium constant fixes the composition. Whenever the composition of the products is to be calculated when the temperature is below 1500°K, K is given its value at 1500°K. This freezes the reaction at the temperature level. Letting $Z = \frac{F \cdot N_c}{CO_2}$ and substituting the mass balance equations in the equation for the equilibrium constant,

$$K = \frac{1/2 \frac{N_h}{N_c} - \frac{N_0}{N_c} + 1 + 1/Z}{\left(\frac{N_0}{N_c} - 1 - 1/Z\right) (Z - 1)}$$

At 1500°K, K is 0.3747⁽²⁹⁾. Solving the above equation and substituting the value of K,

$$Z = \alpha + \sqrt{\alpha^2 + \frac{6.6752}{\left(\frac{N_O}{N_C} - 1\right)}} \quad (2-37)$$

where

$$\alpha = \frac{\left(1 + \frac{N_h}{0.7494 \cdot N_C}\right)}{\left[\frac{N_O}{N_C} - 1\right]} - 1.6688$$

The partial pressures are:

$$CO_2 = \frac{F \cdot N_C}{Z} \quad (2-38)$$

$$H_2O = F(N_O - N_C) - CO_2 \quad (2-39)$$

$$CO = |F \cdot N_C - CO_2| \quad (2-40)$$

$$H_2 = \left|F \cdot \frac{N_h}{2} - H_2O\right| \quad (2-41)$$

$$N_2 = F \cdot \frac{N_h}{2} \quad (2-42)$$

Again absolute values are taken to assure that near the stoichiometric point no small negative partial pressures can be calculated due to round-off errors. The number of moles of each constituent is its partial pressure divided by 'F'.

3. Computer Solution of the Equations of Chemical Equilibrium

The solution technique developed in Part 2 and Appendix E has been programmed on a digital computer. A flow diagram for this program is shown on Figure 23. Appendix E includes a discussion of many important features of the solution. A listing of the program written in the MAD language as an external function called AMOUNT is included in Appendix I.

III. SIMULATION OF THE FUEL-AIR ENGINE CYCLE ON A DIGITAL COMPUTER

A. General

As the initial step toward the development of an improved model for the analysis of the internal combustion engine which could ultimately be programmed on a digital computer, the constant volume fuel-air engine cycle was programmed. The details of the mathematical analysis and computer program are included in Appendix F. The techniques developed in Chapter II have been used to calculate the composition and thermodynamic properties of the working fluid.

The fuel-air cycle is composed of the following idealized processes (see Figure 3).

Process 1-2. An isentropic compression of the working fluid from volume V_1 to V_2 .

Process 2-3. An adiabatic constant volume combustion.

Process 3-4. An isentropic expansion of the working fluid from volume V_2 to V_1 .

Process 4-4. A further expansion to the exhaust pressure. That portion remaining in the cylinder undergoes an isentropic expansion.

Process 4-5. A constant pressure adiabatic expulsion of the burned gas leaving some residual exhaust gas in the clearance volume.

Process 5-1. An adiabatic induction and mixing of the fresh charge with the residual exhaust at the intake pressure, thus completing the cycle.

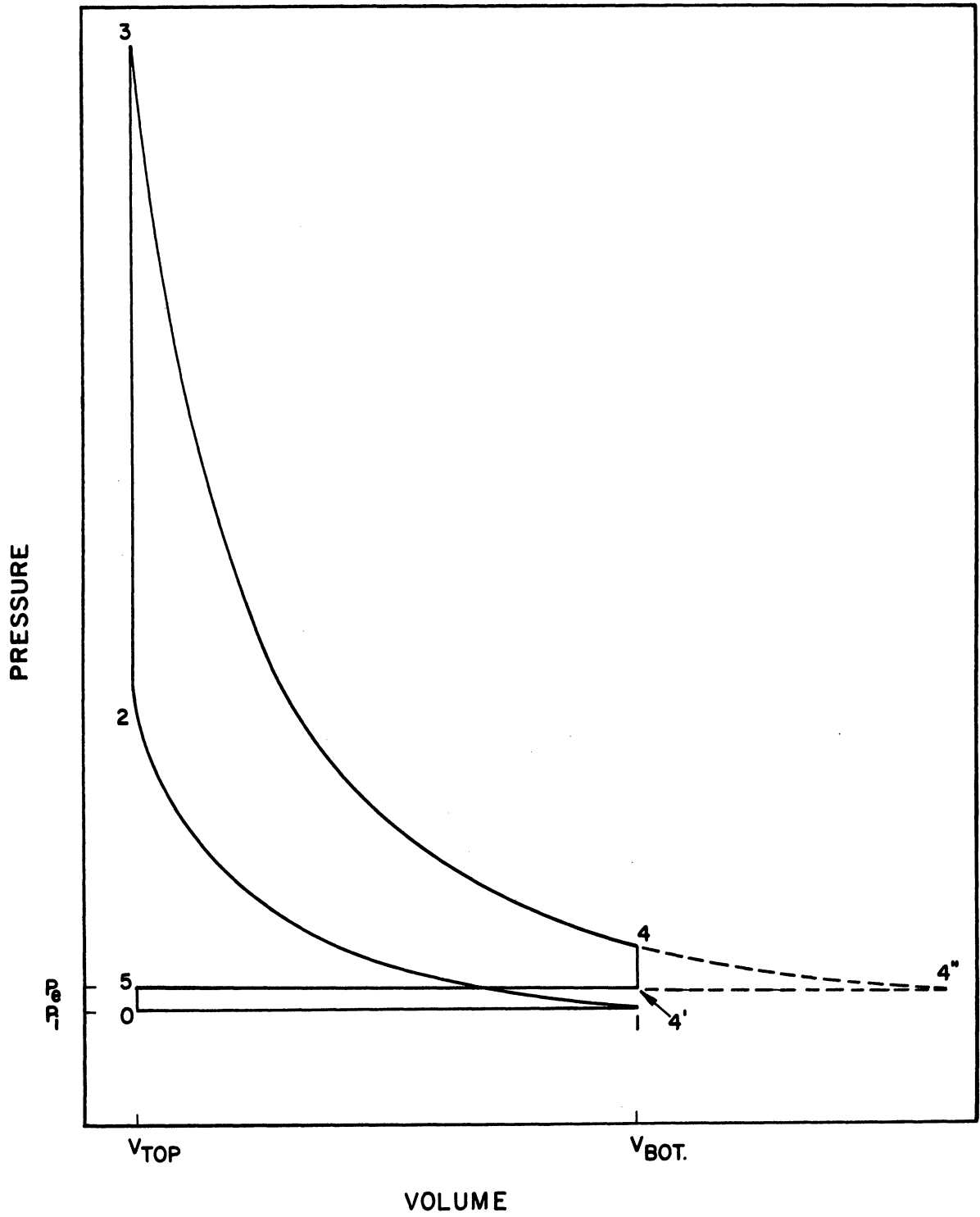


Figure 3. Indicator diagram for the fuel-air cycle engine at part throttle.

The following assumption were made:

1. There is no chemical change in either the fuel or air prior to combustion.
2. After combustion and during expansion, the products are in a state of chemical equilibrium.
3. Equilibrium is frozen below 1500°K.
4. All processes are adiabatic.
5. There are no irreversible fluid flow effects.
6. All valve events occur at top or bottom center.

A complete description of this cycle may be found in References 10 and 45. Calculations made using similar analytical models can be found in References 7, 9, 10, 11, 22, and 45. Reference 22 is of particular interest as the calculations were made on a digital computer.

In as much as a great body of calculations already has been reported using the fuel-air cycle engine model, only selected values of the computer computations will be presented here. These deal with the effects of lean mixtures and high compression ratios on thermal efficiency and mean effective pressure. They are of special interest in light of the increasing effort to raise compression ratios and utilize leaner mixtures in internal combustion engines. Thermodynamic charts available for the calculation of the fuel-air cycle are constructed for mixtures no leaner than 120 percent theoretical air and allow computations for compression ratios up to about 20 to 1⁽¹¹⁾. Both Tizard and Pye⁽⁷⁾ and Goodenough and Baker⁽⁹⁾ presented results of calculations for the effect of very lean mixtures on thermal efficiency. However these results are based upon now outdated heat capacity and dissociation data and their analytical models differ somewhat from that employed here.

B. Results and Discussion

Figures 4 and 5 are plots of the results calculated by the aforementioned computer program for the fuel-air cycle engine. Figure 4 shows the effects of both compression ratio and air-fuel ratio on the thermal efficiency and mean effective pressure of lean octane-air mixtures. As the mixture becomes leaner, the efficiency approaches that of the air-cycle, Equation (1-2), where 'k' is 1.4 for air. It has been proposed by both Tizard and Pye and Goodenough and Baker that the thermal efficiency of the fuel air cycle can be expressed as

$$\eta = 1 - \left(\frac{1}{r}\right)^n$$

where 'n' is a quantity derived from fuel-air cycle calculations. Goodenough and Baker proposed, for lean octance-air mixtures, that

$$n = 0.3867 - \frac{6.5}{100x - 35} - \frac{0.043}{r}$$

where 'x' is the fraction of theoretical air and 'r' is the compression ratio. In order to better fit the curves of Figure 4, this expression was modified slightly. The modified expression,

$$n = 0.374 - \frac{0.1}{x} - \frac{0.043}{r} \quad (3-1)$$

will approximate the thermal efficiencies of Figure 4 within 0.5 percent.

Figure 5 shows the influence of compression ratio on the thermal efficiency and mean effective pressure of a lean octene-air mixture for compression ratios up to 30:1. Edson⁽²²⁾ reports similar calculation for ratios up to 300:1 for a number of fuels and fuel-air ratios. Thermal efficiency and mean effective pressure continually increase as compression ratio is increased. Equation (3-1) is a good approximation for the thermal efficiencies of Figure 5.

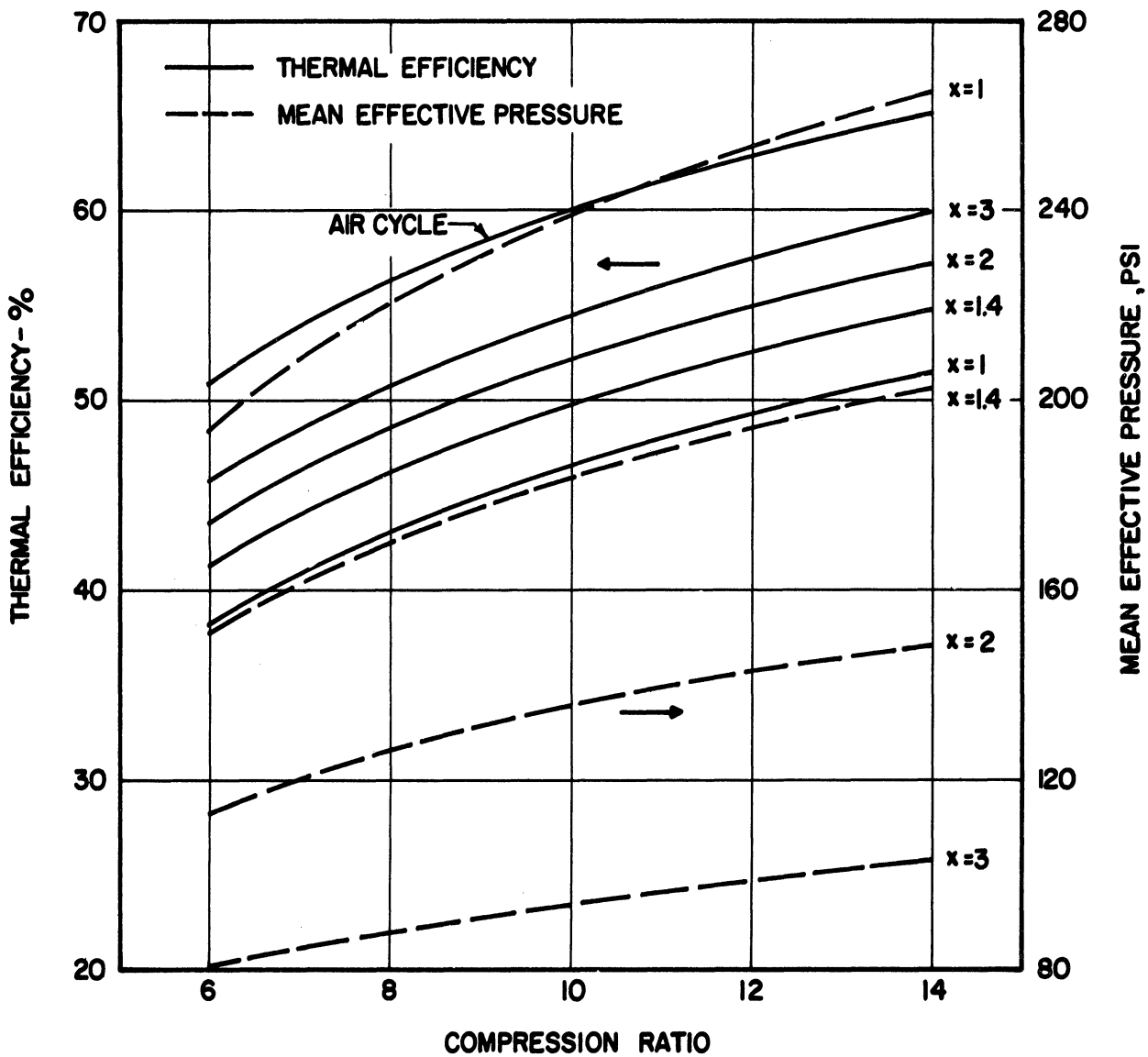


Figure 4. Efficiency and mean effective pressure vs compression ratio for lean mixtures. Fuel-Air cycle analysis. Fuel - octane, $T_i = 520^\circ\text{F}$, $P_i = P_e = 1 \text{ atm}$.

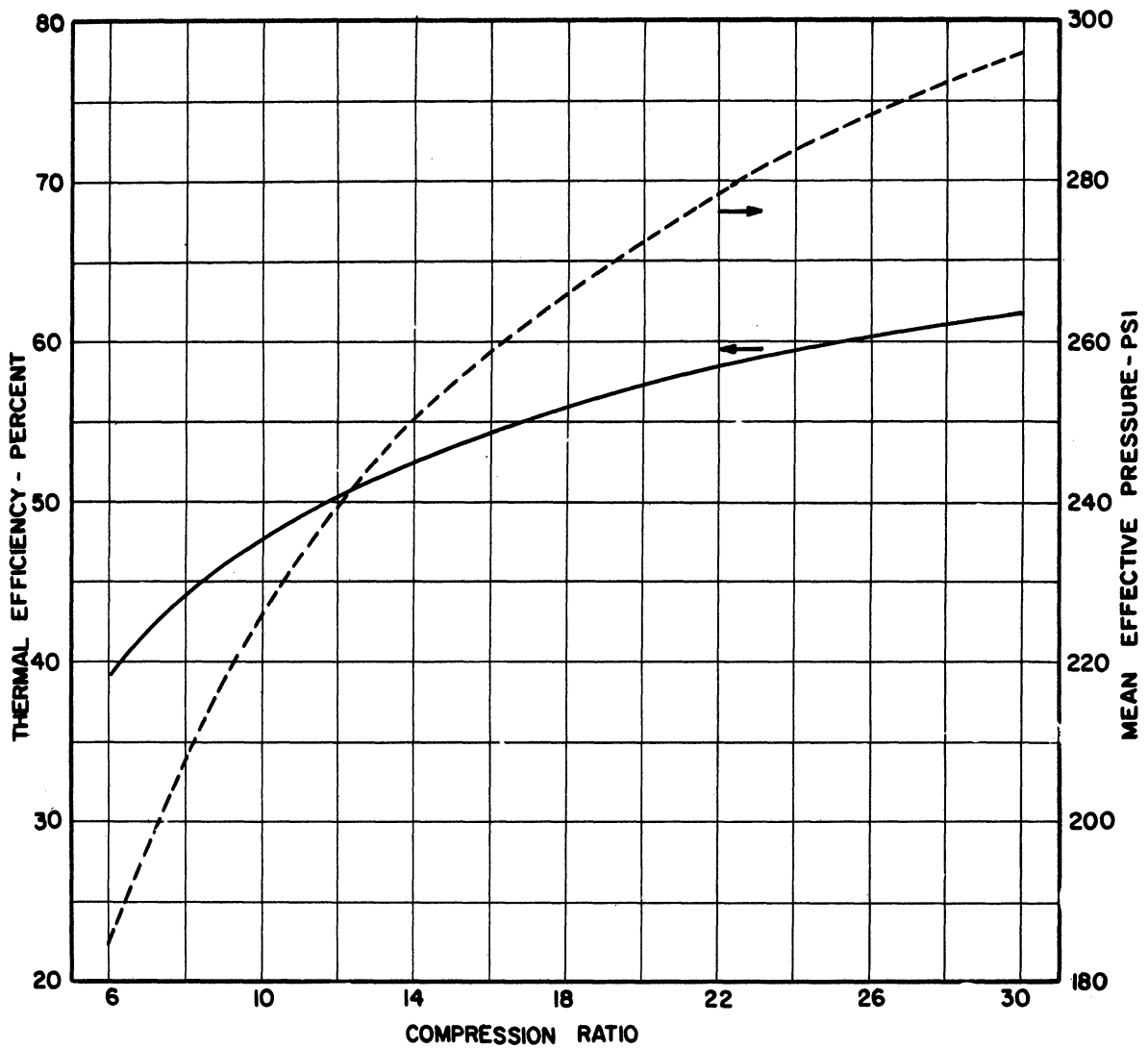


Figure 5. Efficiency and mean effective pressure vs compression ratio for a mixture of octene and 110 percent theoretical air. Fuel-air cycle analysis. $T_1 = 520$, $P_1 = P_e = 1$ atm.

IV. COMBUSTION IN A SPARK-IGNITED ENGINE

Whereas the fuel-air cycle incorporates many important features of the internal combustion engine, it totally disregards two phenomena which are of primary importance. These are the progressive burning of the charge and the heat transfer. Prior to a discussion of their inclusion into the analytical model for the internal combustion engine described in Chapter VI, some discussion of the nature of each is warranted. This chapter deals with the nature of the combustion process and the following chapter deals with that of the heat transfer.

A. General

The combustion process in an internal combustion engine has been investigated photographically and by other methods. The studies of Rassweiler, Withrow, and Cornelius (16, 17, and 18) reveal that in normal combustion the spark induces a kernel of flame which subsequently propagates regularly in all directions. Their photographs show that during combustion two distinct gaseous regions exist in the chamber, a region of burned gas surrounded by a region of unburned gas. The two regions are separated by a definite reaction zone. This reaction zone or flamefront, as it is often called, is generally a spherical shell during the early stages of combustion but becomes distorted as it approaches the wall and tends to assume the shape of the combustion chamber.

The apparent rate of flame propagation in an engine is very high. This is due to both the reaction rate and the expansion of the gases during the combustion process. At the beginning of combustion, the expansion rate is high compared to the reaction rate whereas at the end of combustion

it is low. The fundamental quantity to be investigated is the flame front motion due to the reaction only. This is the flame propagation relative to the unburned gas.

The remainder of this chapter is devoted to obtaining a general expression for the rate of flame propagation in an internal combustion engine. By necessity a number of assumptions have been made in order to simplify the equations involved. Perhaps the most serious of these are those of steady state and one dimensionality. Thus it must be recognized from the outset that the equation presented for the rate of flame propagation is only approximate. Its validity can be determined only by a direct comparison of calculated with experimental burning rates which can be done indirectly through a comparison of calculated and experimental pressure time diagrams for a particular engine.

B. Flame Propagation

Two regimes of flame propagation are recognized, the laminar and the turbulent. Laminar combustion is characterized by a well defined narrow flame front and a relatively low rate of reaction. Turbulent combustion is characterized by a less well defined thicker flame front and a much higher rate of reaction. Three somewhat separable modes of turbulent flame propagation have been observed^(47, 48). They are:

1. Weak turbulence which wrinkles the continuous laminar flame front, thereby causing a higher effective rate of reaction due to the greater surface area through which the basic transport phenomena can occur;

2. Stronger turbulence which disrupts the laminar flame front and creates a distribution of centers of deflagration which can be treated statistically;
3. Still stronger turbulence which shows its effects by locally diluting and preheating the initial centers of deflagration, such that a homogeneous reaction results in the limit.

Observed and calculated rates of flame propagation in internal combustion engines indicate that the combustion is in the turbulent regime (see Rabezanna, Kalmar, and Candelise⁽¹⁹⁾). However little quantitative information concerning the nature of the turbulence in internal combustion engines exists. Semenov⁽⁴⁹⁾ presents the results of some experimental studies which show that the turbulent fluctuations vary from point to point in the cycle, but sheds no light on the nature of the turbulence.

A number of theories have been advanced to explain turbulent flame propagation (see Reference 50). Only the first mode of turbulent flame propagation has been studied analytically. Such studies generally assume that the turbulent burning velocity is related to the laminar velocity by a factor which is proportional to the level of turbulence in the reaction zone.

The theory of Scurlock and Grover⁽⁵¹⁾, one of the most recent, assumes a one dimensional steady state situation in which premixed fuel and air are fed into the reaction zone at a rate such that the zone itself remains fixed relative to the source of the gas. As the level of turbulence is increased, the flow rate must be increased in order to prevent the reaction zone from moving upstream toward the source. The velocity at which

the premixed gases approach the reaction zone is equal in magnitude and opposite in direction to the flame propagation velocity. Scurlock and Grovers' analytical model assumes that the laminar flame front is wrinkled by large scale turbulence like that of the first mode. Figure 6 illustrates this wrinkling. The theory assumes that in this type of turbulent combustion the fundamental transport phenomena remain unchanged and hence the laminar flame speed normal to the flame front is unchanged. The increased rate of flame propagation due to turbulence is presumed to be caused only by the larger surface area over which the reaction can proceed.

The wrinkles in Figure 6 are assumed to be cones with an average height proportional to $\sqrt{\bar{Y}^2}$, the root mean square displacement of a flame element from its mean flame front position, and an average base width proportional to L_{Eu} , the Eulerian scale of turbulence. The ratio of turbulent to laminar flame speed, derived from purely geometric considerations, is

$$\frac{S_t}{S_l} = \frac{A_t}{A_l} = \left[1 + k_3 \left(\frac{\bar{Y}^2}{L_{Eu}^2} \right) \right]^{1/2} \quad (4-1)$$

where $k_3 = 4k_1^2/k_1^2$. Scurlock and Grover consider three effects which are believed important to a determination of \bar{Y}^2 . These are eddy diffusion, flame propagation, and flame generated turbulence.

Bolz and Burlage⁽⁴⁸⁾ have successfully used the theory of Scurlock and Grover to predict the growth of flame kernels in turbulent flow issuing from a jet. Mickelsen and Ernstein⁽⁵²⁾ in a similar experiment found good agreement between flame speeds obtained experimentally and those predicted by the theory of Scurlock and Grover over a range of fuel-air ratios and levels of

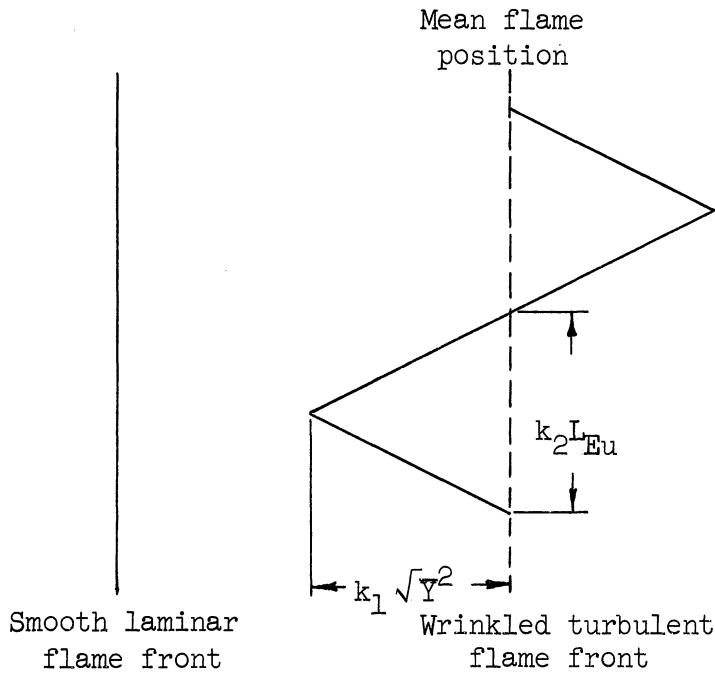


Figure 6. Wrinkled flame front model of Scurlock and Grover⁽⁵¹⁾.

turbulence. However the theoretical values were always higher than the experimental ones. They found that turbulent flame speed is a statistical quantity. Moreover their results indicated that the rate of laminar to turbulent flame speed (at least for propane) is not strongly dependent on fuel-air ratio.

Because quantitative information about the nature of the turbulence in the working fluid of an engine is not available, none of these theories of turbulent flame propagation can be applied directly. However a relationship between the laminar and turbulent flame velocity

can be made. Namely,

$$\frac{S_t}{S_l} = K \quad (4-2)$$

where K is a quantity dependent only on the level of turbulence in the reaction zone. K must be determined from experimental data and is only valid for a given mean level of turbulence. Now it remains to find an expression for the laminar flame propagation velocity.

The principle mechanisms of flame propagation are the transport of heat and active radicals from the reaction zone to the unburned gas ahead of the flame. A number of theories which take into account one or both of these mechanisms have been advanced to predict laminar propagation velocity (see Reference 50 and 53). Comprehensive theories which take into account both phenomena are difficult to apply and require information about reaction kinetics which is not known for many reactions. Hirshfelder et al. (54, 55) have proposed such a theory and have applied it to a few elementary reactions.

In a number of investigations of the burning velocity of hydrocarbon-air mixtures, less complex theories predict burning velocities which often agreed closely with experimental ones. In particular two theories have been widely employed for several investigations (for example Reference 58, 59, 60, and 61). One is the theory of Tanford and Pease⁽⁵⁷⁾ which predicts the velocity from a consideration of the concentration in the reaction zone of certain chemical species, such as H, O, and OH. These are supposed to act as agents to initiate and propagate the reaction. The other is the theory of Semenov⁽⁵⁶⁾ which predicts the burning velocity from purely

thermal consideration. Reference 60 reports particularly good agreement between experimental values and values calculated from the two theories for the maximum burning velocity of iso-octane-air mixtures at one atmosphere pressure over a range of widely different initial mixture temperatures. Whereas the two theories are based on seemingly unrelated phenomena, the consensus of opinion appears to be that no particular advantage is gained using one of these two simplified theories in preference to the other unless it is definitely known that the basic mechanism of the reaction in question favors one model. In general, equally good agreement has been obtained with either theory.

In light of the above discussion a decision was made to use the thermal theory of Semenov to predict the laminar burning velocity of the hydrocarbon-air mixtures considered in this current study. Reference 59, 60, 62, and 63 treat the reaction between a hydrocarbon such as iso-octane and air as second-order bimolecular. Clarke⁽⁶²⁾ reports experimental work which shows this reaction to be bimolecular up to 23 atmospheres.

C. Thermal Theory of Flame Propagation for A Bimolecular Second-Order Reaction

Because the thermal theory of laminar flame propagation is so important to the results calculated from the analytical model of Chapter VI, a general development of this theory is made here for a bimolecular second-order reaction. The original full derivation of this theory is in Reference 56. References 50 and 53 review the derivation.

The thermal theory of flame propagation assumes a one dimensional steady state combustion model (see Figure 7). The reaction is propagated solely by the heating of the unburned gas in Zone I, initially at tempera-

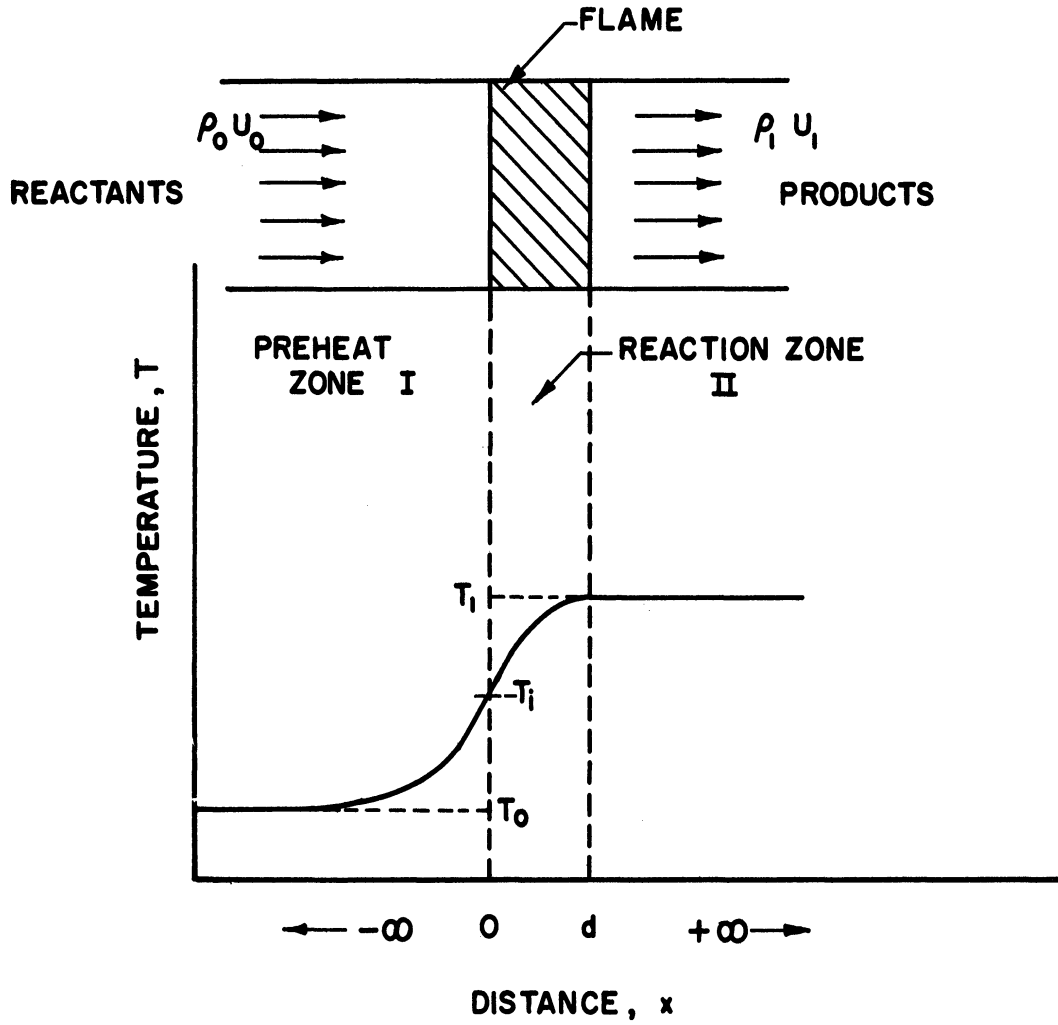


Figure 7. Schematic diagram and temperature profile in the region of the reaction zone.

ture T_0 , to a temperature T_1 at which point the reaction begins. The reaction occurs completely in Zone II, wherein the temperature is raised to T_1 . The width of the reaction zone is 'd'. Beyond Zone II no further reaction occurs.

The notation below is used for the derivation which follows.

Q'	heat of reaction, cal/molecule of reactant
c_p	constant pressure heat capacity, cal/gm-°K
ω	rate of reaction, molecules of reactant/cm ³ -sec
a	concentration of fuel, molecules/cm ³
b	concentration of oxygen, molecules/cm ³
P	steric factor
Z	impact coefficient, number of collisions/cm ³ -sec
d	depth of reaction zone, cm
M	mass rate of flow, gr/cm ² -sec; molecular weight
ρ	density, gr/cm ³
D	diffusion coefficient, cm ² /sec
λ	thermal conductivity, cal/cm-sec-°K

The general equation for a bimolecular second-order reaction is

$$\omega = ab \cdot P \cdot Z \cdot \exp(-E/RT) \quad (4-3)$$

The following assumptions are made in connection with this theory;

1. Pressure is constant.
2. c_p and λ are constant.
3. $\lambda/c_p = D\rho$ (which is exact for the diffusion between two ideal gases whose molecules have the same size and weight).

4. The total number of molecules is constant.
5. Below some temperature, T_i , which is near T_1 the reaction rate is essentially zero.
6. In Zone II, the convection term in the energy balance equation is small with respect to the conduction and heat generation terms.

The basic equations in Zone II are as follow:

Continuity equation for the fuel

$$D\rho \frac{d^2\alpha}{dx^2} - M \frac{d\alpha}{dx} + \omega = 0 \quad (4-4)$$

$$\alpha = \frac{a_0}{\rho_0} - \frac{a}{\rho}$$

Energy equation

$$\frac{\lambda}{c_p} \frac{d^2\theta}{dx^2} - M \frac{d\theta}{dx} + \omega = 0 \quad (4-5)$$

$$\theta = c_p(T - T_0)/Q'$$

Equation of state

$$\rho/\rho_0 = T_0/T \quad (4-6)$$

Conservation of mass

$$M = \rho U = \rho_0 U_0 = \rho_0 S_\ell \quad (4-7)$$

The boundary conditions for these equations are:

$$\text{For } x = -\infty, \quad \alpha = 0, \quad \theta = 0 \quad (4-8)$$

$$\text{For } x = +\infty, \quad \alpha = \frac{a_0}{\rho_0}, \quad \theta = c_p (T_1 - T_0)/Q' \quad (4-9)$$

In consideration of assumption 3 Equations (4-4) and (4-5) are identical in form and have identical solutions if α and θ coincide over the entire reaction zone. That is

$$c_p T + aQ'/\rho = c_p T_0 + a_0 Q'/\rho_0 = c_p T_1 \quad (4-10)$$

This means that the sum of the thermal and chemical energies per unit mass of mixture is constant in the reaction zone. Since this is approximately true, the differential Equation (4-4) can be replaced with the algebraic Equation (4-10). Only the differential Equation (4-5) must now be solved.

In light of assumption 6 Equation (4-5) can be rewritten

$$\frac{d^2 T}{dx^2} + \frac{\omega Q'}{\lambda} = 0 \quad (4-11)$$

where at $x = 0$, $T = T_i$ and at $x = d$, $T = T_1$. In region I, $\omega = 0$ according to assumption 5 and the energy equation is

$$\frac{\lambda}{c_p} \frac{d^2 T}{dx^2} - M \frac{dT}{dx} = 0 \quad (4-12)$$

where at $x = -\infty$, $T = T_0$ and at $x = 0$, $T = T_i$.

The condition which determines the value of 'M' is that the heat transferred from Zone II to Zone I is equal to that received by Zone I.

That is, at $x = 0$

$$\left(\frac{dT}{dx}\right)_{0,I} = \left(\frac{dT}{dx}\right)_{0,II} \quad (4-13)$$

Solving Equations (4-11) and (4-12) and equating their derivatives according to (4-13), the following equation is obtained

$$S_\ell = \frac{M}{\rho_0} = \sqrt{\frac{2\lambda I}{c_p \rho_0 (T_1 - T_0)}} \quad (4-14)$$

where

$$I = \frac{1}{a_0} \int_{T_0}^{T_1} \omega \, dT \cong \frac{1}{a_0} \int_0^{T_1} \omega \, dT \quad (4-15)$$

Since T_i is assumed to be near T_1 , the reaction rate, Equation (4-3), can be approximated in order to facilitate the integration of Equation (4-15).

The approximate equation is

$$\omega \cong ab \cdot P \cdot Z \cdot \exp(-E/RT_1) \cdot \exp(-\sigma E/RT_1^2) \quad (4-16)$$

where $\sigma = T_1 - T$. From Equation (4-10)

$$\frac{a}{a_0} = \frac{T_0 \sigma}{T(T_1 - T_0)} \quad (4-17)$$

Semenov suggests that for a lean mixture the concentration of O_2 in the reaction zone can be approximated by its value in the combustion products.

This quantity is

$$\frac{b}{b_0} = \frac{T_0}{T} \left[1 - \phi \left(1 - \frac{a}{a_0 \frac{T_0}{T}} \right) \right] \quad (4-18)$$

Dugger and Simon⁽⁵⁸⁾ have performed the integration in an approximate manner. Their results are as follows:

$$\int_0^{T_1} \omega \, dT \cong a_{\text{eff}} b_{\text{eff}}^P \cdot Z \cdot \frac{RT_1^2}{E} \cdot \exp(-E/RT_1) \quad (4-19)$$

where ϕ , the equivalence ratio, is the ratio of the actual fuel-air ratio to the stoichiometric fuel-air ratio. For lean mixtures, $\phi < 1$,

$$a_{\text{eff}} = a_0 \frac{T_0}{T_1} \frac{RT_1^2/E}{T_1 - T_0}$$

$$b_{\text{eff}} = b_0 \frac{T_0}{T_1} \left[1 - \phi \left(1 - \frac{RT_1^2/E}{T_1 - T_0} \right) \right]$$

For rich mixtures, the roles of the a's and b's are reversed

$$a_{\text{eff}} = a_0 \frac{T_0}{T_1} \left[1 - \frac{1}{\phi} \left(1 - \frac{RT_1^2/E}{T_1 - T_0} \right) \right]$$

$$b_{\text{eff}} = b_0 \frac{T_0}{T_1} \frac{RT_1^2/E}{T_1 - T_0}$$

The quantity 'Z' is proportional to the square root of temperature and is (Reference 50, p. 151)

$$Z = \left(\frac{d_{\text{col},f} + d_{\text{col},O_2}}{2} \right) \sqrt{8 \pi \frac{R}{A} T \left(\frac{M_f + M_{O_2}}{M_f M_{O_2}} \right)}$$

where d_{col} is the effective collision diameter and M is the molecular weight.

Semenov later relaxes some of the restrictions in assumptions 2, 3, and 4; $\frac{\lambda}{c_p}/D\rho$ is allowed to assume any constant value not necessarily unity; $\frac{\lambda}{c_p}$ is allowed to have the value λ_1/c_{p1} in Zone II; and the number of moles of products to the number of reactants can be the ratio n_0/n_1 .

With the above considered, the final equation for the laminar flame propagation velocity of a second-order bimolecular reaction is

$$S_\ell = \sqrt{\frac{2 \lambda_1 \int_0^{T_1} \omega dT}{a_0 \rho_0 c_p (T_1 - T_0)} \left(\frac{n_0}{n_1}\right)^2 \cdot \left(\frac{\lambda}{D c_p \rho}\right)_1^2} \quad (4-21)$$

Because of the assumptions involved in the integration, Semenov concludes that for a bimolecular reaction, Equation (4-21) is valid for values of $RT_1/E \leq 0.1$. However recent investigations have demonstrated the reliability of the equation for values of RT_1/E up to 0.15 (58, 59, 60).

D. Use of the Thermal Theory Equations to Predict Turbulent Flame Propagation Velocity

According to Equation (4-2)

$$S_t = K \cdot S_\ell \quad (4-22)$$

where S_ℓ can be calculated from Equation (4-21). Because of the unknown constant in Equation (4-22), Equation (4-21) can be considerably simplified by factoring out those quantities which are themselves constants and by expressing other quantities as being proportional to the reaction temperature. In using Equation (4-21), Dugger and Simon⁽⁵⁸⁾ suggest that the thermal conductivity be expressed as the following function

$$\lambda = (C_p + 5/4R)\mu/M$$

and the diffusion coefficient as

$$D = 1.336 \mu/\rho$$

where μ is the viscosity and M is the molecular weight. Dugger⁽⁶¹⁾ proposes that the viscosity can be approximated by the following function of temperature:

$$\mu \propto T^{0.67}$$

Then

$$\frac{\lambda}{c_p D \rho} = (1 + 5/4 R/c_p)$$

and

$$\lambda \propto (c_p + 5/4R)/M \cdot T^{0.67}$$

If c_p is approximated by its value at the flame temperature, c_{p1} , then

$$s_l \propto \sqrt{\frac{T_1^{0.67} \int_0^{T_1} \omega dT}{a_0 \rho_0 (T_1 - T_0)} \left(\frac{n_0}{n_1}\right)^2 \left(1 + 5/4 \frac{R}{c_{p1}}\right)^3} \quad (4-23)$$

The remaining quantities in Equation (4-23) can be calculated by the methods of Chapter II or are known from the initial state of the mixture. The final expression for the turbulent flame speed is

$$s_t = K_1 \sqrt{\frac{a_{eff} b_{eff} T_1^{3.17} \exp(-E/RT)}{a_0 \rho_0 (T_1 - T_0)} \cdot \left(\frac{n_0}{n_1}\right)^2 \cdot \left(1 + 5/4 \frac{R}{c_{p1}}\right)^3} \quad (4-24)$$

where K_1 is determined from experimental data as explained in Chapter VI. K_1 is proportional to the level of turbulence in the reaction zone. For a given value of K_1 , Equation (4-24) is valid for predicting the turbulent flame propagation velocity for a situation or variety of situations in which the level of turbulence remains substantially unchanged.

V. HEAT TRANSFER IN A SPARK-IGNITED ENGINE

A. General

There is no question that the heat transferred from the working fluid of an internal combustion engine measurably lowers the thermal efficiency and power output of the engine. Thus it is an important quantity to consider when analytically studying internal combustion engines. Various investigators have attempted to estimate the magnitude of this heat loss. However since the amount of heat transferred depends a great deal on the engine speed, size, combustion chamber geometry, compression ratio, and operating conditions; as well as the coolant type, temperature, and flow rate; there are often large differences between these various estimates.

Lanchester⁽⁶⁴⁾ estimates the heat loss from the hot gases during combustion and expansion to be about 10 percent of the heat combustion of which 40 percent of this is lost during combustion. Janeway⁽⁶⁵⁾ estimates this heat loss to be 16 to 25 percent of the heat of combustion, depending on the speed, of which a fourth or a fifth is lost during combustion. David and Leah⁽²⁰⁾ suggest that the heat loss, expressed as a percentage of the heat of combustion, is $1500/N$ and $7500/N$ during combustion and expansion respectively where N is the engine speed in revolutions per minute. Pye⁽⁶⁶⁾ estimates that during expansion the heat loss is 10 percent and during combustion is 5 percent of the heat of combustion. Ricardo⁽⁶⁷⁾ suggests that 6 percent is transferred during combustion and 7 percent during expansion.

How much reduction of the thermal efficiency and mean effective pressure of an engine this heat loss effects depends on the crank angle at

which it is transferred. Energy lost near the top center piston position is completely lost in so far as any useful work production is concerned. If an engine without any heat losses has a thermal efficiency of 30 percent and then 5 percent of the heat of combustion is lost at top center, the thermal efficiency will be only 25 percent, a drop of 16 percent. The power output will be reduced by a like amount. However the same heat loss occurring later in the cycle, during expansion, effects a lesser reduction in the thermal efficiency and power output since some work has already been done prior to the decrease of internal energy due to the heat transfer.

Taylor⁽⁴⁵⁾ estimates that the reduction in power and efficiency due to heat loss during combustion and expansion is about 10 percent of the equivalent fuel-air cycle power and efficiency. It is generally agreed that the magnitude of the heat transfer during compression prior to combustion is much smaller than that during combustion and expansion. To a first approximation the compression process can be treated as adiabatic.

In addition to the heat transferred directly from the working fluid during compression, combustion, and expansion, additional heat is transferred to the coolant primarily during the exhaust stroke and as a result of piston friction. Because this study is mostly concerned with the analysis of the compression, combustion, and expansion processes, these additional heat quantities are not considered.

The heat transferred from the working fluid to the cylinder surfaces is primarily convective in nature, although some (estimated to be 5 to 10 percent of the total) is lost by radiation. The radiation losses will be discussed later in this chapter. A number of investigators have found the principal heat transfer mechanism to be turbulent forced

convection, an observation which correlates well with the nature of the flame propagation described in Chapter IV. These investigators correlated the average gas side Nusselt number with the Reynolds and Prandtl numbers as is commonly done with forced convection through tubes or over flat plates etc.⁽⁶⁸⁾. This relationship is

$$\frac{hL}{k} = C \left(\frac{GL}{\mu} \right)^a \left(\frac{C_p \mu}{k} \right)^b$$

where

$$\frac{hL}{k} = \text{Nusselt number}$$

$$\frac{GL}{\mu} = \text{Reynolds number}$$

$$\frac{C_p \mu}{k} = \text{Prandtl number}$$

and C, a, and b are experimentally determined constants. Pinkel⁽⁶⁹⁾ developed a method for determining the average heat transfer coefficient in an internal combustion engine which is based upon this forced convection theory. This method with some modification has been used successfully to predict average heat transfer coefficients in many investigations^(69, 70, 71, 72, 73, 74, and 75). Taylor⁽⁴⁵⁾, Chapter 8, describes this theory completely. After examining experimental results from 16 different engines, Taylor and Toong⁽⁷⁵⁾ proposed the following expression

$$\frac{U_e}{k_g} = 10.4 (Re_g)^{0.75} \quad (5-1)$$

U_e is the overall average heat transfer coefficient and Re_g , the gas side Reynolds number, is equal to Gb/μ , where b is the bore and G is the time averaged mass flow into the cylinder divided by the piston area.

While these investigations shed light on the overall heat transfer process, showing that forced convection is the predominate heat transfer mechanism, they yield no information about the instantaneous heat transfer rates or how the convective heat transfer coefficient depends upon the state of working fluid.

Analytical solutions which consider the multi-dimensional nature of the problem as well as the variation of the heat transfer coefficient can not be obtained. However an analytical solution based on a simplified one dimensional model with a constant heat transfer coefficient can be obtained and does yield a great deal of information about the nature of the forced convection.

B. Convective Heat Transfer in Engines

1. Temperature Distribution and Heat Flow in a Finite Plate in Contact With a Uniform Medium Whose Temperature Varies Periodically

To a first approximation, the working fluid in an internal combustion engine can be assumed to be contained in a hollow cylinder capped at both ends by flat plates. Dahl⁽⁷⁸⁾ has shown that in considering the temperature distribution in a hollow cylinder, the curvature may be neglected for many practical applications. If the ratio of the outside to the inside cylinder radius, r_o/r_i , is less than 1.4, the error in neglecting the curvature is approximately $25(r_o/r_i - 1)$ percent. For the internal combustion engine, where the radius ratio is only slightly greater than one, the error in neglecting the curvature will probably not exceed 3 percent.

Thus the one dimensional flat plate analysis is equally valid for any combustion chamber surface which is either thin walled or has little curvature.

Figure 8 shows the model upon which the heat transfer analysis is based. It has the following characteristics. The wall temperature on the coolant side, T_c , is constant. This approximation has been born out by experimental studies. The gas temperature varies about its mean value, T_0 , cosinusoidally. For this one dimensional situation, the equation for heat conduction in the solid wall is⁽⁷⁸⁾

$$\frac{d^2T}{dx^2} = \frac{1}{\alpha} \frac{dT}{dt} \quad (5-2)$$

where $\alpha = c_p \rho / k$. This quantity is approximately constant over the range of temperatures experienced by the wall and is assumed to be constant for the solution. The gas temperature is presumed to vary as

$$T_g = T_0 + \theta \cos(\omega t) \quad (5-3)$$

The solution of this problem for large values of time is composed of the sum of two solutions, the steady state and the steady periodic. That is,

$$T(x,t) = T_s(x) + T_p(x,t) \quad (5-4)$$

The boundary conditions are

$$T(d,t) = T_c$$
$$-k \frac{dT(0,t)}{dx} = h \left[T_0 + \theta \cos(\omega t) - T_s(0) - T_p(0,t) \right] \quad (5-5)$$

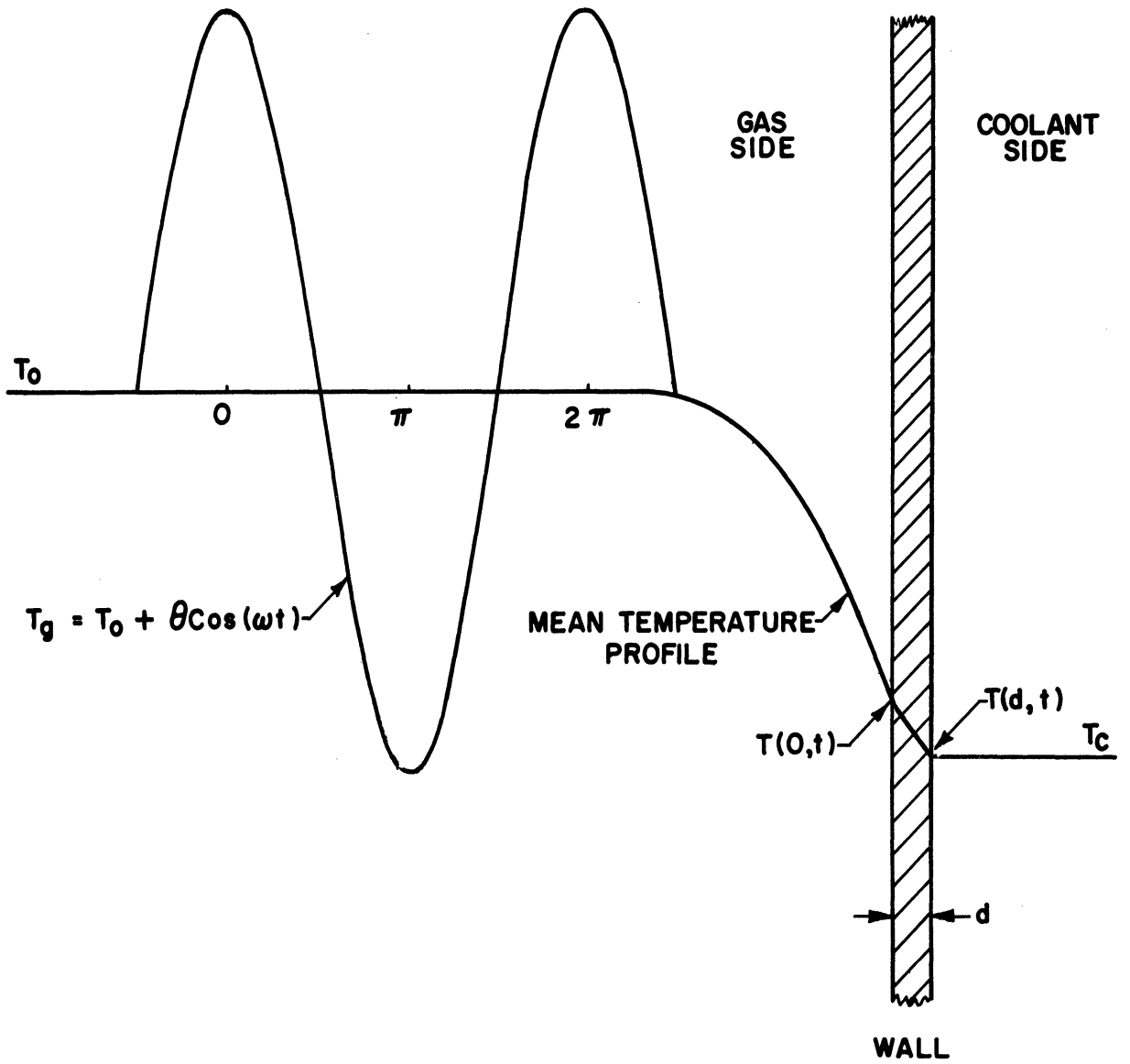


Figure 8. Heat transfer model for the internal combustion engine.

By considering the solution at large values of time, the transient starting solution is eliminated as well as one boundary condition, leaving the two boundary conditions (5-5). To obtain the solution (5-4), the boundary conditions(5-5) can be broken up into two pairs, one for each solution.

For the steady state solution, the differential equation is

$$\frac{d^2 T_s}{dx^2} = 0 \quad (5-6)$$

and the boundary conditions are

$$\begin{aligned} T_s(d) &= T_c \\ -k \frac{dT_s(0)}{dx} &= h [T_0 - T_s(0)] \end{aligned} \quad (5-7)$$

The solution of (5-6) with these boundary conditions is easily obtained as

$$T_s(x) = T_c + [T_0 - T_c] \frac{h(d-x)/k}{hd/k + 1} \quad (5-8)$$

For the periodic part, the differential equation is

$$\frac{d^2 T_p}{dx^2} = \frac{1}{\alpha} \frac{dT_p}{dt} \quad (5-9)$$

and the boundary conditions are

$$\begin{aligned} T_p(\infty, t) &= 0 \\ -k \frac{dT_p(0, t)}{dx} &= h [\Theta \cos(\omega t) - T_p(0, t)] \end{aligned} \quad (5-10)$$

The first boundary condition is approximate only. However it will be seen later that the temperature fluctuations in the wall are damped out so quickly that the wall need not be very thick to approximately satisfy this boundary condition. The solution of this equation, according to Jakob⁽⁷⁹⁾ is

$$T_p(x,t) = \theta \eta e^{-mx} \cos(\omega t - mx - \epsilon)$$

where

$$\eta = \sqrt{\frac{1}{1 + 2m/b + 2m^2/b^2}} \quad (5-11)$$

$$\epsilon = \tan^{-1} \frac{1}{1 + b/m}$$

and

$$m = \sqrt{\frac{\omega}{2\alpha}}$$

$$b = h/k$$

The complete solution is

$$T(x,t) = T_c + \left[T_0 - T_c \right] \frac{b(d-x)}{bd + 1} + \theta \eta e^{-mx} \cos(\omega t - mx - \epsilon) \quad (5-12)$$

Jakob⁽⁷⁹⁾ presents a table of η and ϵ for various values of b/m .

By substituting some typical numerical values for iron in Equation (5-12), the temperature distribution in an engine wall can be approximated. For iron, $\alpha = 0.5 \text{ ft}^2/\text{hr}$ and $k = 36 \text{ Btu/hr-ft}^2\text{-}^\circ\text{F}$ ⁽⁸⁰⁾. A typical engine wall thickness is 0.25 in and speed is 2000 rpm or 4000π radians per min. The gas side heat transfer coefficient varies a good deal, but during the early portion of the expansion stroke, a value of $360 \text{ Btu/hr-ft}^2\text{-}^\circ\text{F}$ is

representative⁽⁷⁷⁾. Thus $b = 10/\text{ft}$, $m = 700/\text{ft}$, $\epsilon \cong \pi/4$, and $\eta = 0.01$.

The average coolant side surface temperature is constant at a given point on the wall and 220°F is a reasonable value when the coolant is water⁽⁷⁷⁾.

If θ is 2000°F and T_0 is 2060°F , Equation (5-12), with the above numerical substitutions, is

$$T(x,t) = 220 + 1840 \cdot \frac{(0.208 - 10x)}{1.208} + 20e^{-700x} \cos(4000\pi t - 700x - \pi/4)^\circ\text{F}$$

A number of conclusions can be drawn immediately:

1. At the wall ($x = d = 0.25/12$ ft) the boundary condition is satisfied. The temperature fluctuations are reduced by a factor of $e^{+14.5}$ which is approximately 10^{+8} . Thus treating the wall as a very thick plate for the purposes of obtaining the periodic part of the solution is a good assumption.
2. At the wall ($x = 0$) the fluctuations are reduced to 1 percent of those in the gas or to 20°F for the present example. This magnitude is in excellent agreement with experimentally measured temperature fluctuations reported in References 77 and 81.
3. At the wall ($x=0$) the fluctuations lag those in the gas by 45° . Because of the great simplification in the gas temperature fluctuation made for the derivation of (5-12), the 45° figure is too large. Although this lag is always a positive quantity it is considerably less than 45° . The data of Oguri⁽⁸²⁾ indicates that half this value may be a better estimate.

4. The average surface temperature ($x = 0$) is about 540°F . This is much too high and reflects the fact that the temperature fluctuations are not cosinusoidal in nature. The relatively low temperatures during the exhaust and induction processes greatly reduce the average surface temperature appreciably. For a water side wall temperature of 220°F , Reference 77 reports a gas side wall temperature of 300°F . This value varies somewhat from point to point on the wall.

Thus while Equation (5-12) must be recognized as being very approximate, it yields wall temperature distributions which at least reflect the more complex processes occurring within the cylinder. It shows that the average wall temperatures are much lower than the average gas temperatures and that the temperature fluctuations in the wall are a miniature replica of those in the gas.

The heat transferred to the wall per unit area is

$$q/A = -k \frac{dT(0,t)}{dx} = k \left[T_0 - T_c \right] \frac{b}{bd+1} + 2m\theta\eta \cos(\omega t - \epsilon - \pi/4) \quad (5-13)$$

This equation shows that the heat transfer is composed of a steady part plus a periodically varying part which is positive as often as it is negative. Substituting numerical values

$$\begin{aligned} q/A &= 36 \cdot \left[15250 + 19750 \cos(\omega t) \right] \\ &= 350 \text{Btu/ft}^2\text{-sec} \quad (\text{maximum}) \end{aligned}$$

A number of conclusions can be drawn immediately:

1. The periodic term is of the same magnitude as the steady state term and during periods of low temperature some heat is transferred from the wall back to the working fluid (for example during the compression process). Thus the wall acts as a sort of flywheel in this respect.
2. For the cosinusoidal gas temperature variation assumed, the heat transfer fluctuations are in phase with the temperature fluctuations of the gas. However because of the many simplifications introduced into the solution this phase relationship is not exactly correct and in fact the gas temperature fluctuations will lead the heat transfer fluctuations by a few degrees.
3. The heat transfer rate calculated in the numerical example, 350 Btu/ft²-sec, compares favorably with the magnitude of the rates measured experimentally in Reference 77 for comparable engine operating conditions.

A more complete solution of (5-2) can be made if it is assumed that the gas temperature varies as a Fourier sine and cosine series as follows

$$T_g = T_0 + \sum_{n=0}^{\infty} \left[A_n \cos(n\omega t) + B_n \sin(n\omega t) \right] \quad (5-14)$$

where T_0 is still the time averaged gas temperature. The complete solution

is

$$T(x,t) = T_c + \left[T_0 - T_c \right] \frac{b(d-x)}{bd + 1} + \sum_{n=0}^{\infty} \eta e^{-m'x} \left[A_n \cos(\omega t - \epsilon - m'x) + B_n \sin(\omega t - \epsilon - m'x) \right] \quad (5-15)$$

where

$$m' = \sqrt{\frac{\omega n}{2\alpha}}$$

The other quantities remain the same as they were in Equation (5-10) except that m is replaced by m' in every case.

If, in an actual engine, the heat transfer fluctuations and the gas temperature fluctuations are not out of phase too much, which appears to be the case (see Oguri⁽⁸²⁾), then to a first approximation the heat transfer rate can be expressed as

$$q/A = h(T_g - T_w) \quad (5-16)$$

where T_w is the average wall temperature and T_g is the instantaneous gas temperature of that gas in contact with the surface A . Since it has been shown that T_w is approximately constant at a point on the wall even for widely fluctuating gas temperatures, Equation (5-16) should predict the rate of heat transfer reasonably well when the appropriate value of ' h ' is employed in the computation.

2. The Convection Heat Transfer Coefficient

The applicability of the exact solution of the preceding section is limited, in part, due to the wide fluctuation in the value of the heat

transfer coefficient in the working fluid of an internal combustion engine. It may vary by as much as a factor of ten during the cycle. A number of investigators have attempted to express this variation in terms of the temperature, pressure, and density of the working fluid.

Of the several expressions in the literature, two are of special interest. Nusselt⁽⁸³⁾ was among the first to formulate an expression for the heat transfer coefficient in an internal combustion engine. His expression, based on experimental observation, included radiation as well as convection effects. It is

$$h = \underbrace{0.99 \sqrt[3]{P^2 T} (1 + 1.24w)}_{\text{convection}} + \underbrace{0.362 \frac{(T/100)^4 - (T_w/100)^4}{T - T_w}}_{\text{radiation}} \text{ kcal/m}^2\text{-hr-}^\circ\text{K} \quad (5-17)$$

where 'w' is the average piston speed in meters per sec. Eichelberg⁽⁸¹⁾ also deduced a formula from experimental data for the heat transfer coefficient. It is

$$h = 2.1 \sqrt[3]{w} \sqrt{PT} \text{ kcal/m}^2\text{-hr-}^\circ\text{K} \quad (5-18)$$

Although these two equations are somewhat different in form, they have certain features in common. For example, both depend on the temperature and pressure of the working fluid and the engine speed. Moreover since the cycle temperature and pressure vary in a similar manner the square root of PT and the cube root of P²T tend to predict similar variations in the heat transfer coefficient. Oguri⁽⁸²⁾ reports that for 'w' in the range of 4-6 meters per sec., both expressions give comparable values of h over the compression, combustion, and expansion processes. Some recent investiga-

tions have employed the expression of Eichelberg to predict heat transfer coefficients. Vincent and Henein⁽⁸⁵⁾ used Eichelberg's equation to predict thermal loadings of diesel engine cylinder walls. Overbye and Co-workers⁽⁷⁷⁾ determined 'h' for a wide range of engine operating conditions. However not particularly good agreement between experimental and calculated values of 'h' was found. On the other hand Oguri⁽⁸²⁾ reported reasonably good agreement between experimental and calculated (by Eichelberg's equation) values of 'h', at least for the combustion and expansion portions of the cycle. Oguri also presents a dimensionless correlation for the gas side Nusselt number but the expression contains constants which must be determined for each particular engine.

Consequently, at this time it appears as if the formula of Eichelberg is about the best expression available for the convective heat transfer coefficient. In light of this, the equation for the convection heat transfer rate per unit area is presumed to be

$$q/A = 2.1 \sqrt[3]{w} \sqrt[2]{PT} [T - T_w] \text{ kcal/m}^2\text{-hr-}^\circ\text{K} \quad (5-19)$$

This equation will be used to calculate the rate of heat transfer during the combustion and expansion processes. The compression process, prior to combustion, will be treated as reversible and adiabatic.

C. Radiant Heat Transfer in Engines

The radiant component of heat transfer is small compared to the convective component in internal combustion engines exhibiting normal combustion. Nusselt⁽⁸³⁾ found that the radiant heat transfer was about 5 percent or less of the total heat transfer (see Equation (5-17)).

Overbye and Co-workers deduced from their experimental observations that the radiant heat transfer was negligible. Baker and Laserson⁽⁸⁶⁾ found experimentally that the radiant heat transfer was quite low. In a 4.6 to 1 compression ratio engine the radiant heat transfer amounted to only 22 Btu/ft²-sec. at its peak. This would be less than 10 percent of the total heat transfer. However for higher compression ratios it is expected that this percentage would increase. Thus a simple calculation is made below to see if this is the case.

Of the principal constituents present in the products of an internal combustion engine, only CO₂ and H₂O emit any appreciable amount of radiation during normal combustion. McAdams⁽⁶⁸⁾ and Hottel⁽⁸⁷⁾ indicate a procedure by which the amount of radiation from these two constituents may be approximated. Unfortunately the curves from which the estimation is made do not extend into the range of temperatures and pressures found in a modern high compression ratio engine. Thus extensive and often questionable extrapolation is necessary. In Appendix H, a calculation is made of the maximum rate of heat transfer due to radiation by the method of these men. An engine operating on the fuel-air cycle at a compression ratio of 10 to 1 was selected for the calculations. The heat transfer was calculated immediately after the adiabatic combustion at top center in order to obtain the conditions where the radiation would be most severe. It was found that under these conditions the radiant heat transfer was still only 12 percent of the total at most. Because of the idealizations of the fuel-air cycle engine, the percentage is higher than it would be in an actual engine.

Thus it appears that in the operating range of the present day spark-ignited internal combustion engine, the amount of heat transferred by radiation during normal combustion is a small proportion of the total. Because of this, radiation will be neglected in the analytical model for the internal combustion engine being proposed in this study and Equation (5-19) will be employed for all heat transfer calculations.

VI. THE ANALYTICAL MODEL FOR SPARK IGNITED ENGINE CYCLE CALCULATIONS

A. General

The discussion of Chapter I indicates that the fuel-air cycle greatly oversimplifies many internal combustion engine processes in that it neglects fundamental phenomena which significantly affect engine performance. Of these phenomena, two are of particular importance, namely the heat transfer and the flame propagation. The purpose of this chapter is to describe an analytical model for internal combustion engine cycle calculations which includes these phenomena. Their inclusion into the cycle analysis, especially during the compression, combustion, and expansion processes, makes possible the calculation of a stepwise continuous pressure time diagram which shows many important features of the experimental by determined diagram. Moreover, this allows an explicit evaluation of the effects of these two phenomena on the efficiency and power output of the engine. A consideration of such variables as engine speed, spark advance, and combustion chamber geometry further enhances the similarity between the analytical model and the actual engine.

The analytical model employs many features of the fuel-air cycle, especially in its treatment of the working fluid. The charge is presumed to be composed of a mixture of fuel and air which is burned during the cycle. The following assumptions, made in connection with the fuel-air cycle analysis in Chapter III, are also made for the analytical model:

1. There is no chemical change in either fuel or air prior to combustion.
2. All products of combustion are in a state of chemical equilibrium at temperatures in excess of 1500°K and are in a state of frozen equilibrium below that temperature.
3. Induction, compression, and exhaust processes are adiabatic.
4. There are no irreversible fluid flow effects.
5. All valve events occur at top and bottom center.

The thermodynamic properties of the working fluid as well as the composition of the products of combustion are computed by the methods illustrated in Chapter II, just as they were for the fuel-air cycle engine analysis of Chapter III.

It is the combustion and expansion processes wherein the analytical model differs greatly from the fuel-air cycle. Both processes are assumed to occur in a stepwise manner and allowance for heat transfer is made during each. Figure 9 illustrates the pressure volume diagram for the assumed processes of this analytical model.

B. The Cycle Processes

1. Process 1-2: Compression

The working fluid is compressed from Point 1 (bottom center) to Point 2. Point 2 is an arbitrary crank angle where combustion begins. The compression is treated as a reversible and adiabatic process.

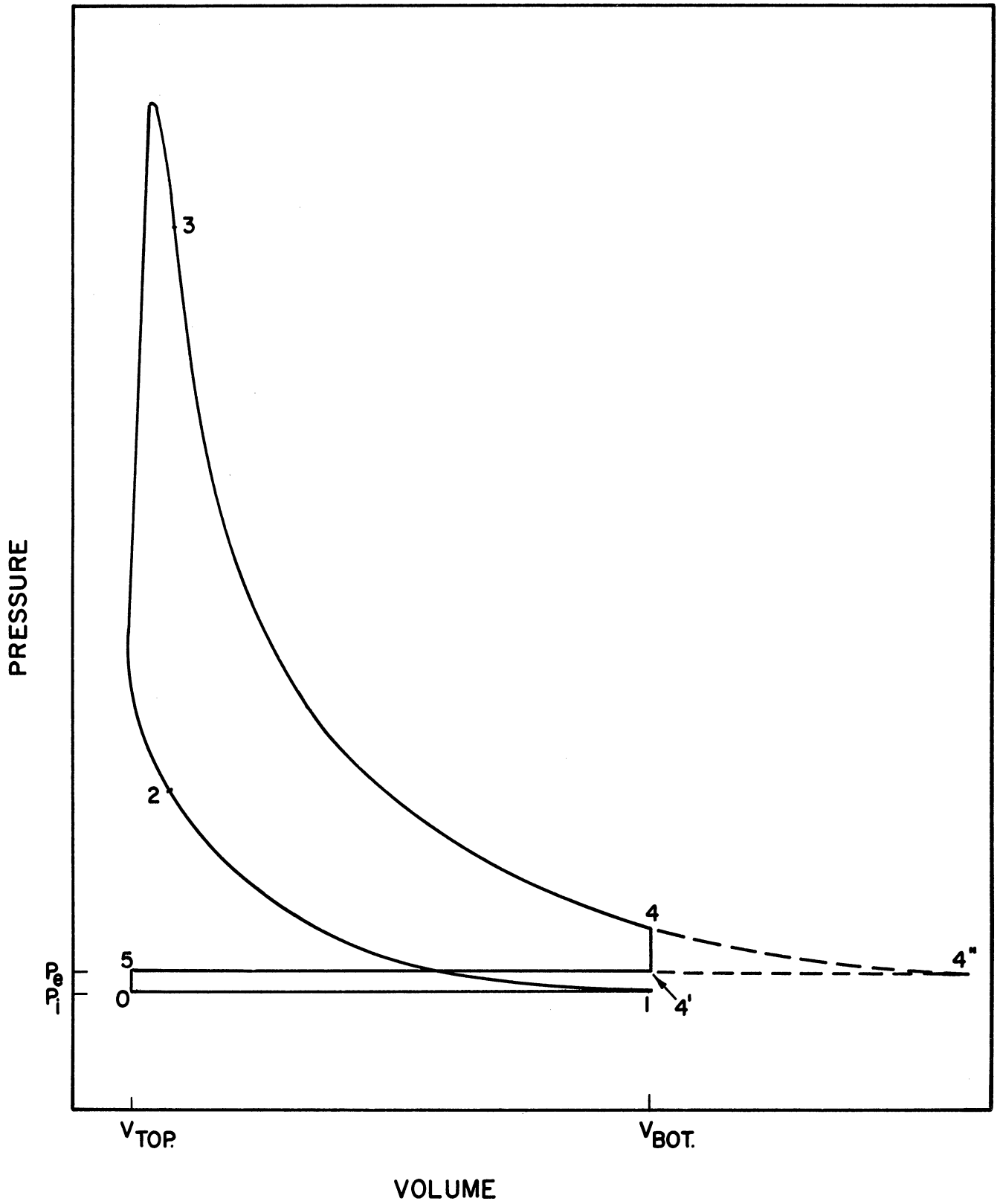


Figure 9. Indicator diagram for the engine on the "analytical model" at part throttle.

Experimental studies have confirmed the general validity of this assumption. They show that during a normal engine cycle little heat is transferred during compression. The state of the system during this process is calculated at regular crank angle intervals in order to construct a pressure time diagram for this part of the cycle. The expression for the volume of the cylinder (including the clearance volume) as a function of crank angle and engine geometry is as follows:

$$V = V_{\text{top}} + \pi \frac{\text{Bore}^2}{4} \left[\frac{\text{Stroke}}{2} (1 + \cos \theta) + \text{Rod} \cdot \left(1 - \sqrt{1 - \left(\frac{\text{Stroke}}{2 \text{rod}} \sin \theta \right)^2} \right) \right]$$

From this expression, the successive volume changes for the isentropic compression processes can be calculated. The total work done during the compression process is

$$W_{1-2} = U_1 - U_2$$

The total heat transferred is zero.

2. Process 2-3: Combustion

The combustion process begins at some crank angle or spark advance corresponding to Point 2. Strictly speaking, this may not be the true spark advance as no provision is made for any ignition delay. Combustion is assumed to progress in a stepwise manner with an arbitrary fraction of the total mass burning during each increment. A single increment is composed of seven steps. During each increment some heat is transferred and some piston motion occurs.

The working fluid during combustion is assumed to be composed of two distinct gaseous regions, one of burned and one of unburned gas.

These are physically separated by a flame front. The unburned gas is compressed continuously during combustion due to the greater volume occupied by the hot products of combustion. Thus each increment that burns does so from slightly different initial conditions. Because of this, a temperature stratification is formed in the burned portion of the gas.

At this stage in the analysis of the combustion process, a number of assumptions must be made, for it is impossible to consider in detail the mixing and heat transfer which may occur within the burned region and between the burned and unburned regions. Within the burned region it must be assumed that either complete uniformity exists because of rapid mixing and heat transfer or that no mixing or heat transfer occur whatsoever, and each burned increment follows an isentropic path subsequent to its combustion. This later assumption was made in an analysis by Hershey.⁽¹⁰⁾ He cites experimental temperature measurements which confirm the stratification although these show it to be less than that calculated. It is not felt that an individual treatment of each burned increment in this manner will lead to significantly different or more meaningful results. Therefore in this analysis complete uniformity will be assumed in the burned region as well as in the unburned region.

It will be assumed however that there is no mixing between the burned and unburned portions. Photographs of normal combustion processes show regular continuous flame fronts and do not indicate that appreciable mixing occurs.

The unburned portion of the gas will be assumed to follow an isentropic path. Pressure and temperature measurements taken in this region by Livengood et al.,⁽⁸⁸⁾ show this to be a reasonably good assumption. This does not mean that there is no heat transfer to or from the unburned gas, but only that the net amount is very small. Thus one final assumption is made for the combustion process, namely, that heat is transferred from the burned to the unburned gas in an amount exactly equal to that transferred from the unburned gas to the combustion chamber surfaces. In effect this means that the burned gas will be charged with the entire internal energy decrease due to heat transfer and the unburned gas will always follow an isentropic path.

Figures 10 a-h illustrates the seven steps comprising the combustion of the i 'th mass increment. To the right of each figure, the total system (burned + unburned) thermodynamic properties are designated by appropriate symbols whenever a single property pertains to both regions. The two homogeneous regions are designated B (burned) and U (unburned). The cross-hatched area represents the currently inflamed mass element and is labeled F (flame element). Let the system pressure, volume, and internal energy at the completion of the $(i-1)$ 'st increment be P_0 , V_0 , and U_0 . Figure 10a illustrates the state of the system in this condition.

Step 1 - Figure 10b - Constant pressure combustion of the i 'th mass increment. A thin shell of gas whose thickness is $(R'_0 - R_0)$ is burned. The system volume increases during this constant pressure

combustion to V_1 . The state of the previously burned and still unburned portions remains unchanged. The flame element undergoes an isenthalpic process, burning to its equilibrium adiabatic flame temperature.

Step 2 - Figure 10c - Constant pressure mixing of the flame element with the previously burned gases. The current flame element is mixed adiabatically with the previously burned portion. The end of this isenthalpic process finds a single homogeneous region of burned gas. The state of the unburned gas remains unchanged.

Step 3 - Figure 10 d - Recompression to volume V_0 . The volume of the system is changed back to V_0 in such a way that the unburned gas follows an isentropic path and the total internal energy of the system after recompression is U_0 . During this step the pressure rises to P_3 . At this point in the process it is as if the system had experienced the combustion of a small increment of mass adiabatically and at constant volume.

Step 4 - Figure 10e - Isentropic change of state due to piston motion. The combustion of the i 'th mass increment has taken a finite amount of time. Call this Δt . During this interval the piston moves causing a change in the volume of the system. Each gaseous region is presumed to undergo an isentropic change of state. The system volume is changed to V_4 and the pressure to P_4 . Δt is figured in Step 7 from a knowledge of the flame speed. An initial assumption is made for Δt . The work done in this step, which is the total work done during the combustion of the i 'th mass increment, is

$$W_i = U_{i0} - U_{i4}$$

Step 5 - Figure 10f - Constant volume heat transfer. Heat is transferred from both regions to the combustion chamber surfaces and from the burned to the unburned gas. The volume of each region remains the same. Because the amount of heat transferred to the unburned gas is just off-set by that from the unburned gas to the walls, the unburned gas undergoes no change of state whatsoever during this step. The burned gas undergoes a constant volume process in which heat is transferred to the walls and to the unburned gas. Thus the internal energy of the gas is reduced by the amount of this heat transfer.

Since heat is transferred from both burned and unburned regions, it is necessary to know, along with Δt , the amount of wall and piston area exposed to both burned and unburned gases. These areas differ with the flame front position and the crank angle and depend on the combustion chamber geometry. During a given combustion increment, subsequent to Step 4, the burned volume occupies a certain fraction of the entire combustion space. Corresponding to this volume fraction there is a certain mean flame front position as well as a definite wall and piston surface area exposed to both burned and unburned regions. By initially assuming that combustion occurs at top center and that the flame front progresses spherically from the center of ignition, these area-length-volume relationships can be found by graphical construction. Such quantities reduced to fractional form are presented in Table VII for the combustion chamber of Figure 1. The proper values can then be found by interpolation in the table (a three point interpolation is used in this work between the length-volume entries and a linear interpolation between the length-area entries).

It is often assumed that a small amount of piston motion does not change these area-length-volume relationships. Taub⁽⁸⁹⁾ reports that the result of detailed calculations of the possible effects of piston motion on these relationships indicates that no appreciable changes occurred for moderate amounts of piston motion, at least for the particular geometry investigated. However, for larger piston displacements their validity is questionable and a more detailed analysis of the combustion chamber may be necessary. This is the case for the combustion chamber of Figure 11. To account for the effect of piston displacement on the position of the burned gas region in this chamber, the following assumptions were made. Prior to top center piston position the relations of Table VIII are presumed valid. After top center some burned gas is found in a right cylinder whose height is just equal to the piston displacement from top center and whose radius is equal to the radius of the flame front. The remaining burned gas occupies the combustion chamber according to the volume-length relationships of Table VIII. Referring to Figure 11, for a small amount of piston displacement the burned gas region would look like a short right cylinder which is shaped like the combustion chamber head on one end and has a hemispherical cap on the other, at least prior to the time when the hemispherical dish is filled with burned gas.

Having determined the position of the flame front by the above assumptions, the wall, head, and piston areas exposed to the burned gas can then be calculated. The latter two are calculated from

a table such as Table VIII and the wall area is calculated from the crank angle as follows:

$$A_{wt} = \pi \cdot \text{Bore} \left[\frac{\text{Stroke}}{2} (1 + \cos\theta) + \text{Rod} \left(1 - \sqrt{1 - \frac{\sin^2\theta \cdot \text{Stroke}^2}{4 \cdot \text{Rod}^2}} \right) \right] \quad (6-2)$$

where the proportion of this exposed to the burned gas can be calculated geometrically. In a similar manner the areas exposed to the unburned gas can be found. For heat transfer calculations the heat and cylinder wall areas are added together.

In Chapter V a decision was made to use the formula of Eichelberg⁽⁸¹⁾, Equation (5-19), to compute the heat transfer. Considering heat transfer from both regions to the walls and piston, the expression for the total heat transfer during the *i*'th combustion increment is:

$$Q_i = 2.1 \sqrt[3]{w} \sqrt{P_4} \left[\sqrt{T_{4b}} (A_{wb} (T_{4b} - T_{\bar{w}}) + A_{pb} (T_{4b} - T_p)) + \sqrt{T_{4u}} (A_{wu} (T_{4u} - T_{\bar{w}}) + A_{pu} (T_u - T_p)) \right]_i \cdot \Delta t \quad (6-3)$$

Equation (6-3) assumes that all the combustion chamber surfaces with the exception of the piston surface are at some constant average temperature and that the piston surface temperature is at some other constant average temperature. In Chapter V, a discussion of the magnitude of the temperature fluctuations at the wall surface indicated that any point on the wall has a relatively constant temperature. However, no two points necessarily have the same temperature. Nevertheless an average overall wall temperature, while a great simplification, would still appear to be a reasonable assumption.

At the end of this step, the burned gas is in a state of chemical equilibrium at volume V_4 . Its internal energy has been decreased during this process by Q_i . The pressure in each of the two regions is different.

Step 6 - Figure 10g - Adiabatic constant energy readjustment to a uniform pressure. During this step the two regions are allowed to attain pressure equilibrium adiabatically, without a change in the total system volume, and therefore at constant internal energy. The unburned gas undergoes an isentropic change of state. The final pressure is P_6 . This step completes the combustion of the i 'th increment. However, the time interval Δt is still not known.

Step 7 - Figure 10h - Determination of the time interval required for combustion. The flame element of Figure 10b is isentropically compressed to the final pressure P_6 . It is assumed that this equilibrium state represents the average burned state for the i 'th increment. The values of T_7 , C_{p7} , and n_7 are substituted into Equation (4-24) as T_1 , C_{p1} , and n_1 . The quantities needed in this equation for the state prior to combustion are those in the unburned gas just prior to Step 1. The flame speed, S_t , is now computed from Equation (4-24) with the following consideration. For air-fuel ratios in the region of five percent lean to five percent rich concentrations a_{eff} and b_{eff} are taken as the average of the concentration at these two extremes. This is because in the derivation of these quantities, it was assumed that the concentration of the fuel was much greater than that of the oxygen or vice-versa. This refinement is suggested by Dugger and Simon. (58)

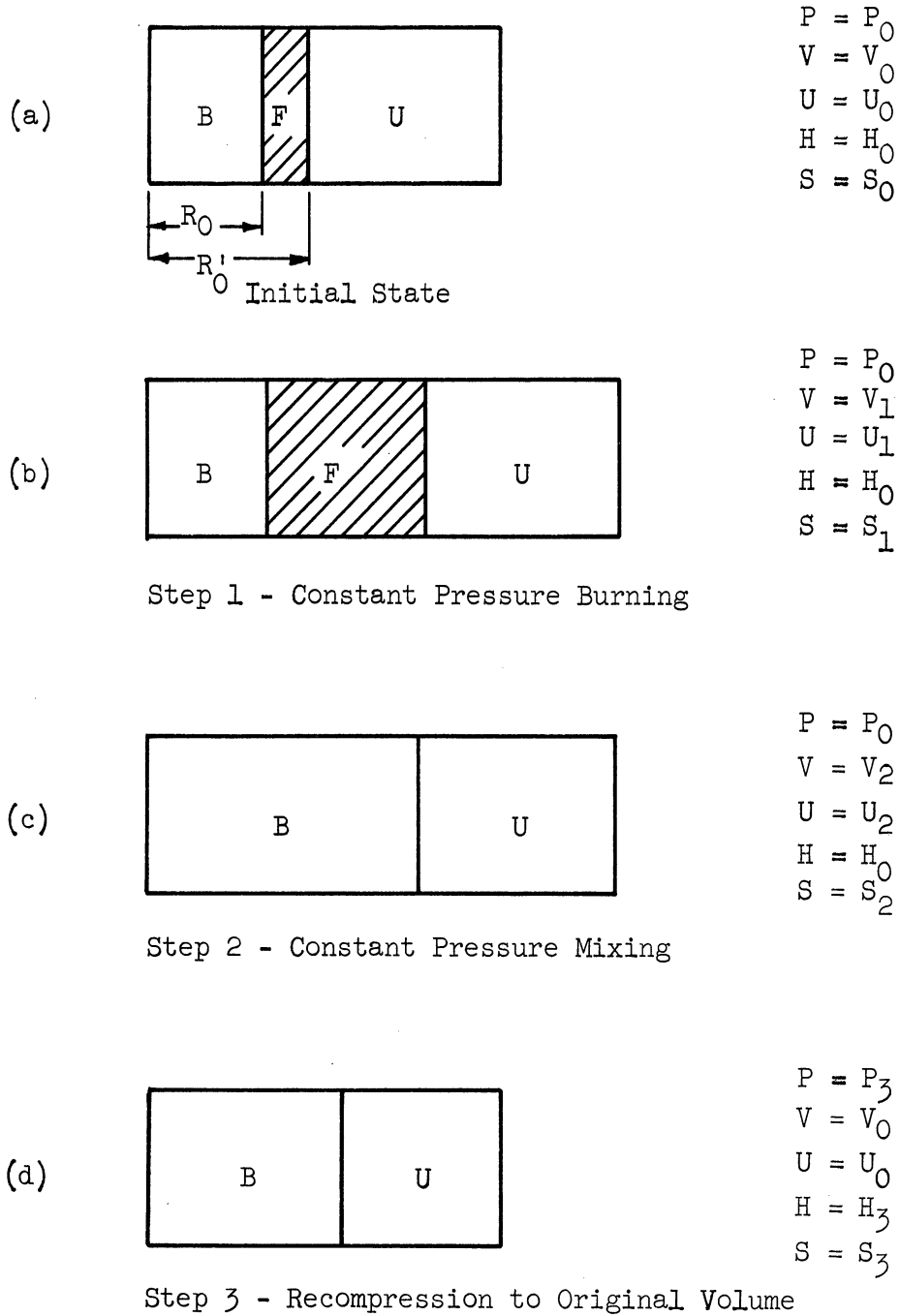
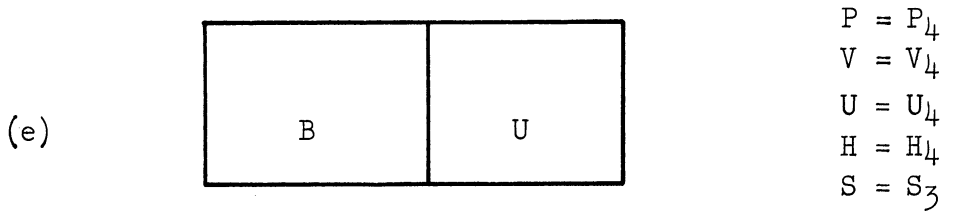
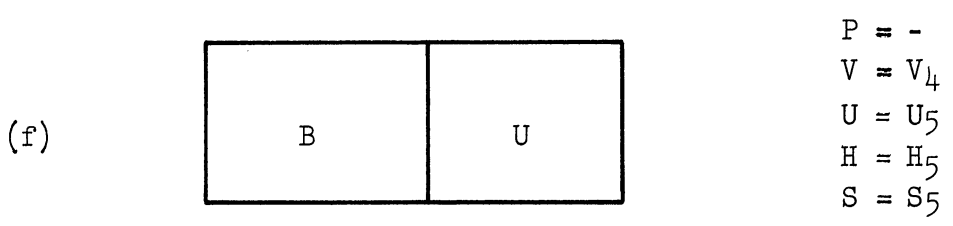


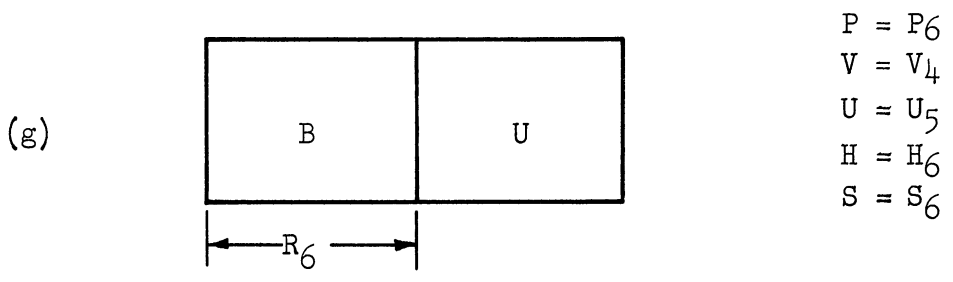
Figure 10. Mathematical Model for Combustion and Heat Transfer.



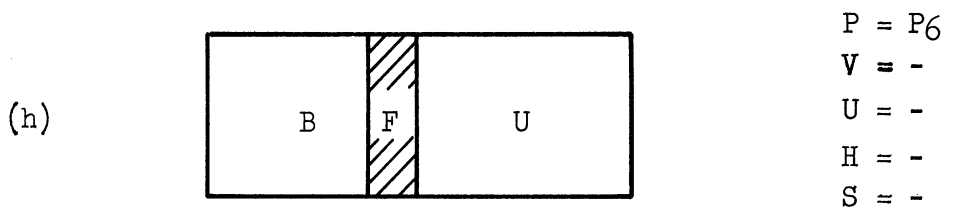
Step 4 - Change of Volume Due to Piston Motion



Step 5 - Constant Volume Heat Transfer



Step 6 - Adiabatic Readjustment to Uniform Pressure



Step 7 - Recompression of Flame Element to Final Pressure

Figure 10. (Continued)

Corresponding to the volume fraction inflamed at the end of the (i-1)'st increment, there was some characteristic distance R_0 , the radius of the spherical flame front. For the combustion chamber of Figure 11, Table VIII includes values of the flame front radius divided by the longest distance the flame travels if the combustion occurs at top center. The effects of small piston displacements on this volume fraction-distance fraction relationship are neglected. At the end of the combustion of the i'th increment, the flame front is at a distance R_G from the center of ignition (see Figure 10g). The difference, $R_G - R_0$ is the total extension of the flame front during the combustion of the i'th increment. Part of this is due to expansion and part to flame propagation. That due to flame propagation is

$$\Delta l = (R_0' - R_0) \quad (6-4)$$

For the first increment this quantity is

$$\Delta l = (3/4 \cdot \pi \cdot V_{f0})^{1/3} \quad (6-5)$$

wherein it is assumed that after the combustion of the first increment the burned gas is spherical in shape. V_{f0} is the volume of the first mass increment prior to combustion.

The time required for the increment to burn is

$$\Delta t = \Delta l / S_t \quad (6-6)$$

If the Δt assumed in Step 4 is not suitably close to that just calculated, Steps 4, 5, 6, and 7 are repeated until the change in successive Δt 's is within a specified limit. When this is the case the i 'th increment is completed and the $(i+1)$ 'st begun. The combustion process is continued until all the mass is consumed. This occurs at Point 3 on Figure 9. For this process the total heat transferred is

$$Q_{2-3} = \sum_{\text{combustion}} Q_i$$

The total work is

$$W_{2-3} = \sum_{\text{combustion}} W_i$$

If the flame speed constant, K_1 , Equation (4-24), is not known; some experimental data must be used to establish its value. The necessary information may be obtained from the experimental pressure-time diagram for the particular engine under consideration. It is necessary to know only the spark advance and the crank angle where the maximum pressure is obtained.

The constant, K_1 , is determined as follows. The incremental combustion process is presumed to begin at the known spark advance. After the combustion of several increments the cylinder pressure will be observed to have reached a peak and then begin to decline. If the crank angle at which this peak occurs is not sufficiently close to the crank angle where the peak occurs experimentally, then the constant can be corrected by a factor which is the ratio of the experimentally determined number of degrees from the beginning of combustion to the

point where the maximum pressure occurs to this same quantity just determined analytically. The combustion process is then repeated and the constant recorrected until the two peaks are suitably close together. When this is the case, the combustion process is then continued. Varying K_1 primarily affects only the crank angle where the peak pressure occurs but indirectly affects the magnitude of the peak pressure in that the cylinder volume is a function of the crank angle.

3. Process 3-4: Expansion

Some expansion has already occurred during the latter stages of the combustion process. However, Process 3-4 can be distinguished in that combustion is already completed and a single homogeneous region of burned gas is present. The expansion process is presumed to occur also in a stepwise manner. A single expansion increment consists of two steps. It corresponds to an incremental change in crank angle $\Delta\theta_e$. It is assumed that prior to the i 'th expansion increment the pressure, temperature, and internal energy are P_0 , T_0 , and U_0 .

Step 1 - Constant volume heat transfer. At volume V_0 heat is transferred from the hot gases, causing a reduction in the internal energy of the gases. The amount of this heat transfer is

$$Q_i = 2.1 \sqrt[3]{V} \sqrt{P_0 T_0} [A_{wt}(T_0 - T_w) + A_{pt}(T_0 - T_p)] \cdot \Delta t \quad (6-7)$$

where the total exposed wall area, A_{wt} , is computed from Equation (6-2) for the current crank angle and $\Delta t = \Delta\theta_e/6/\text{rpm}$. The entropy at the end of Step 1 is S_1 and the internal energy is $U_1 = U_0 - Q_i$.

Step 2 - Isentropic change of state due to piston motion. During the time interval Δt the system undergoes a change of state due to piston motion. The working fluid is presumed to expand isentropically to this new volume. At the end of Step 2, $S_2 = S_1$ and the internal energy is U_2 . The work done during this expansion increment is

$$W_i = U_2 - U_1$$

The expansion process is continued until the piston reaches bottom center, Point 4 on Figure 9. The total heat transferred during this process is

$$Q_{3-4} = \sum_{\text{expansion}} Q_i$$

The total work is

$$W_{3-4} = \sum_{\text{expansion}} W_i$$

4. Process 4-4: Expansion to the Exhaust Pressure

At bottom center the exhaust valve opens and the gases expand to the exhaust pressure. Those which remain in the cylinder are presumed to have undergone an isentropic change of state. This process is identical to that of the fuel-air cycle and is more fully described in Appendix F, Part B, Section 7.

5. Process 4'-5: Exhaust

The piston returns to top center, expelling all the gas except that remaining in the clearance volume. This process is presumed

to be adiabatic. The gases remaining in the clearance volume undergo no change of state. The fraction of gas remaining in the clearance volume is $F = V_5/V_4$. This process is identical to that of the fuel-air cycle and is discussed in Appendix F, Part B, Section 8.

6. Process 5-1: Induction and Mixing

This adiabatic process is the same as that for the fuel-air cycle and is described in Appendix F, Part B, Section 2.

C. The Cycle Characteristics

1. Net Work

The net cycle work is the sum of the work performed during compression, combustion, and expansion plus any pumping work. This quantity is

$$W_t = W_{1-2} + W_{2-3} + W_{3-4} + (V_{\text{bot}} - V_{\text{top}})(P_i - P_e)J \quad (6-8)$$

2. Net Heat Transferred

The net heat transferred is the sum of the heat transferred during combustion and expansion. This quantity is

$$Q_t = Q_{2-3} + Q_{3-4} \quad (6-9)$$

3. Thermal Efficiency

The indicated thermal efficiency is the net work divided by the enthalpy of combustion of the fuel consumed. This is

$$\eta = \frac{W_t \cdot 100}{\Delta H_c^\circ \cdot F_0} \quad (6-10)$$

where ΔH_c° is the enthalpy of combustion per mole of fuel at 298.16°K and 1 atmosphere pressure and F_0 is the moles of fuel inducted per cycle.

4. Mean Effective Pressure

The mean effective pressure is

$$\text{mep} = \frac{W_t}{(V_{\text{bot}} - V_{\text{top}})J} \quad (6-11)$$

5. Volumetric Efficiency

The volumetric efficiency is defined as the quotient of the actual mass of fresh charge inducted to that which would just fill the displaced volume at the inlet density. This quantity is

$$\eta_v = \frac{F_0 \cdot Z \cdot (cx + x) \cdot RT_i \cdot 100}{P_i(V_{\text{bot}} - V_{\text{top}})} \quad (6-12)$$

D. Computational Method

The analytical model has been programmed on an IBM 709 digital computer. The MAD compiler language was used throughout. Appendix I contains a description of the program, a flow diagram, and a listing of the MAD statements. The following chapter reports the results of the computer calculations.

VII. CALCULATED RESULTS FROM THE ANALYTICAL MODEL

A. General

The scope of the calculations presented herein employing the analytical model is designed to accomplish three purposes. The first is to compare the characteristics of a calculated and an experimental cycle for a particular engine. The second is to predict performance characteristics for the engine operating under conditions for which no experimental data are available, assuming the flame speed and heat transfer representation is correct. Finally, the third is to study the availability and irreversibility associated with an engine cycle.

For this work, a single engine was employed. This engine was selected because its combustion chamber geometry was relatively easy to describe mathematically and because its high compression ratio and advanced design were characteristic of current practice. Moreover the high compression ratio would diminish the influence of the exhaust residual whose composition and temperature were not known and could not be calculated accurately by the analytical model. Figure 11 is a sketch of the combustion chamber of this engine. It has a relatively flat head with an off-set hemispherical dish in the piston. Pertinent characteristics of the chamber are listed in Table VII. Table VIII lists its volume-area-length relationships. The compression ratio is 10 to 1. Experimental data for this engine were obtained through Professor Jay A. Bolt of the University of Michigan Department of Mechanical Engineering.

It is known that there will be differences between the calculated and experimental cycle characteristics. These are due to both over-simplifications in the analytical model and differences in operating conditions between the analytical model and the actual engine. Those due to over-simplifications in the model will be discussed later. Differences in operating conditions are due to either a difference in fuel type or to the fact that no experimental value is available for the parameter in question. The experimental data were taken using a blended hydrocarbon of approximate composition $C_{7.35}H_{14.7}$ whereas all calculations were made assuming iso-Octane, C_8H_{18} as a fuel. The thermodynamic properties of the fuel were approximated by those of n-Octane (Tables II and III) and the activation energy was assumed to be 39000 cal/mole as reported in NACA Report 1300.⁽⁵⁰⁾ While it is felt the discrepancies between the analytical and experimental results introduced by this fuel difference are small, no investigation of the effects of fuel type has been conducted.

Certain parameters which are necessary in the calculations were given assumed values because no experimental data were available. These are the wall, piston, and exhaust temperatures. They were given the values 420°K, 500°K, and 1000°K respectively and all calculations reported are made with these values unless otherwise noted.

B. The "Normal Cycle"

A single cycle was selected as a normal cycle against which the resulting analytical model calculations could be compared. Table

IX lists the operating conditions under which the actual engine was run and those for which calculations have been made. For the calculated cycle, the heat transfer was determined directly from Equation (5-19) and the flame speed from Equation (4-24) with K_1 equal to 135,000. This is equivalent to a ratio of turbulent to laminar flame speed of slightly under 4 to 1. The numerical value was arrived at by comparing the experimental with the calculated pressure-time diagram and attempting to achieve good agreement over the early portion of the combustion process. This procedure is explained in Chapter VI. The heat transfer and flame speed calculated in this manner are termed "normal" and the resulting cycle is termed the "normal cycle."

The curve designated as that for normal heat transfer in Figure 12 is the pressure-time curve resulting from the calculations. Figure 13 shows the corresponding pressure-volume diagram. The experimental curves are shown in each case. It must be remembered that these experimental curves represent average values for a large number of cycles and do not necessarily conform to any one single cycle. The complete calculated data for these operating conditions are presented in the first set of data in Data Set A, Appendix K.

C. Comparison of the Analytical and Experimental "Normal Cycles"

1. The Pressure-Time and Pressure-Volume Diagrams

a. The compression process

On both Figures 12 and 13 it is seen that the assumption of a reversible and adiabatic compression process is a close approximation to the real process.

b. The combustion process

1) The ignition delay. No ignition delay was provided for in the calculations. The agreement between the calculated and experimental curves over the early portion of the combustion process indicates that the ignition delay is very small. This observation is in good agreement with the recent findings of Curry⁽⁹⁰⁾ who experimentally found the ignition delay to be very small.

2) First portion of the combustion process. Up to about 8 degrees after top center the calculated pressure-time diagram corresponds closely to the experimental one. At this point about one half of the mass of mixture has been consumed. During this early portion of the combustion process the flame front is passing through the hemispherical dish in the piston but has not yet reached the surface of the piston. Thus, very little heat is transferred as the hot gases are in contact with relatively little wall area. Figure 21 shows the magnitude of the flame speed as a function of crank angle. During this portion of the combustion process the flame speed nearly doubles. The close agreement between the experimental and calculated pressure-time diagrams in this region tends to validate Equation (4-24) for the flame speed.

3) Last portion of the combustion process. At about 8 degrees after top center, the hemispherical dish in the piston becomes completely filled with burned gas. In Figure 12 it is seen that at this point in the cycle the calculated curve abruptly departs from the experimental curve. The analytical model predicts that once the

piston cavity is filled with burned gas, the burning of successive mass increments requires a longer period of time. This is because the flame front must move further in the narrow quench region to burn equal mass increments than it did when the flame was propagating into the hemispherical dish. The experimental curve does not show this.

Because of the many assumptions and simplifications inherent in the analytical model it is difficult to attribute the above deviation to a single factor. Nevertheless some comments are in order. First it must be recognized that the actual flame propagation does not occur spherically as shown in Figure 12 but occurs in such a way that the shape of the flame front tends to reflect the shape of the enclosing combustion chamber. Thus the surface area over which the reaction can occur becomes somewhat larger than that assumed by the analytical model. Moreover, the burned gas tends to move in mass toward a more central position in the chamber (experimentally shown by ionization gap studies) and this causes more surface area over which reactions can proceed than the analytical model assumes. Both these effects would tend to increase the effective burning rate slightly and in particular would cause the hemispherical dish to be completely filled by burned gas slightly later in the cycle. Thus, the inability to describe accurately the shape and position of the burned gas region leads to discrepancies between the calculated and experimental pressure-time diagrams. This is due to a necessary simplification in the analytical model. Second, as the hemispherical dish is filled, the flame front arrives at a sudden change in combustion chamber contour. Rabezanna et al. ⁽¹⁹⁾ have experimentally

studied chambers with sudden changes in contour and have found that in every case there is a sudden appreciable increase in the flame propagation rate whenever such a section change is encountered. This increase which is normally attributed to an increase in turbulence may be as much as 25 or 30 percent of the original flame speed. It is suspected that this latter over-simplification of the combustion process is the major cause of the differences observed between the experimental and calculated pressure-time curves during the last portion of the combustion process.

In consequence of these factors the analytical model predicts a lower peak pressure and longer combustion time than is experimentally observed. The observed peak pressure is 875 psia whereas the calculated is 795 psia. The observed duration of combustion is 45 crank degrees whereas the calculated is 54 degrees.

c. The Expansion Process

During the expansion portion of the cycle, the calculated curve lies above that of the experimental curve. This indicates that its internal energy is greater. The difference between the two curves is due primarily to three factors. First, the analytical model gives less work done by the engine near the top center piston position and consequently more internal energy is present in the gas during the expansion stroke than would be the case if the experimental peak pressure were attained. This is clear from Figure 13 in which the area under the curve is proportional to the work. Second, complete combustion is assumed in the analytical model. In this respect the model represents an optimum condition which is approached but never

attained in practice. This assumption of complete combustion would tend to raise the calculated pressure-time curve higher than the measured one during the expansion stroke. Third, the volumetric efficiency of the engine represented by the calculated curve is slightly greater than that actually measured (97.6 percent verses a measured 94 percent). Thus there is slightly more mass present in the cylinder. Again this would tend to raise the calculated expansion curve higher than the measured curve.

2. Relationship Between Mass Burned and Volume of the Burned Charge

Figure 22 shows the relationship between the mass burned and the volume of the burned charge. Good agreement is seen between the calculated data and the curve presented by Rabezanna et al.⁽¹⁹⁾

3. Heat Transfer

The total heat transferred during the cycle is 17.5 percent of the heat of combustion of which about one third of this is transferred during the combustion process. Although no experimental data are available concerning the actual heat transfer in this engine, these figures agree well with the estimates of Pye⁽⁶⁶⁾, Ricardo⁽⁶⁷⁾, and Janeway⁽⁶⁵⁾. From this and the shape of the pressure-time diagram it appears as if Equation (5-18), the expression proposed by Eichelberg⁽⁸¹⁾, provides a reasonably good estimate of the convective heat transfer coefficient.

4. Rate of Pressure Rise

The calculated pressure-time curve shows good agreement with the experimental curve especially in so far as the rate of pressure rise is concerned. Additional analysis with different combustion chamber

geometries is needed before the analytical model can be used in combustion chamber design.

5. Power and Efficiency

The calculated thermal efficiency of the cycle is 36.5 percent and the measured is 35.8 percent (see Table IX). Most of this difference can be attributed to incomplete combustion of the charge in the actual engine. The differences present in fuel type and volumetric efficiency are believed to affect the thermal efficiency only slightly. If slightly incomplete combustion had been considered and the shape of the pressure volume diagram had remained essentially the same, the calculated thermal efficiency would have been slightly less instead of slightly greater than that measured.

The calculated mean effective pressure is 182 psia whereas that measured was 166 psia. Most of this difference can be attributed to three factors, incomplete combustion, difference in volumetric efficiency, and fuel type.

D. Effect of Exhaust Temperature

Because the analytical model does not treat the exhaust process in detail, the temperature of the exhaust becomes a variable of the problem. For the high compression ratio engine considered in this study the effects of the exhaust residual on the temperatures and pressures of the cycle is slight and thus the temperature and composition of the exhaust residual do not have to be known with great accuracy. However the effect of a 200°K variation in the exhaust temperature was studied.

A single cycle was calculated with all operating conditions identical to those of the normal cycle except for the exhaust temperature. Instead of 1000°K, the exhaust temperature was assumed to be 1200°K. The calculated results for this cycle are presented as Data Set E, Appendix K. This change in exhaust temperature produced no significant changes in any of the cycle calculations.

E. Influence of Heat Transfer and Flame Propagation Rates

This section describes the results of calculations which show the influence of changes in heat transfer and flame propagation rates on the engine cycle characteristics. The "normal" cycle was presumed to be the standard against which the other cycles could be compared. The operating conditions were exactly the same as those for the "normal cycle" with the exception that both the heat transfer coefficient and the flame speed were changed. In particular three different heat transfer rates were considered, no heat transfer, normal heat transfer as defined above, and twice normal heat transfer. Three different flame propagation rates were considered in combination with the three heat transfer rates, a normal rate of flame propagation as defined above, 115 percent of normal, and 85 percent of normal. The results of these calculations are presented in Data Sets A, B, and C of Appendix K.

Figures 12 and 13 are the resulting pressure-time and pressure-volume diagrams for the normal rate of flame propagation. Figures 14 and 15 are the pressure-volume diagrams for the rapid and reduced rates of flame propagation respectively.

Looking at Figure 12, it is seen that the pressure-time curves for the three heat transfer rates become distinguishable near the peak pressure shortly after the hemispherical dish is filled with burned gas. This is the first point in the cycle where burned gas comes into contact with an appreciable wall area. The effects of the three different rates of heat transfer on the pressure-volume diagrams are similar for each of the three flame speeds considered. Increasing the amount of heat transfer causes the expansion curve to fall below that for a lesser amount of heat transfer. In each case the peak pressure changes only slightly.

Figure 16 shows the effect of changing the rate of flame propagation while maintaining a normal rate of heat transfer. It is seen that the pressure-time diagrams and peak pressures are quite different for each different rate of flame propagation.

Figure 17 shows the thermal efficiency and mean effective pressure for these nine sets of operating conditions. The most significant changes in thermal efficiency and mean effective pressure are caused by the changes in heat transfer rate. The changes effected by the differences in flame propagation rate are not large. It must be noted however, that the magnitude of the changes in heat transfer rate is much larger than that of the changes in flame propagation rate. From these results it appears that cycle to cycle variations in the flame propagation rate effect thermal efficiency and power output less than the pressure-time diagram appears to indicate. This is because moderate variations in the pressure-volume diagram do not cause large

variations in either thermal efficiency or mean effective pressure. However, it is apparent that appreciable increases in these quantities can be effected by reducing the heat transfer.

On Figure 17 are indicated the mean effective pressure and thermal efficiency calculated by a fuel-air cycle analysis, the results of which are included in Appendix K, Data Set D and are summarized in Table IX. These idealized values are approached by the analytical model engine with no heat transfer and rapid combustion. In fact the analytical model engine would attain these values of thermal efficiency and mean effective pressure if the combustion rate were greatly increased and the spark advance decreased toward zero degrees. The calculated peak pressure for the cycle with no heat transfer and rapid flame propagation was only about 835 psia whereas that of the fuel air cycle was 1326 psia. Thus a finite combustion time is very beneficial in that it keeps the peak pressures at reasonable levels and yet only slightly lowers thermal efficiency and mean effective pressure.

Table X presents the amount of heat transferred as a percent of the heat of combustion for the two non-zero heat transfer rates and the three flame propagation rates. It is seen that the more rapid the rate of flame propagation, the higher the heat transfer. This is because the highest flame speed cycles were associated with the highest pressures and since the heat transfer coefficient is proportional to the square root of the cylinder pressure the amount of heat transfer increased as the flame speed increased. From this it can be inferred that the highest heat transfer during normal combustion is associated with the most efficient cycles.

Figure 18 is the entropy-volume diagram for the normal rate of flame propagation. Since the compression process is presumed to be reversible and adiabatic no entropy increase occurs. However the combustion process is highly irreversible as it occurs at a finite rate. Thus a large increase of entropy occurs during combustion. For the adiabatic cycle no further entropy change occurs during expansion. However, when heat transfer is taken into account the entropy is seen to decrease during the expansion stroke. This is because heat is transferred from the hot gases and thus the entropy change is negative.

F. Availability and Irreversibility

The availability is the maximum work which can be done by a system when it changes from a given state to a standard state. This maximum work can be realized only through reversible processes including reversible heat transfer to the surroundings. When irreversibilities occur the work realized is less than the maximum. The difference between the maximum and the actual work is termed the irreversibility of the process. The system in question is the closed system comprising the contents of the engine cylinder. For such a system the availability is:

$$\text{Availability} = (U - T_0S) - (U - T_0S_0)$$

where the subscript refers to the standard state.

Because the composition of the system changes due to combustion and dissociation, some difficulty arises in the definition of the standard state. For this work the standard state is taken as the

products of combustion at 1 atmosphere pressure and 298.16°K. The products are presumed to consist only of N_2 , CO_2 , H_2O , and CO for the rich mixture considered here.

The lower curve of Figure 19 shows the availability versus the crank angle for the operating conditions reported in the first set of data of Data Set A, Appendix K. These were conditions of normal heat transfer and normal flame propagation. Under these conditions the mole fractions constituting the products are as follows:

N_2	0.718
CO_2	0.105
CO	0.028
H_2O	0.150

and the total number of moles is 0.02059. The molal properties of these constituents in the standard state are given in Table II. The value of $U_0 - T_0S_0$ is then -702.7 cal.

During the compression portion of the cycle, prior to combustion, the state changes are presumed to be reversible and adiabatic and thus the availability is increased by the amount of work done on the system. During combustion and expansion the availability decreases. If these processes were reversible the expansion work would be equal to the change in availability. However, this is not the case. The combustion process is highly irreversible as the entropy-volume diagram of Figure 18 shows. Both during combustion and expansion additional irreversibility is introduced due to heat transfer. This is shown in Figure 19. Note that subsequent to the completion of the combustion process no additional irreversibility is introduced from this source and further increases are due to heat transfer only.

Figure 20 shows the availability as a function of crank angle for the normal rate of flame propagation with no heat transfer. For this cycle the only irreversibility is due to combustion.

It is interesting to note the large amount of availability existent at the end of the expansion stroke in both Figures 19 and 20. It is a substantial fraction of the original availability. The thermal efficiency of the engine could be greatly increased if some method were found for taking advantage of all or a part of this potential energy.

G. The Analytical Model and the Fuel-Air Cycle

It has already been shown that the fuel-air cycle is the limiting case of the analytical model for rapid rates of flame propagation. In turn it was shown that the air cycle is the limiting case of the fuel-air cycle for extremely lean mixtures.

The number of additional parameters included in the analytical model, not the least of which are the heat transfer and flame propagation, allow a mathematical analysis whose results show close similarity to the experimental data in many respects. Thus the analytical model provides a distinct improvement over the fuel-air cycle for internal combustion engine analysis. It must be pointed out however, that the analytical model calculations must be made on a digital computer whereas those of the fuel-air cycle require no such aid.

Table IX lists the operating conditions and experimental results for the normal cycle, the calculated results from the analytical model, and those from the fuel-air cycle for similar engine operating

conditions. A comparison of the quantities presented indicates the improvement in the cycle analysis made possible by the analytical model.

H. Concluding Remarks

The correlation between the calculated and experimental data was largely satisfactory for the normal cycle considered. The major discrepancies appear to be caused by two factors. The first is the inability to describe accurately the shape and location of the burned gas region. The second and most influential factor is the inability to predict the effect of section changes on turbulent flame propagation rate. In those portions of the cycle where the effects of these two factors are small the heat transfer and flame propagation appear to be described satisfactorily.

Before the analytical model can be employed to evaluate new engine designs or to even aid in the design of combustion chambers, much more work will be needed to verify the calculated results. In particular the influence of combustion chamber geometry, different fuel type, and different engine operating conditions must be studied.

More information is needed to describe the combustion process accurately. The expression employed in this study for the flame propagation rate needs improvement. In particular more information is needed about the turbulence in an engine so that the flame speed can be predicted directly from engine operating conditions without relying on an experimentally determined constant whose value under new operating conditions is highly questionable.

In summary the analytical study of the internal combustion engine must proceed along two paths. First the analytical model must be studied by employing it to calculate engine cycles under widely varying operating conditions. Second, a research program must be undertaken to obtain the following information:

1. The nature of the turbulence in the cylinder of a spark ignited engine.
2. The influence of section change on the turbulent flame propagation rate.
3. The introduction of the preceding information into the analytical model for the prediction of the turbulent flame speed.

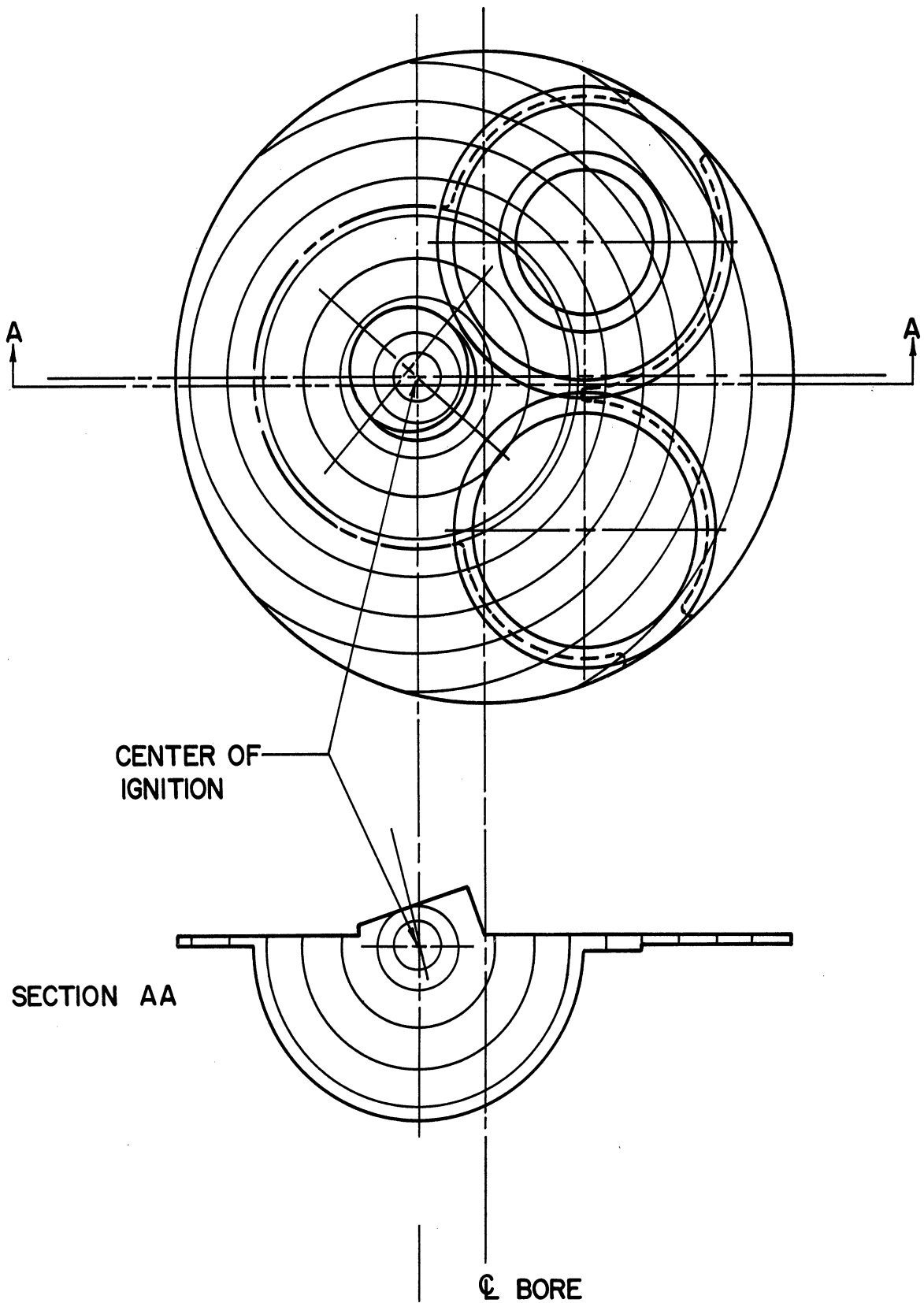


Figure 11. View of engine combustion chamber showing spherical flame propagation pattern.

TABLE VII
PERTINENT DIMENSIONS FOR ENGINE OF FIGURE 11

Dimension	Symbol	Numerical Value	
Bore	Bore	3.94 in	10 cm
Stroke	Stroke	2.46 in	6.25 cm
Connecting Rod	Rod	7.065 in	17.9 cm
Radius of Hemispherical Dish		1.06 in	2.79 cm
Quench Height		0.05 in	1.27 cm
Compression Ratio	r	10 to 1	10 to 1
Clearance Volume	V_{top}	3.33 in ³	54.6 cm ³
Swept Volume	V_{bot}	30.00 in ³	492 cm ³
Head Area	A_h	12.13 in ²	78.4 cm ²
Piston Surface Area	A_p	15.35 in ²	99.0 cm ²
Maximum Flame Travel Length If Combustion Occurs at Top Center	L_t	2.4 in	6.1 cm

TABLE VIII

VOLUME-AREA-LENGTH RELATIONSHIPS FOR THE COMBUSTION CHAMBER OF
FIGURE 11

L/L_t	A_{hb}/A_{ht}	A_{pb}/A_{pt}	V/V_t		V/V_t	L/L_t
0.0	0.0	0.0	0.0	0	0.0	0.0
0.05	0.0	0.0	0.00285	1	0.05	0.150
0.1	0.0	0.0	0.01875	2	0.1	0.196
0.15	0.0335	0.0	0.05353	3	0.15	0.230
0.2	0.0596	0.0	0.10629	4	0.2	0.260
0.25	0.0934	0.0	0.18093	5	0.25	0.282
0.3	0.1341	0.0	0.29034	6	0.3	0.304
0.35	0.1840	0.0	0.43688	7	0.35	0.322
0.4	0.2385	0.0	0.63774	8	0.4	0.340
0.45	0.299	0.449	0.77300	9	0.45	0.354
0.5	0.369	0.504	0.80343	10	0.5	0.368
0.55	0.446	0.566	0.83313	11	0.55	0.380
0.6	0.533	0.634	0.85971	12	0.6	0.392
0.65	0.625	0.706	0.88276	13	0.65	0.403
0.7	0.709	0.770	0.90933	14	0.7	0.414
0.75	0.777	0.825	0.93317	15	0.75	0.430
0.8	0.837	0.874	0.94724	16	0.8	0.490
0.85	0.894	0.917	0.96561	17	0.85	0.580
0.9	0.938	0.954	0.98163	18	0.9	0.686
0.95	0.979	0.985	0.99101	19	0.95	0.804
1.0	1.0	1.0	1.0	20	1.0	1.0

TABLE IX

NORMAL ENGINE CYCLE OPERATING CONDITIONS
(wide open throttle)

Compression Ratio: 10 to 1

Operating Condition	Experimental	Analytical Model	Fuel-Air Cycle
Fuel	Blended	Iso-Octane	Iso-Octane
Air-Fuel Ratio (fraction Theoretical air)	.92	.92	.92
Piston Temperature	Unknown	500°K	----
Wall Temperature	Unknown	420°K	----
Manifold Temperature	320°K	320°K	320°K
Manifold Pressure	.99 atm	.99 atm	.99 atm
Exhaust Temperature	Unknown	1000°K	1240°K
Exhaust Pressure	1.03 atm	1.03 atm	1.03 atm
Spark Advance	16°BTC	16°BTC	----
Speed	2800 rpm	2800 rpm	----
Peak Pressure	875 psia	795 psia	1326 psia
Max. Rate of Pressure Rise	85 psi/deg	85 psi/deg	∞
Duration of Combustion	46 degrees	54 degrees	----
Thermal Efficiency	35.8 percent	36.5 percent	42.7 percent
Mean Effective Pressure	166 psia	182 psia	213 psia
Volumetric Efficiency	94 percent	97 percent	97.5 percent

TABLE X

AMOUNT OF HEAT TRANSFERRED
(Percent of heat of combustion)

Compression Ratio: 10 to 1
Fuel: Iso-Octane
Speed: 2800 rpm
Ignition: 16° BTC

Rate of Flame Propagation	Normal Heat Transfer			Twice Normal Heat Transfer		
	During Combustion	During Expansion	Total	During Combustion	During Expansion	Total
Normal	6.06	11.4	17.5	11.8	17.4	29.2
Rapid	5.71	12.3	18.0	11.2	18.8	30.0
Reduced	6.90	9.5	16.4	13.2	14.4	27.6

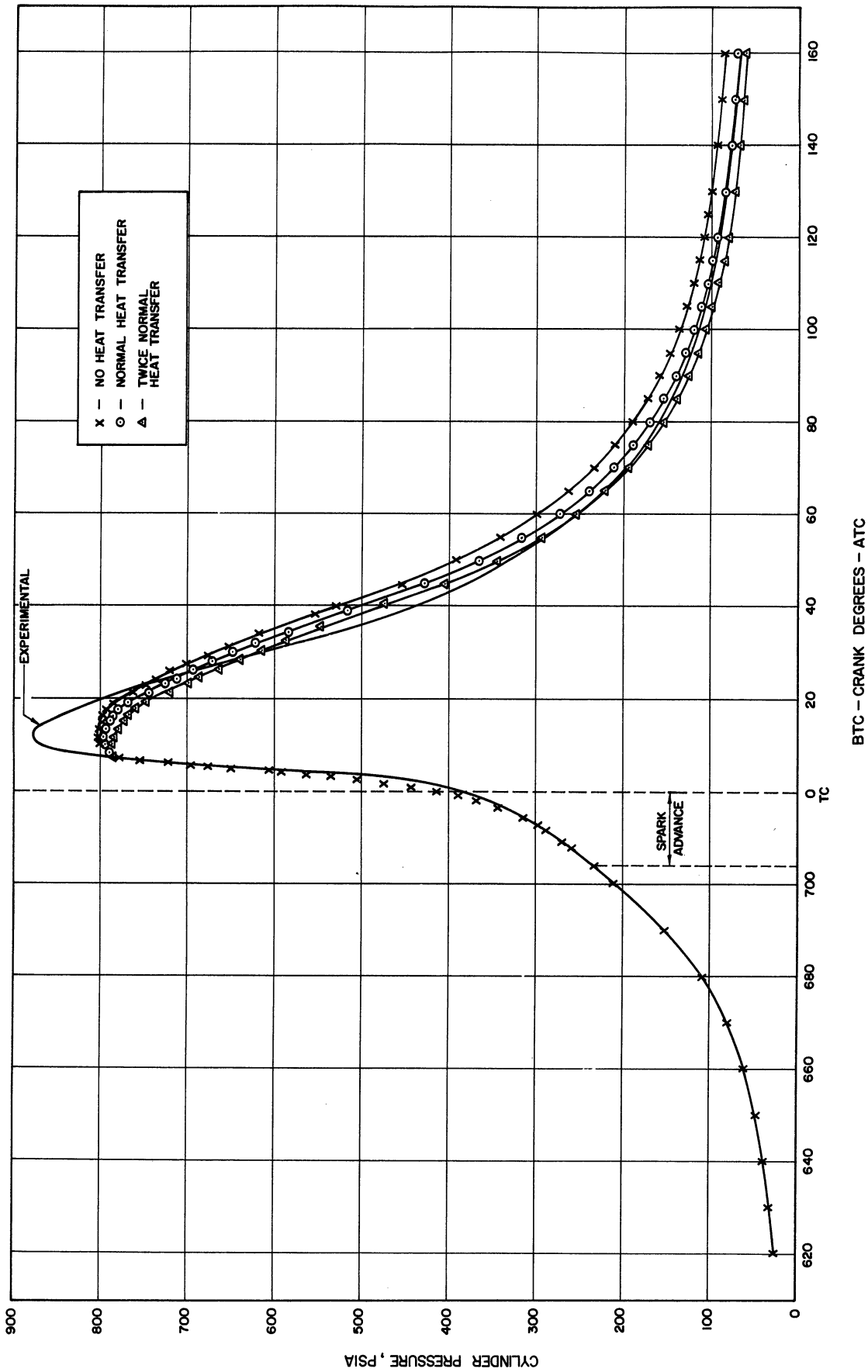


Figure 12. Cylinder pressure vs crank angle for a 10:1 compression ratio engine with normal rate of flame propagation (Data Set A); operating conditions listed in Table IX.

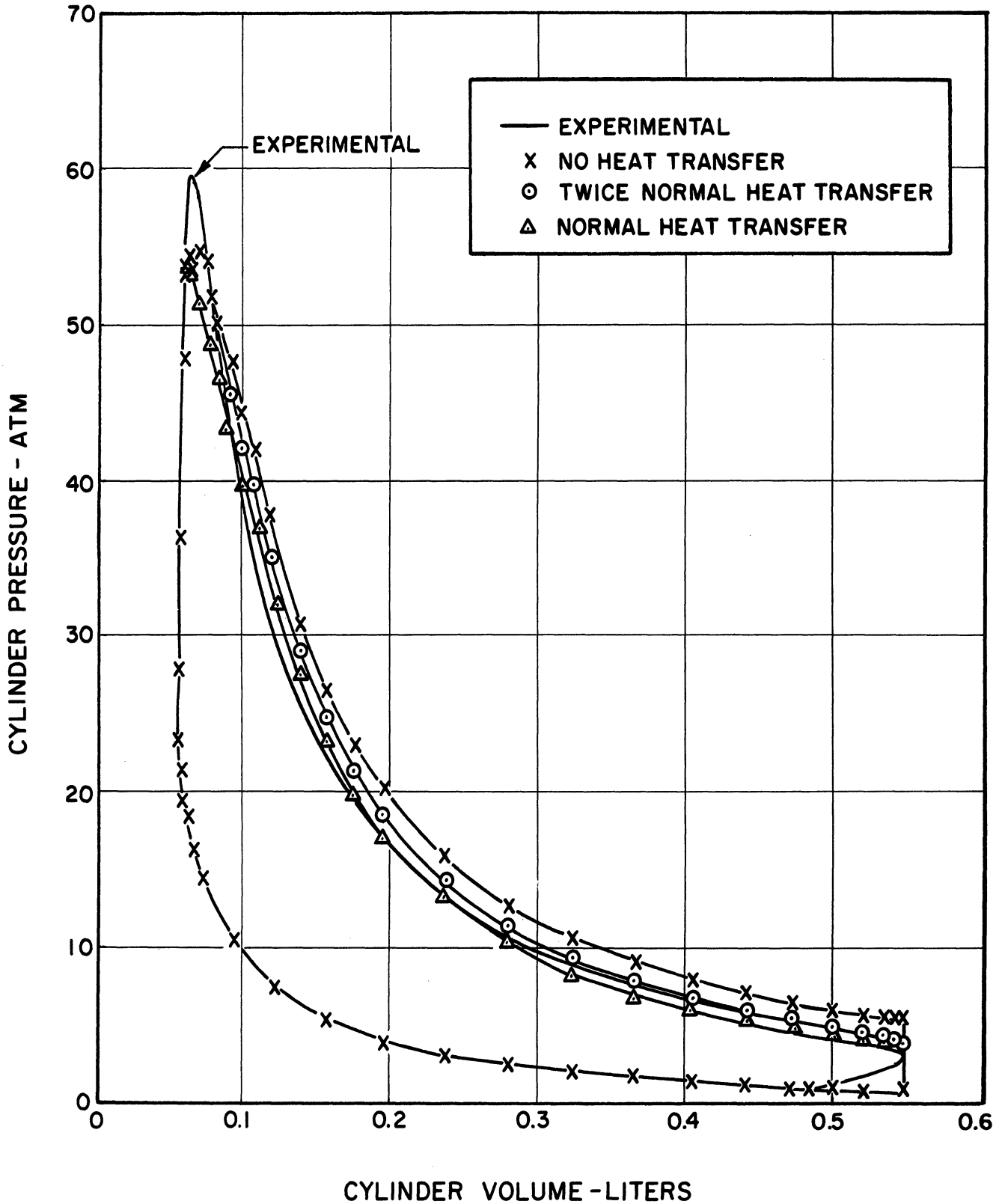


Figure 13. Pressure-volume diagram for a 10:1 compression ratio engine with normal rate of flame propagation (Data Set A). Operating conditions listed in Table IX.

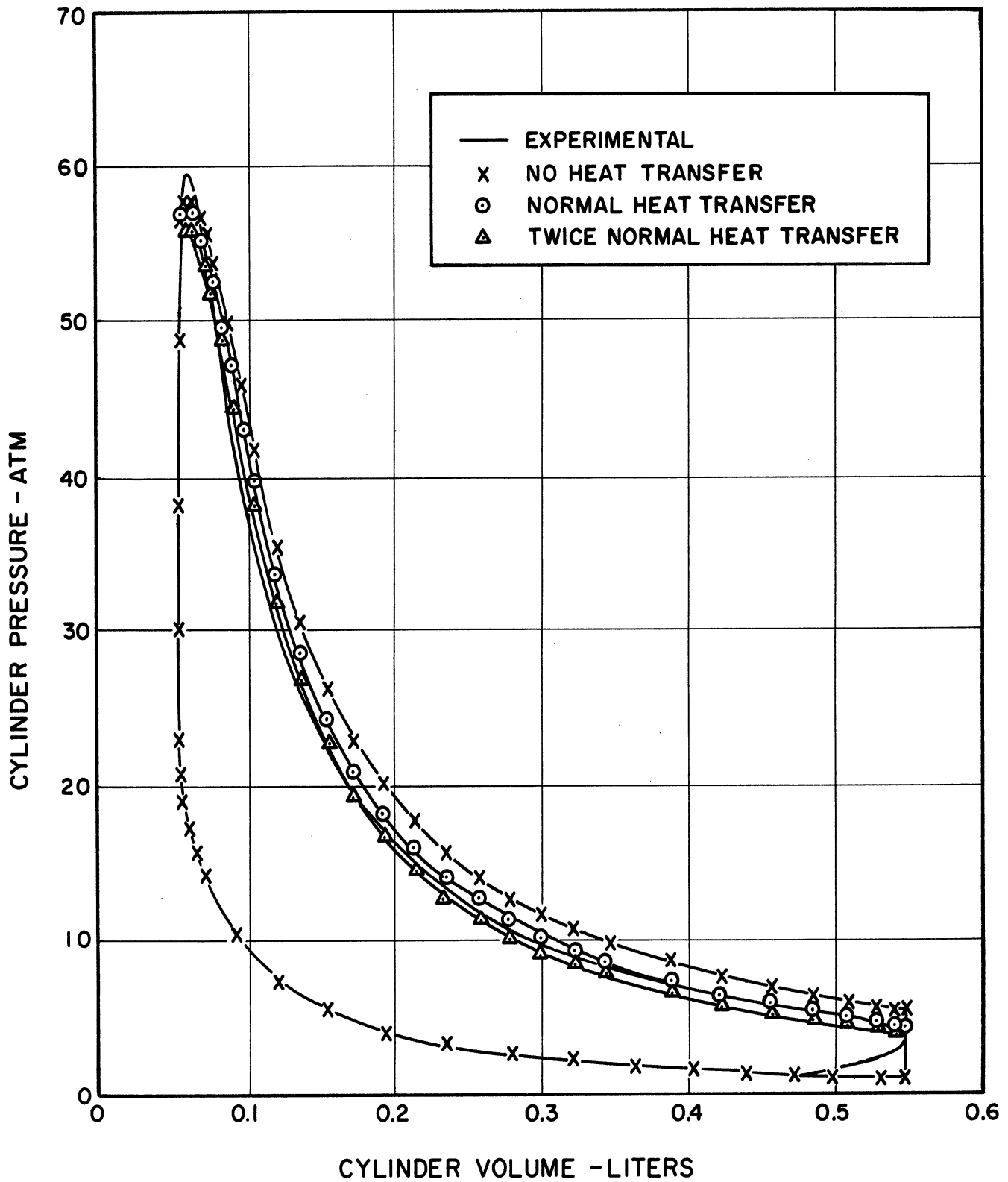


Figure 14. Pressure-volume diagram for a 10:1 compression ratio engine with rapid rate of flame propagation (Data Set B). Operating conditions listed in Table IX.

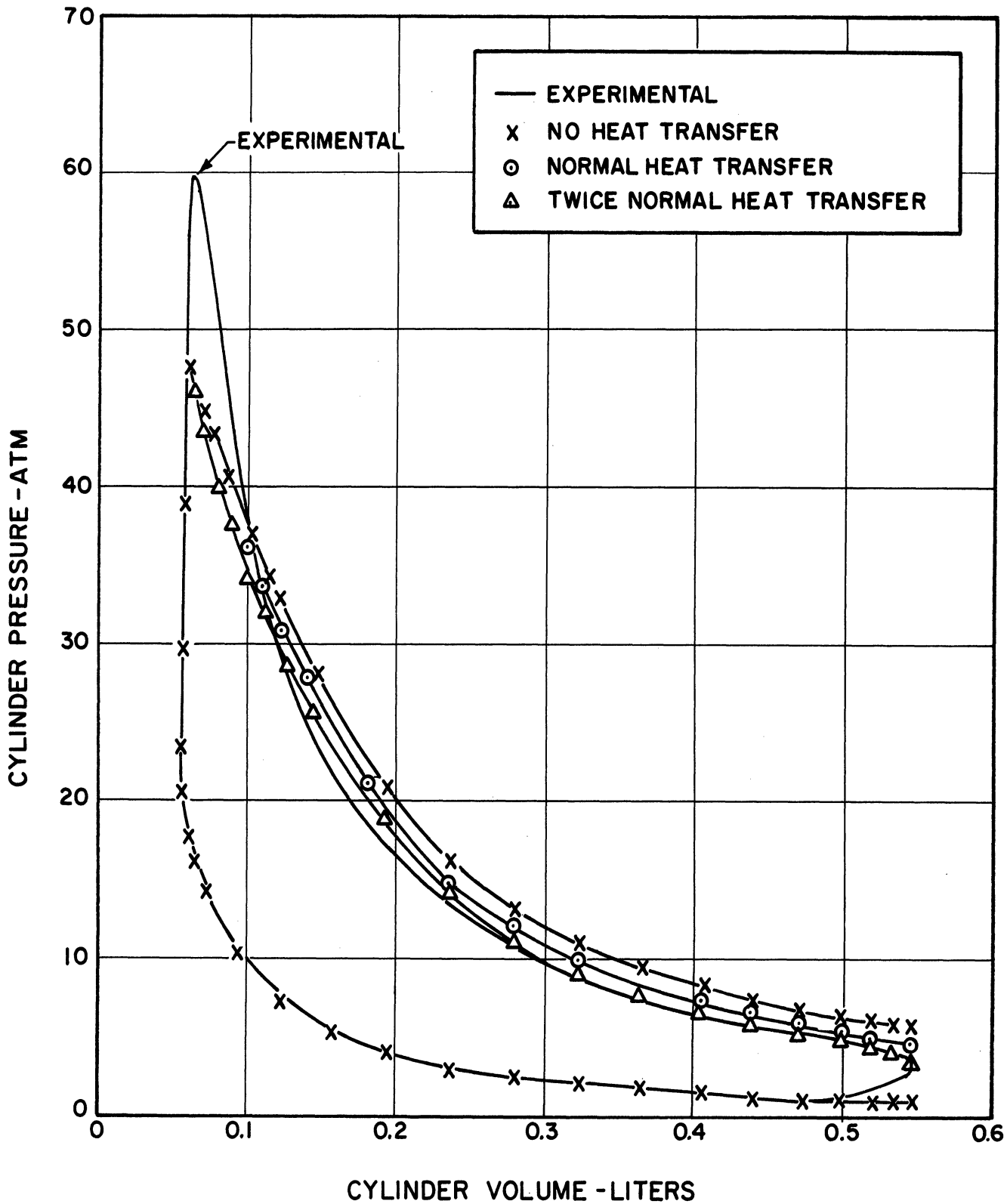


Figure 15. Volume diagram for 10:1 compression ratio engine with reduced rate of flame propagation (Data Set C). Operating conditions listed in Table IX.

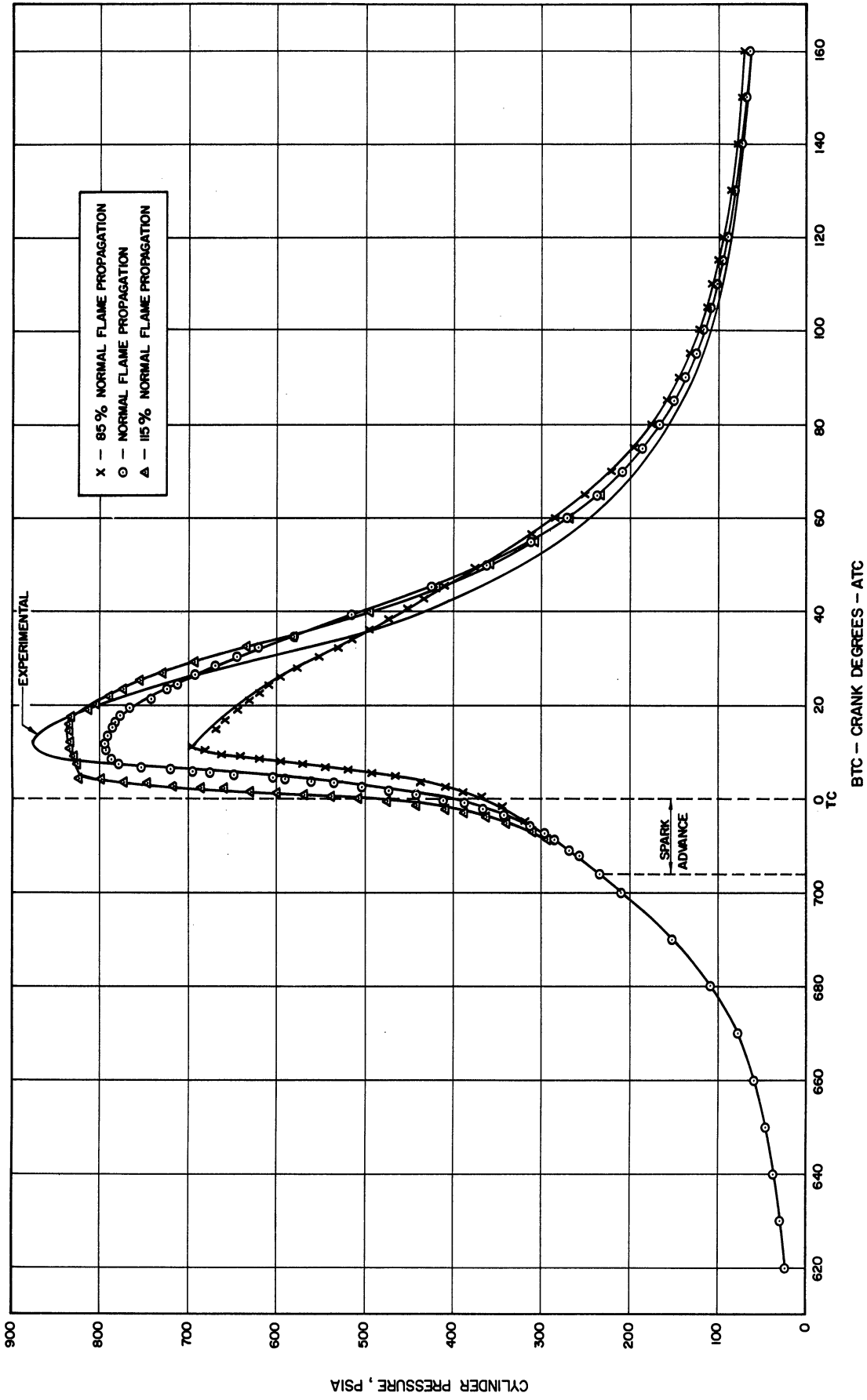


Figure 16. Cylinder pressure vs crank angle for a 10:1 compression ratio engine with normal rate of heat transfer. Operating conditions listed in Table IX.

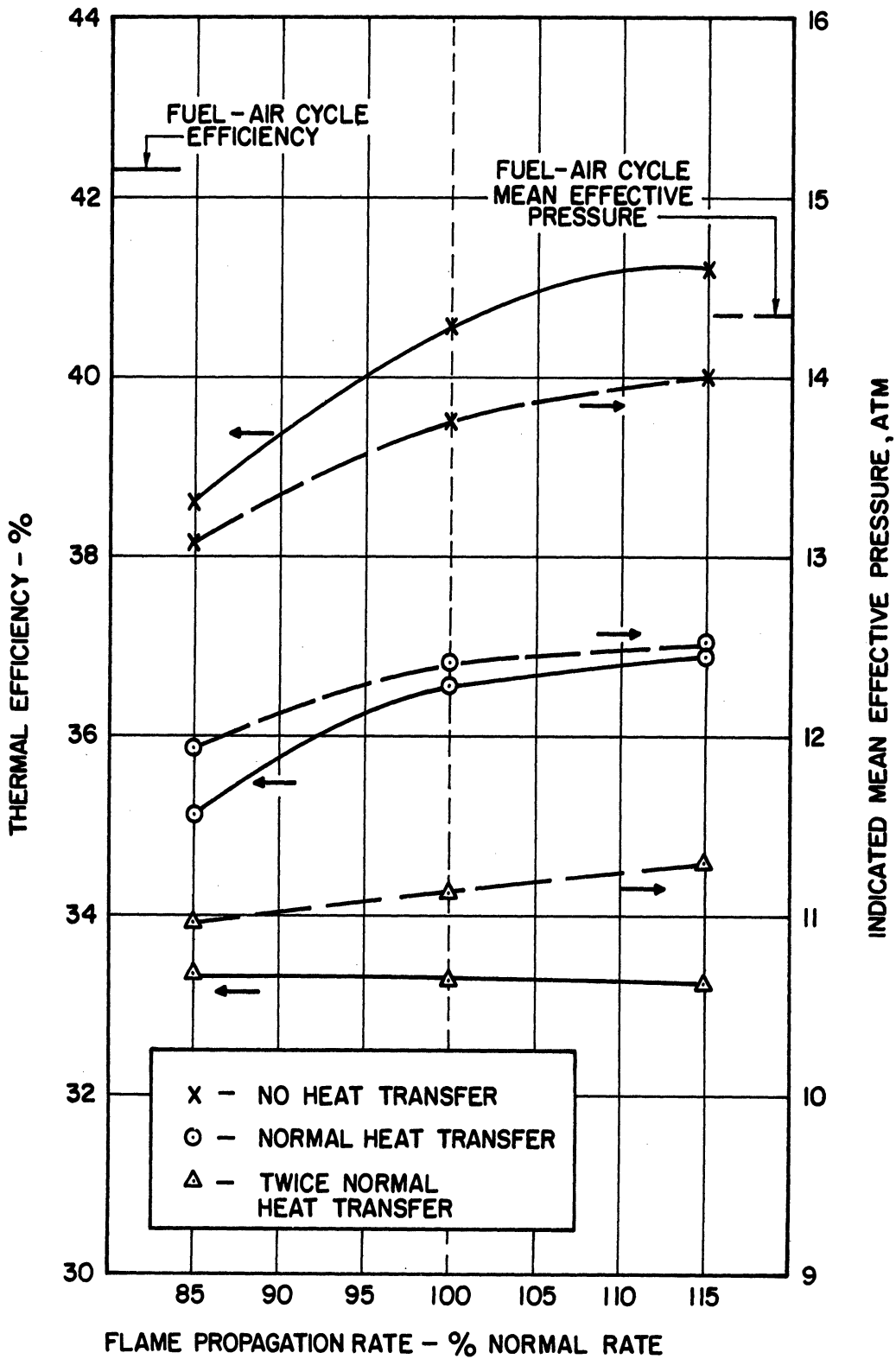


Figure 17. Calculated thermal efficiency and mean effective pressure vs flame propagation rate for a 10:1 compression ratio engine. Operating conditions listed in Table IX.

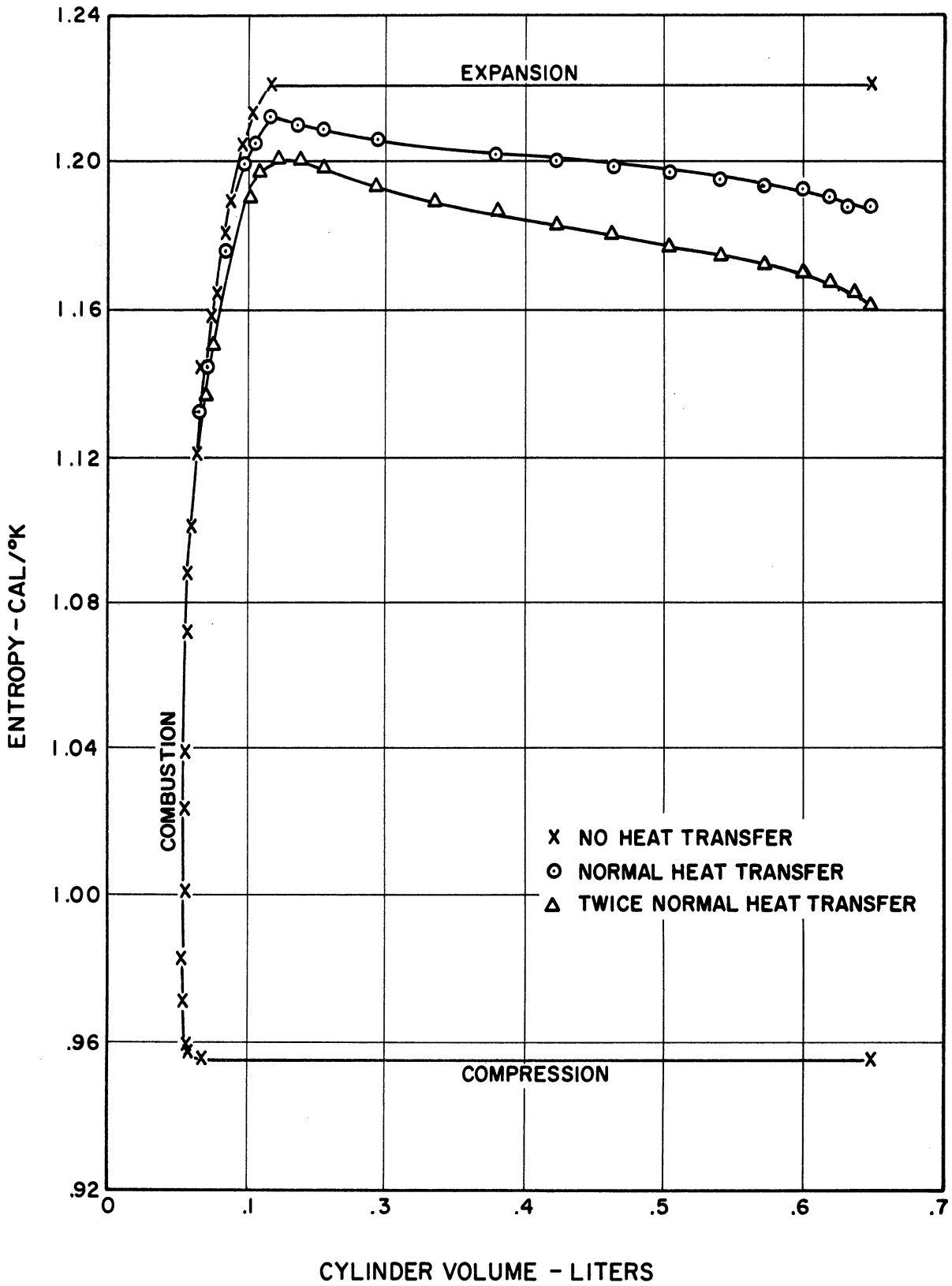


Figure 18. Entropy-volume diagram for a 10:1 compression ratio engine with normal rate of flame propagation (Data Set A).

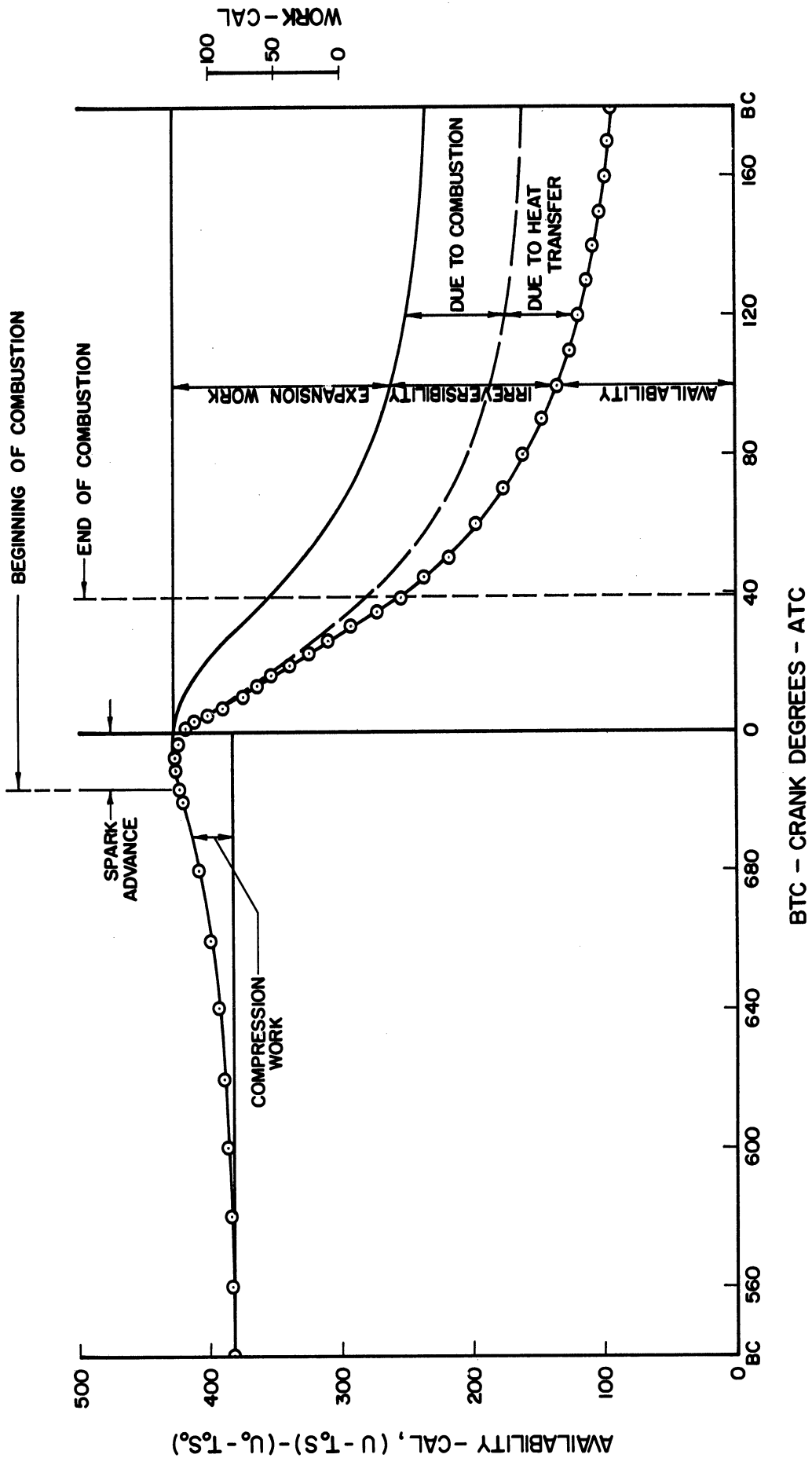


Figure 19. Availability vs crank angle for a 10:1 compression ratio engine with normal rate of flame propagation and heat transfer (Data Set A).

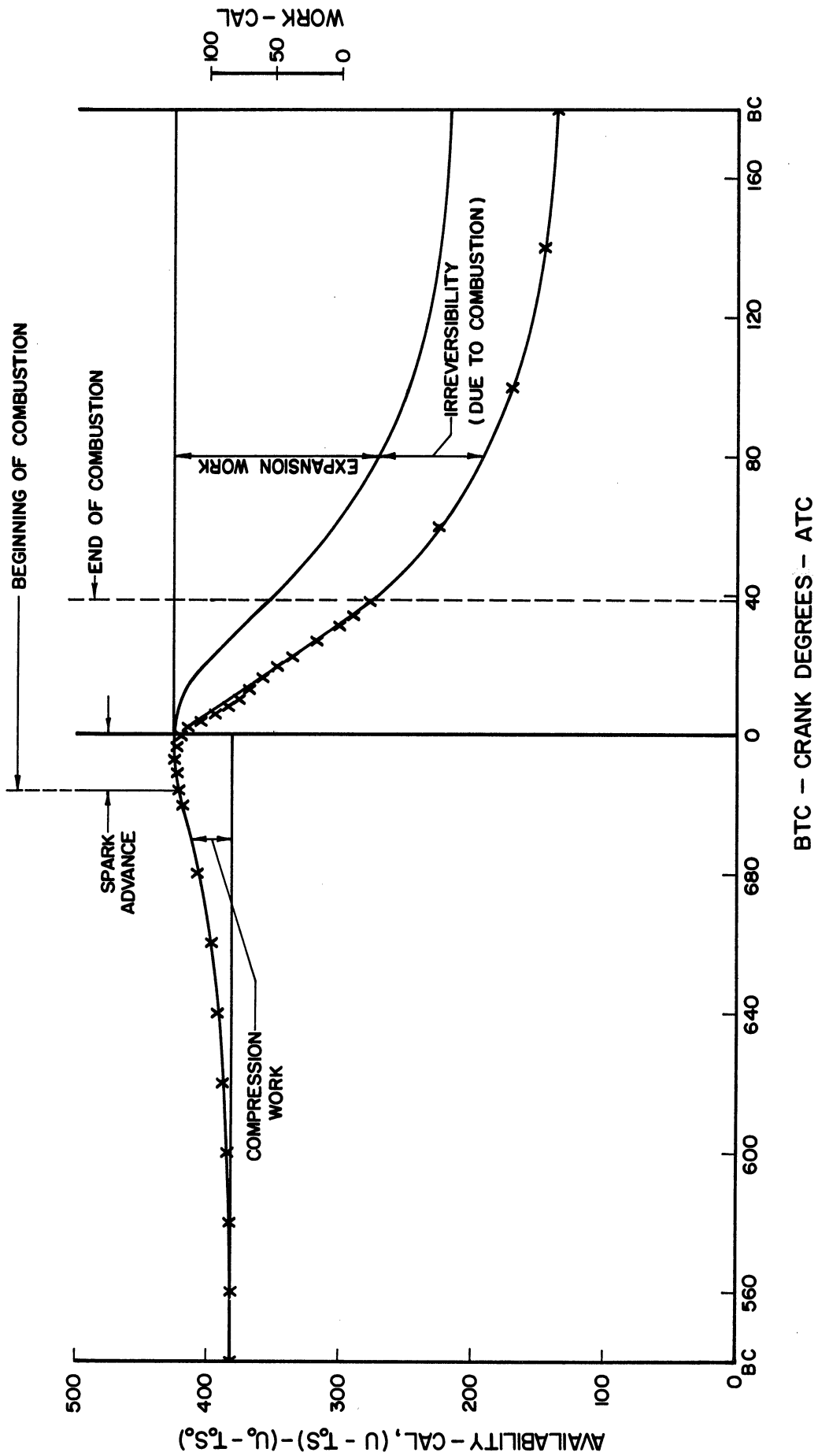


Figure 20. Availability vs crank angle for a 10:1 compression ratio engine with no heat transfer and normal flame propagation rate (Data Set A).

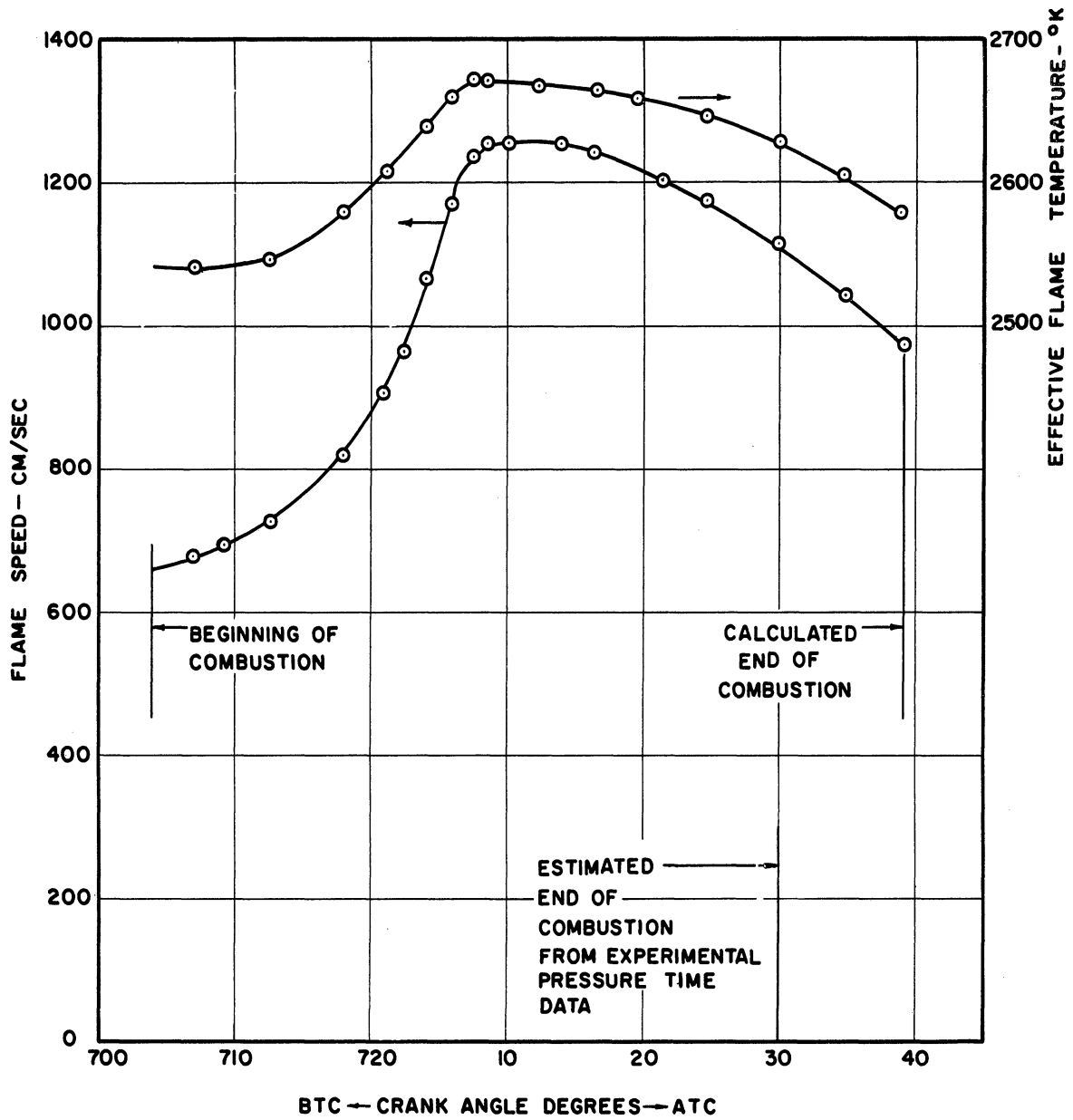


Figure 21. Flame speed and effective flame temperature vs crank angle for a 10:1 compression ratio engine with normal flame propagation and heat transfer rates (Data Set A).

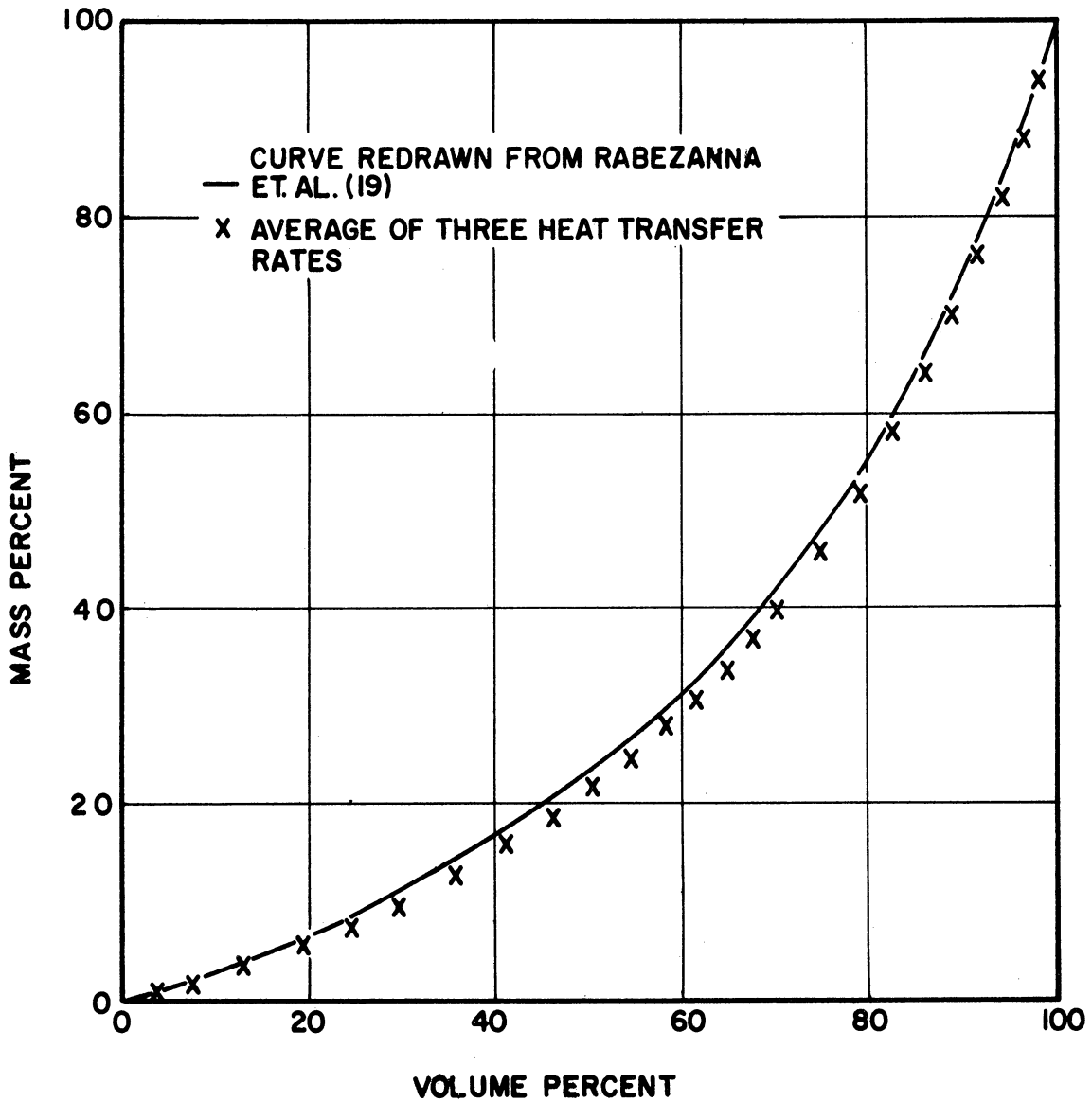


Figure 22. Relationship between mass and volume of the burned charge for a 10:1 compression ratio engine with normal rate of flame propagation (Data Set A).

APPENDIX A

POLYNOMIAL APPROXIMATION BY THE METHOD OF LEAST SQUARES

In the discussion below the most important features of polynomial approximation by the method of least squares are presented along with an adaptation of this general method to the specific problem of obtaining empirical heat capacity equations from tabular data. For a complete treatment of the least squares method see a standard work on numerical analysis such as Reference 31.

If a function $f(x)$ is specified at a set of $m + 1$ equally reliable discrete points, x_0, x_1, \dots, x_m , an n 'th degree polynomial of the form

$$f(x) \approx a_0 + a_1x + a_2x^2 + \dots = \sum_{k=0}^n a_k x^k = y(x)$$

may be used to approximate $f(x)$ over the set of points.

Defining the error ' v ' by the equation

$$v = f(x) - y(x) = f(x) - \sum_{k=0}^n a_k x^k$$

the principle of least squares defines the value of the a_k 's in such a way that the sum of the v^2 's is a minimum. Relating v_i to x_i , this principle may be formally stated as follows:

$$\sum_{i=0}^m v_i^2 = \sum_{i=0}^m \left[f(x_i) - \sum_{k=0}^n a_k x_i^k \right]^2 = \text{A Minimum}$$

For a given set, x_0, x_1, \dots, x_m , the above expression is a function of the a_k 's only.

To determine this minimum value, this function is differentiated with respect to each of the a_k 's and the resulting expressions equated to zero.

Thus:

$$2 \sum_{i=0}^m v_i = 0 \quad (k = 1)$$

$$2 \sum_{i=0}^m v_i x_i = 0 \quad (k = 2)$$

$$\vdots \quad \quad \quad \vdots$$

$$2 \sum_{i=0}^m v_i x_i^n = 0 \quad (k = n)$$

Noting that

$$v_i = f(x_i) - y(x_i) = y_i - \sum_{k=0}^n a_k x_i^k$$

where $y_i = f(x_i)$, the above equations can be rewritten as follows:

$$a_0(m+1) + a_1 \sum x_i + \dots + a_n \sum x_i^n = \sum y_i$$

$$a_0 \sum x_i + a_1 \sum x_i^2 + \dots + a_n \sum x_i^{n+1} = \sum x_i y_i$$

$$\vdots \quad \quad \quad \vdots \quad \quad \quad \vdots \quad \quad \quad \vdots$$

$$a_0 \sum x_i^n + a_1 \sum x_i^{n+1} + \dots + a_n \sum x_i^{n+n} = \sum x_i^n y_i$$

The summations are over all $m + 1$ points. The above set of $n + 1$ equations can then be solved for the $n + 1$ unknowns (the a_k 's).

Note the coefficient matrix.

$$\begin{array}{cccc}
 m+1 & \sum x_i & \dots & \sum x_i^n \\
 & \sum x_i & \sum x_i^2 & \dots & \sum x_i^{n+1} \\
 & \vdots & \vdots & & \vdots \\
 & \vdots & \vdots & & \vdots \\
 & \sum x_i^n & \sum x_i^{n+1} & \dots & \sum x_i^{n+n}
 \end{array}$$

is symmetric.

For the special case when $f(x) = C_p$, $x = T$, and $n = 3$, so that

$$C_p = a_0 + a_1 T + a_2 T^2 + a_3 T^3$$

the above set of equations reduces to:

$$\begin{aligned}
 a_0 (m+1) + a_1 \sum T + a_2 \sum T^2 + a_3 \sum T^3 &= \sum C_p \\
 a_0 \sum T + a_1 \sum T^2 + a_2 \sum T^3 + a_3 \sum T^4 &= \sum C_p T \\
 a_0 \sum T^2 + a_1 \sum T^3 + a_2 \sum T^4 + a_3 \sum T^5 &= \sum C_p T^2 \\
 a_0 \sum T^3 + a_1 \sum T^4 + a_2 \sum T^5 + a_3 \sum T^6 &= \sum C_p T^3
 \end{aligned}$$

Hildebrand⁽³¹⁾, Chapter 10, describes a number of methods for solving such a system of non-homogeneous linear equations for a_0 , a_1 , a_2 , and a_3 .

The root mean square error involved in such a polynomial approximation is defined as:

$$\text{RMS Error} = \left[\sum_{i=0}^m v_i^2 / (m+1) \right]^{1/2}$$

APPENDIX B

EQUILIBRIUM CONSTANT AS A FUNCTION OF TEMPERATURE

Introduction

The reader is referred to a text on chemical engineering thermodynamics for a more complete treatment of this subject. See, for example, Weber and Meissner⁽³²⁾, Chapter 21, Lewis and Randall⁽³³⁾, Chapter 20, or Hougen, Watson, and Ragatz⁽²⁵⁾, Chapter 25.

If \bar{G} is defined as the partial molal free-energy of a substance in any given state of pressure and temperature and \bar{G}° that in the standard state, then

$$\bar{G} - \bar{G}^\circ = RT \ln f/f^\circ \quad (\text{B-1})$$

where f is the fugacity in the given state and f° is the fugacity in the standard state. For a chemical reaction of the form



where l, m, \dots, q, r, \dots are the moles of component L, M, \dots Q, R, \dots respectively, the change in free-energy for this reaction in any state is

$$\Delta G = (q\bar{G}_Q + r\bar{G}_R + \dots) - (l\bar{G}_L + m\bar{G}_M + \dots) \quad (\text{B-3})$$

By definition,

$$a = f/f^\circ \quad (\text{B-5})$$

where 'a' is the activity in the nonstandard state. Combining Equations (B-1) through (B-5), the following equation is obtained:

$$\Delta G = \Delta G^\circ + RT \ln \frac{a_Q^q a_R^r \dots}{a_L^l a_M^m \dots} \quad (\text{B-6})$$

When the reaction has reached a point such that there is no net change in the number of moles of any constituent, it is said that chemical equilibrium has been attained. The criterion for this equilibrium is that at constant temperature and pressure, there can be no change in the free-energy of the system. Consequently, when these limitations are imposed on Equation (B-6), ΔG must be zero for equilibrium and the following familiar expression for the equilibrium constant is obtained:

$$\begin{aligned} \Delta G^\circ &= - RT \ln K_a \\ K_a &= \frac{a_Q^q a_R^r \dots}{a_L^l a_M^m \dots} \end{aligned}$$

For a system in which the behavior of each component can be described by the equation of state

$$P v = RT \quad (\text{B-7})$$

the fugacity of each component becomes identically equal to its partial pressure in the system at all temperatures and pressures for which (B-7) is valid. When this is the case, the equilibrium constant is denoted by K_p . Choosing the standard state as the pure component at one atmosphere pressure, the equations become,

$$\begin{aligned} \Delta G^\circ &= - RT \ln K_p \\ K_p &= \frac{p_Q^q p_R^r \dots}{p_L^l p_M^m \dots} \end{aligned} \quad (\text{B-8})$$

where p_i is the partial pressure of the i 'th component. K_p is then referred to as the equilibrium constant at one atmosphere. The free-energy change is termed the standard free-energy change of the reaction. This quantity depends upon temperature, the definition of the standard state of each component, and the number of moles entering into the stoichiometric Equation (B-2). Once the standard state pressure is fixed, the equilibrium constant becomes independent of the system pressure.

Analogically to expression (B-4), expressions for ΔH° and ΔS° , the standard state change in enthalpy and entropy for the reaction can be found. It immediately follows that

$$\Delta G^\circ = \Delta H^\circ - T\Delta S^\circ$$

Effect of Temperature on K_p

To determine the effect of temperature on the equilibrium constant, Equation (B-8) can be differentiated as follows:

$$\begin{aligned} \left[\frac{\partial}{\partial T} (\ln K_p) \right]_P &= - \left[\frac{\partial}{\partial T} \left(\frac{\Delta G^\circ}{RT} \right) \right]_P \\ &= - \frac{1}{R} \left[\frac{1}{T} \frac{\partial \Delta H^\circ}{\partial T} - \frac{\Delta H^\circ}{T^2} - \frac{\partial \Delta S^\circ}{\partial T} \right]_P \end{aligned}$$

where $\frac{\Delta G^\circ}{T} = \frac{\Delta H^\circ}{T} - \Delta S^\circ$.

However since $\left(\frac{\partial \Delta H^\circ}{\partial T} \right)_P = T \left(\frac{\partial \Delta S^\circ}{\partial T} \right)_P$, the above simplifies to:

$$\left[\frac{\partial}{\partial T} (\ln K_p) \right]_P = \left[\frac{\Delta H^\circ}{RT^2} \right]_P$$

Since the quantity $\frac{\Delta H^\circ}{RT^2}$ is independent of pressure, this restriction can be removed. Then:

$$\frac{d}{dT} (\ln K_p) = \frac{\Delta H^\circ}{RT^2}$$

Integrating this equation between T and T₀, where T₀ is some reference temperature, and recalling that:

$$\Delta H^\circ = \Delta H_{T_0}^\circ + \int_{T_0}^T \sum_{\text{Products}} n_i C_{pi}^\circ dT + \int_T^{T_0} \sum_{\text{Reactants}} n_i C_{pi}^\circ dT$$

the following equation for equilibrium constant as a function of temperature is obtained:

$$\begin{aligned} \ln K_{p_T} - \ln K_{p_{T_0}} &= \frac{\Delta H_{T_0}^\circ}{R} \left[\frac{1}{T_0} - \frac{1}{T} \right] \\ &+ \int_{T_0}^T \frac{1}{RT^2} \left[\int_{T_0}^T (\sum_{\text{Products}} n_i C_{pi}^\circ - \sum_{\text{Reactants}} n_i C_{pi}^\circ) dT \right] dT \end{aligned} \tag{B-9}$$

APPENDIX C

CALCULATION OF THERMODYNAMIC PROPERTIES
FROM EMPIRICAL HEAT CAPACITY EQUATIONS

A. General

For a system of 's' chemical species, it is assumed that the ideal gas molal heat capacity at one atmosphere of the i-th component can be expressed as follows:

$$C_p^{\circ} = a_{0i} + a_{1i}T + a_{2i}T^2 + a_{3i}T^3$$

298.16 - 1500°K

$$C_p^{\circ} = b_{0i} + b_{1i}T + b_{2i}T^2 + b_{3i}T^3$$

1500 - 3500°K

and that the molal enthalpy of formation and entropy at 298.16 and 1 atm pressure are $(\Delta H_{f298}^{\circ})_i$ and $(S^{\circ}_{298})_i$. The standard state is the ideal gas at one atmosphere pressure. The calculation of the thermodynamic functions is made most conveniently by matrix algebra.

The following matrix is defined for the temperature range 298.16 - 1500°K:

$$A = \begin{vmatrix} a_{01} & a_{02} & \dots & \dots & a_{0s} \\ a_{11} & a_{12} & \dots & \dots & a_{1s} \\ a_{21} & \cdot & \cdot & \cdot & \cdot \\ a_{31} & \cdot & \cdot & \cdot & \cdot \\ (\Delta H_{f298}^{\circ})_1 & \cdot & \cdot & \cdot & \cdot \\ (S^{\circ}_{298})_1 & \cdot & \cdot & \cdot & \cdot \end{vmatrix}$$

and for the range 1500 - 3500°K:

$$B = \begin{vmatrix} b_{01} & b_{02} & \dots & b_{0s} \\ b_{11} & b_{12} & \dots & b_{1s} \\ b_{21} & \cdot & \dots & \cdot \\ b_{31} & \cdot & \dots & \cdot \\ (H_{1500}^{\circ})_1 & \cdot & \dots & \cdot \\ (S_{1500}^{\circ})_1 & \cdot & \dots & \cdot \end{vmatrix}$$

where

$$(H_{1500}^{\circ})_i = (\Delta H_{f298}^{\circ})_i + (H_{1500}^{\circ} - H_{298}^{\circ})_i$$

For a selected number of substances the appropriate constants for matrix A and B are presented in Tables I and II.

It is assumed that the composition of the system of 's' species whose properties are desired is known and that there are n_i moles of component 'i'. The following matrix will then be defined to represent the composition of the system:

$$N = \begin{vmatrix} n_1 \\ n_2 \\ \cdot \\ \cdot \\ \cdot \\ n_s \end{vmatrix}$$

B. Method of Calculation

1. Computation of the Ideal Gas Enthalpy

$$H = \sum n_i (\Delta H_{f298}^{\circ})_i + \sum n_i (H_T^{\circ} - H_{298}^{\circ})_i$$

Consider the second summation. For the temperature range 298.16 to 1500°K

$$\begin{aligned} \sum n_i (H_T^{\circ} - H_{298}^{\circ})_i &= \sum n_i \int_{T_0=298.16}^T c_{p_i}^{\circ} dT \\ &= \sum n_i \left[T_0 a_{0i} (\theta_A - 1) + \frac{T_0^2 a_{1i}}{2} (\theta_A^2 - 1) + \frac{T_0^3 a_{2i}}{3} (\theta_A^3 - 1) + \frac{T_0^4 a_{3i}}{4} (\theta_A^4 - 1) \right] \end{aligned}$$

For the temperature range 1500 to 3500°K

$$\begin{aligned} \sum n_i (H_T^{\circ} - H_{298}^{\circ})_i &= \sum n_i \left[\int_{T'_0=1500}^T c_{p_i}^{\circ} dT + (H_{1500}^{\circ} - H_{298}^{\circ})_i \right] \\ &= \sum n_i \left[T'_0 b_{0i} (\theta_B - 1) + \frac{T'^0_0{}^2 b_{1i}}{2} (\theta_B^2 - 1) + \frac{T'^0_0{}^3 b_{2i}}{3} (\theta_B^3 - 1) + \frac{T'^0_0{}^4 b_{3i}}{4} (\theta_B^4 - 1) \right. \\ &\quad \left. + (H_{1500}^{\circ} - H_{298}^{\circ})_i \right] \end{aligned}$$

where

$$\theta_A = T/298.16$$

$$\theta_B = T/1500$$

If the matrices K_A and K_B are defined as follows:

$$K_A = \begin{vmatrix} 298.16 (\theta_A - 1) \\ \frac{(298.16)^2}{2} (\theta_A^2 - 1) \\ \frac{(298.16)^3}{3} (\theta_A^3 - 1) \\ \frac{(298.16)^4}{4} (\theta_A^4 - 1) \\ 1 \\ 0 \end{vmatrix} = \begin{vmatrix} 298.16 (\theta_A - 1) \\ 44,449.69 (\theta_A^2 - 1) \\ 8,835,413.6 (\theta_A^3 - 1) \\ 19,757,752 \cdot 10^2 (\theta_A^4 - 1) \\ 1 \\ 0 \end{vmatrix}$$

$$K_B = \begin{vmatrix} 1500 (\theta_B - 1) \\ \frac{(1500)^2}{2} (\theta_B^2 - 1) \\ \frac{(1500)^3}{3} (\theta_B^3 - 1) \\ \frac{(1500)^4}{4} (\theta_B^4 - 1) \\ 1 \\ 0 \end{vmatrix} = \begin{vmatrix} 1500 (\theta_B - 1) \\ 1125 \cdot 10^3 (\theta_B^2 - 1) \\ 1125 \cdot 10^6 (\theta_B^3 - 1) \\ 1,265,625 \cdot 10^6 (\theta_B^4 - 1) \\ 1 \\ 0 \end{vmatrix}$$

the enthalpy of the mixture then can be calculated as the product of matrices.

$$H = \begin{cases} K_A^t \cdot A \cdot N \\ K_B^t \cdot B \cdot N \end{cases} \begin{cases} 298.16 \leq T \leq 1500 \\ 1500 < T \leq 3500 \end{cases} \quad (C-1)$$

(in cal)

2. Computation of the Ideal Gas Entropy

$$S = \sum n_i S_{298}^0 + \sum n_i (S_T^0 - S_{298}^0)_i - R \sum n_i \ln(x_i P)$$

where

x_i = mole fraction of the i-th component

P = total system pressure

Consider the second summation. For the temperature range 298.16 to 1500°K

$$\begin{aligned} \sum n_i (S_T^0 - S_{298}^0)_i &= \sum n_i \int_{T_0=298.16}^T \frac{C_{p_i}^0}{T} dT \\ &= \sum n_i \left[a_{0i} \ln \theta_A + T_0 a_{1i} (\theta_A - 1) + \frac{T_0^2 a_{2i}}{2} (\theta_A^2 - 1) + \frac{T_0^3 a_{3i}}{3} (\theta_A^3 - 1) \right] \end{aligned}$$

For the temperature range 1500 to 3500°K

$$\begin{aligned} \sum n_i (s_T^o - s_{298}^o)_i &= \sum n_i \left[\int_{T'_0=1500}^T \frac{C_{p_i}}{T} dT + (s_{1500}^o - s_{298}^o)_i \right] \\ &= \sum n_i \left[b_{0i} \ln \theta_B + T'_0 b_{1i} (\theta_B - 1) + \frac{T'_0{}^2 b_{2i}}{2} (\theta_B^2 - 1) + \frac{T'_0{}^3 b_{3i}}{3} (\theta_B^3 - 1) \right. \\ &\quad \left. + (s_{1500}^o - s_{298}^o)_i \right] \end{aligned}$$

where

$$\theta_A = T/298.16$$

$$\theta_B = T/1500$$

If the matrices K_A and K_B are defined now as follows:

$$K_A = \begin{vmatrix} \ln(\theta_A) \\ 298.16(\theta_A - 1) \\ 44449.693(\theta_A^2 - 1) \\ 8835413.6(\theta_A^3 - 1) \\ 0 \\ 1 \end{vmatrix} \quad K_B = \begin{vmatrix} \ln(\theta_B) \\ 1500(\theta_B - 1) \\ 1125 \cdot 10^3(\theta_B^2 - 1) \\ 1125 \cdot 10^6(\theta_B^3 - 1) \\ 0 \\ 1 \end{vmatrix}$$

the entropy of the mixture then can be calculated as follows:

$$S = \left\{ \begin{matrix} K_A^t \cdot A \cdot N \\ K_B^t \cdot B \cdot N \end{matrix} \right\} - R \sum n_i \ln(x_i P) \quad \begin{cases} 298.16 \leq T \leq 1500^\circ\text{K} \\ 1500 < T \leq 3500^\circ\text{K} \end{cases} \quad (\text{C-2})$$

(in cal per °K)

3. Computation of $(\frac{\partial H}{\partial T})_P$

The procedure here is the same as in Parts 1 and 2. If a matrix K is defined as follows:

$$K = \begin{vmatrix} 1 \\ T \\ T^2 \\ T^3 \\ 0 \\ 0 \end{vmatrix}$$

$(\frac{\partial H}{\partial T})_P$ then can be computed as the following product:

$$\begin{aligned} (\frac{\partial H}{\partial T})_P &= C_p = K^t \cdot \begin{Bmatrix} A \\ B \end{Bmatrix} \cdot N & \begin{cases} 298.16 \leq T \leq 1500^\circ\text{K} \\ 1500 < T \leq 3500^\circ\text{K} \end{cases} & (C-3) \\ \text{(in cal per } ^\circ\text{K)} & & \end{aligned}$$

4. Computation of $(\frac{\partial S}{\partial T})_P$

Identically to Part 3, if a matrix K is defined as follows:

$$K = \begin{vmatrix} 1/T \\ 1 \\ T \\ T^2 \\ 0 \\ 0 \end{vmatrix}$$

$(\frac{\partial S}{\partial T})_P$ then can be computed as the following product:

$$\begin{aligned} (\frac{\partial S}{\partial T})_P &= K^t \cdot \begin{Bmatrix} A \\ B \end{Bmatrix} \cdot N & \begin{cases} 298.16 \leq T \leq 1500^\circ\text{K} \\ 1500 < T \leq 3500^\circ\text{K} \end{cases} & (C-4) \\ \text{(in cal per } ^\circ\text{K}^2) & & \end{aligned}$$

5. Computation of the Equilibrium Constant K_p

A general expression for K_p as a function of T with reference state at T_0 is as follows: (see Equation (B-9))

$$\begin{aligned} \ln K_{p_T} - \ln K_{p_{T_0}} &= \frac{\Delta H_{T_0}^{\circ}}{R} \left[\frac{1}{T_0} - \frac{1}{T} \right] + \int_{T_0}^T \frac{1}{RT^2} \left[\left(\sum_{\text{Products}} n_i C_{p_i}^{\circ} - \sum_{\text{Reactants}} n_i C_{p_i}^{\circ} \right) dT \right] dT \\ &= \frac{\Delta H_{T_0}^{\circ}}{RT_0} \left[1 - \frac{1}{\theta} \right] + \frac{1}{R} \left[c_1 \left(\ln \theta + \frac{1}{\theta} - 1 \right) \right. \\ &\quad \left. + \frac{c_2 T_0}{2} \left(\theta + \frac{1}{\theta} - 2 \right) + \frac{c_3 T_0^2}{6} \left(\theta^2 + \frac{2}{\theta} - 3 \right) + \frac{c_4 T_0^3}{12} \left(\theta^3 + \frac{3}{\theta} - 4 \right) \right] \end{aligned}$$

where

$$\theta = T/T_0$$

$$c_j = \sum_{\text{Products}} n_i b_{ji} - \sum_{\text{Reactants}} n_i b_{ji} \quad (j = 0, 1, 2, 3)$$

For the temperature range $1500 \leq T \leq 3500^\circ\text{K}$, let $T_0 = 1500^\circ\text{K}$, then

$\theta = \theta_B = T/1500$ and note that

$$\ln K_{p_{1500}} = \frac{-\Delta G_{1500}^{\circ}}{R \cdot 1500}$$

Then substituting

$$\begin{aligned} \ln K_{p_T} &= \frac{-\Delta G_{1500}^{\circ}}{2980.785} + \frac{\Delta H_{1500}^{\circ}}{2980.785} \left[1 - \frac{1}{\theta_B} \right] + \frac{1}{R} \left[c_1 \left(\ln \theta_B + \frac{1}{\theta_B} - 1 \right) \right. \\ &\quad \left. + 750 \cdot c_2 \left(\theta_B + \frac{1}{\theta_B} - 1 \right) + 375 \cdot 10^3 \cdot c_3 \left(\theta_B^2 + \frac{2}{\theta_B} - 3 \right) \right. \\ &\quad \left. + 28125 \cdot 10^4 \cdot c_4 \left(\theta_B^3 + \frac{3}{\theta_B} - 4 \right) \right] \end{aligned} \quad (C-5)$$

where ΔG_{1500}° and ΔH_{1500}° are the standard free-energy and enthalpy of reaction at 1500°K in cal per mole and are listed in Table II for selected reactions.

C. Computer Programs

The computation of the thermodynamic properties and equilibrium constants by a computer is most conveniently made if the computational schemes are incorporated into a number of computer sub-programs or external functions. Such programs have been written in the MAD language for an IBM 704 or 709 computer and are listed in Appendix I. The functions for computing enthalpy, entropy, $\frac{dH}{dT}$, and $(\frac{\partial S}{\partial T})_p$, are called ENTHAL., ENTROP., HPRIME., and SPRIME., respectively. That for computing the equilibrium constants is called EQUIL.

APPENDIX D

THE TIME RATE OF TEMPERATURE CHANGE DURING EXPANSION

In order to calculate the time rate of temperature change, $\partial T/\partial t$ during the expansion process of an internal combustion engine, the overall expansion can be approximated by a polytropic process. The maximum rate of change of volume with time is

$$\frac{dV}{dt} = (2\pi r N / 60) (\pi b^2 / 4)$$

where

r = the crank radius

b = the bore

N = the engine speed, rpm

The temperature in the polytropic process is

$$T = T_1 \left(\frac{V}{V_1} \right)^{1-n}$$

and

$$\frac{\partial T}{\partial t} = T (1-n) \left(\frac{V}{V_1} \right)^{-n} \cdot \frac{d}{dt} \left(\frac{V}{V_1} \right)$$

Substituting

$$- \frac{\partial T}{\partial t} = \frac{T_1}{V_1} (n-1) \left(\frac{V}{V_1} \right)^{-n} \cdot \pi^2 r b^2 N / 120 \quad (D-1)$$

where

T_1 = the temperature at TC

V_1 = the volume at TC

n = the polytropic exponent

Some typical values of the constants are as follows:

$$b = 4 \text{ in}$$

$$r = 1.25 \text{ in}$$

$$V_1 = 3.33 \text{ cu in}$$

$$T_1 = 2500^\circ\text{K}$$

$$N = 3000 \text{ rpm}$$

$$n = 1.3$$

Substituting these values in (D-1),

$$= \frac{\partial T}{\partial t} \approx 10^6 \cdot \left(\frac{V_1}{V}\right)^{1.3} \text{ }^\circ\text{K/sec}$$

$\left(\frac{V_1}{V}\right)^{1.3}$ will be between 0.1 and 1 so that,

$$= \frac{\partial T}{\partial t} \approx 10^5 \text{ to } 10^6 \text{ }^\circ\text{K/sec}$$

APPENDIX E

METHOD OF SOLUTION OF THE EQUATIONS OF SIMULTANEOUS GAS EQUILIBRIUM

Equations (2-19) through (2-21) are repeated here for convenience

$$Z = -\frac{X}{4K_4} + \sqrt{\left(\frac{X}{4K_4}\right)^2 + \frac{F \cdot N_n}{2}} \quad (2-19)$$

$$Y = \frac{-(X/K_3 + 1/K_6) + \sqrt{(X/K_3 + 1/K_6)^2 + 8(1 + X/K_2) \cdot F \cdot N_n}}{4(1 + X/K_2)} \quad (2-20)$$

$$F \cdot N_0 = \frac{X/K_1 \cdot F \cdot N_c}{1 + X/K_1} + \frac{Y^2 X}{K_2} + 2X^2 + \frac{YX}{K_3} + \frac{ZX}{K_4} + F \cdot N_c + \frac{X}{K_5} \quad (2-21)$$

The problem is to find the value of X which satisfies Equation (2-21). This equation can be restated as follows:

$$f(X) = 0 \quad (E-1)$$

For any estimated value X_1 the right hand side in general is not zero,

$$f(X_1) \neq 0$$

and some correction to $f(X_1)$, $\Delta f(X_1)$ must be made in order to equate it to zero.

$$f(X_1) + \Delta f(X_1) = 0 \quad (E-2)$$

From (E-2), it is hoped that a correction ΔX_1 to X_1 can be found giving a value X_2 which is closer to the correct answer. This correction is

$$\Delta f(X_1) \cong \left(\frac{df(X)}{d(\ln X)}\right)_1 \cdot \Delta \ln X \quad (E-3)$$

Equating Equations (E-2) and (E-3),

$$f(X_1) = - \left(\frac{df(X)}{d(\ln X)} \right)_1 \cdot \Delta \ln X = - \frac{f'(X_1)}{X_1} \cdot \Delta \ln X$$

Re-arranging,

$$\ln X_2 - \ln X_1 = - \frac{f(X_1)}{X_1 f'(X_1)}$$

or

$$X_2 = X_1 \cdot \exp \frac{-f(X_1)}{X_1 f'(X_1)}$$

In general, for the i-th trial,

$$X_{i+1} = X_i \cdot \exp \frac{-f(X_i)}{X_i f'(X_i)} \quad (E-4)$$

The reason for generating X_{i+1} in this particular manner is to assure that it will always be positive if X_1 initially is assumed positive. Thus the possibility of obtaining a solution with some negative partial pressures is eliminated.

The calculation of $f'(X)$ presents some problem. In the following equations the argument X will be dropped for convenience.

$$f' = \frac{df}{dX} = \left(\frac{\partial f}{\partial Y} \right) \left(\frac{dY}{dX} \right) + \left(\frac{\partial f}{\partial Z} \right) \left(\frac{dZ}{dX} \right) + \frac{\partial f}{\partial X} \quad (E-5)$$

From Equation (2-21)

$$\begin{aligned} \frac{\partial f}{\partial Y} &= \frac{2XY}{K_2} + \frac{X}{K_3} & \frac{\partial f}{\partial Z} &= \frac{X}{K_4} \\ \frac{\partial f}{\partial X} &= \frac{f \cdot N_c \cdot K_1}{(X + K_1)^2} + 4X + \frac{Y^2}{K_2} + \frac{Y}{K_3} + \frac{Z}{K_4} + \frac{1}{K_5} \end{aligned} \quad (E-6)$$

From Equation (2-20)

$$\frac{dY}{dX} = \left\{ -\alpha \left[\frac{1}{K_3} - \left(\frac{\delta}{K_3} + \frac{4F \cdot N_h}{K_2} \right) / \gamma \right] - 4/K_2 (\gamma - \delta) \right\} / \alpha^2 \quad (E-7)$$

where

$$\begin{aligned} \alpha &= 4 (1 + X/K_2) \\ \delta &= X/K_3 + 1/K_6 \\ \gamma &= \sqrt{\delta^2 + 2\alpha F \cdot N_h} \end{aligned}$$

From Equation (2-19)

$$\frac{dZ}{dX} = \frac{X/K_4\beta - 1}{4K_4} \quad (E-8)$$

where

$$\beta = \sqrt{\left(\frac{X}{K_4}\right)^2 + 8 F \cdot N_h}$$

If $f(X)$ and $f'(X)$ are evaluated at X_i by Equations (2-21) and (E-5), a new values of X , X_{i+1} , can be calculated from Equation (E-4). The process can be repeated until $\left| \frac{X_{i+1} - X_i}{X_i} \right|$ is less than some prescribed value ϵ_2 . The number of iterations required for convergence is about four, plus an additional four for each power of ten from which the estimate differs from the correct answer. Table XI gives the accuracy with which the partial pressures are calculated for various values of ϵ_2 . It is assumed that the partial pressures are of the form $0.\text{XXXXXXXX} \cdot 10^{\text{XX}}$. The number entered in the table is the digit which changes when the value of ϵ_2 was decreased by successive powers of 10.

In connection with the actual mechanics of the computer subroutine (see the flow diagram, Figure 23) which solves these equations for the partial pressures and moles of each component, it should be pointed out that when Equation (E-4) is being computed and the quantity $\left| \frac{f(X_i)}{X_i f'(X_i)} \right|$ exceeds 16, X_i is multiplied or divided by the quantity 'e', the base of the natural logarithms. This precludes any wild fluctuation of the successive estimates for X . In the many thousands of times which this subroutine has been called upon to compute the equilibrium composition, it has never failed to converge on the correct answers regardless of the value of the initial estimate.

TABLE XI

DIGIT CHANGED IN THE PARTIAL PRESSURE OF EACH CONSTITUENT
FOR SELECTED VALUES OF RELATIVE ERROR

Stoichiometric Octene-Air Mixture
Temperature, 2500°K
Pressure, 19 Atm

Relative Error $\epsilon_2 \leq \frac{X_{i+1} - X_i}{X_i}$	O ₂	N ₂	CO ₂	H ₂ O	CO	H ₂	OH	NO	O	H
0.1	-	-	-	-	-	-	-	-	-	-
0.01	4	7	5	6	4	4	4	4	4	4
0.001	5	8	6	7	5	5	5	5	5	5
0.0001	7	8	NC	7	7	7	7	7	7	7
0.00001	NC	NC	NC	NC	NC	8	NC	8	8	8

NC - No Change

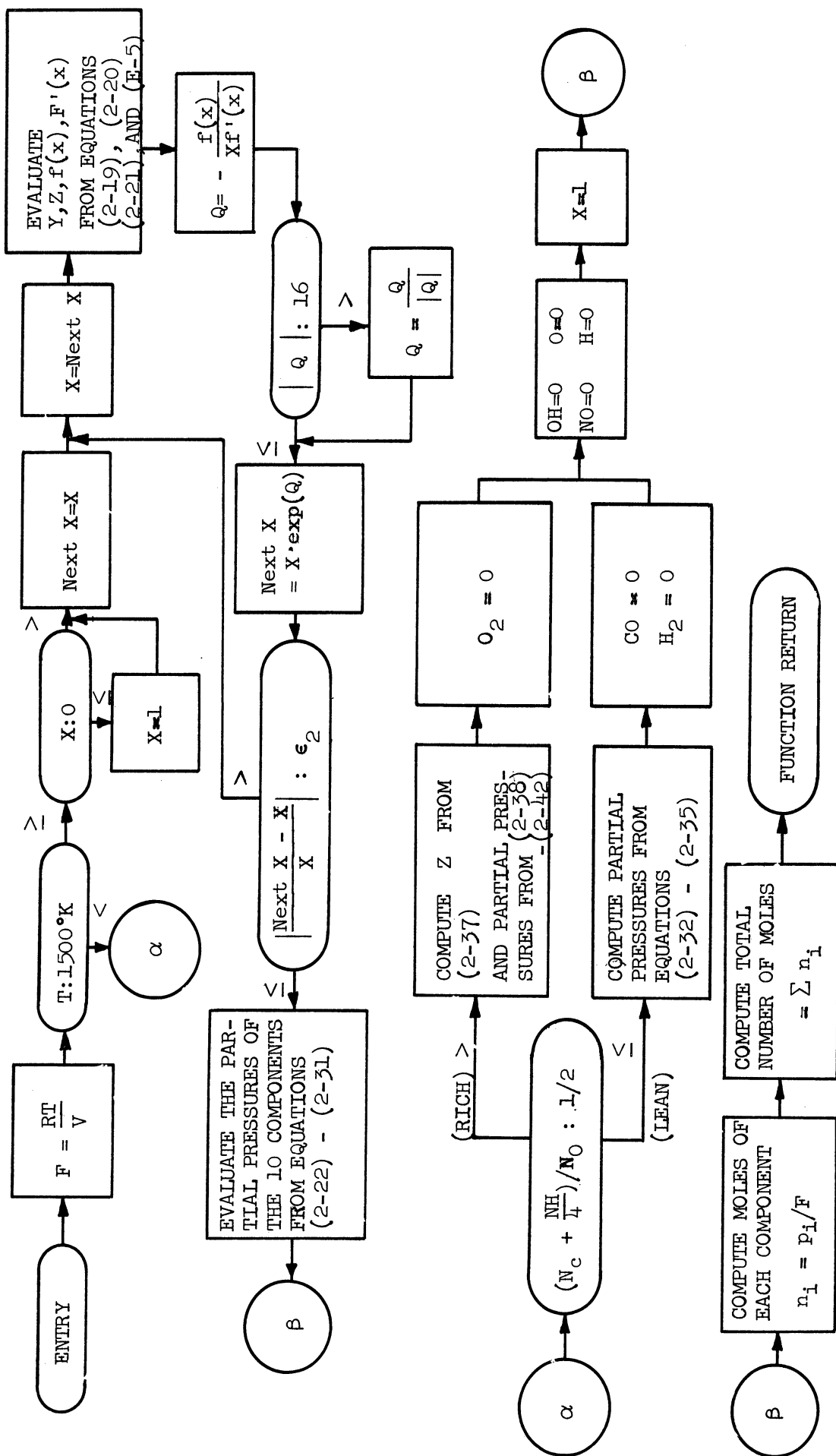


Figure 23. Flow diagram for the subroutine computing the composition of the products of combustion.

APPENDIX F

MATHEMATICAL ANALYSIS, FLOW DIAGRAM, AND COMPUTER PROGRAM LISTING FOR THE FUEL-AIR CYCLE

A. General

The mathematical analysis of the fuel-air cycle treats a number of quantities as variables. These are the fuel, the air-fuel ratio, the manifold temperature and pressure, the exhaust pressure, and the compression ratio. The fuel is described by specifying the number of carbon and hydrogen atoms per molecule of fuel; its enthalpy of combustion at the reference state; the constants for its empirical heat capacity equation, Table I; and its enthalpy of formation and entropy at the reference state, Table II. Two additional variables arise because of the iterative nature of some of the calculations. These are ϵ_1 , the allowable relative error in those thermodynamic properties calculated by trial and error methods, and ϵ_2 , the allowable error in the equilibrium composition (see Appendix E).

Among the quantities calculated from the cycle analysis are the pressure, temperature, enthalpy, entropy, and internal energy at each point of Figure 3. The indicated thermal efficiency, mean effective pressure, and exhaust gas residual fraction are also calculated.

The analysis assumes that all enthalpies and entropies and their derivatives are computed by the methods of Chapter II, Section B, and the equilibrium composition of the products of combustion is computed by those of Chapter II, Section D. Internal energies and their derivatives are computed from the corresponding quantities for the enthalpy.

For a perfect gas, these are

$$U = H - nRT \text{ and } \frac{dU}{dT} = \frac{dH}{dT} - nR$$

Subscripts on thermodynamic properties refer to points on Figure

3. The metric system is used throughout.

B. Mathematical Analysis

1. Exhaust Residual - Initial Approximation

Computation begins with an assumption of the residual exhaust gas temperature T_5 . If the clearance volume is one liter and the pressure is P_e , the total number of moles remaining in the clearance volume is

$$n_5 = P_e / RT_5 \quad (\text{F-1})$$

An approximate exhaust gas composition can be calculated if it is assumed that the total number of moles of products per mole of fuel burned is the sum of the coefficients of the products in Equation 2-2. This quantity is

$$n_{t0t} = a + b/2 + (a + b/4)(cx + x - 1) \quad (\text{F-2})$$

The number of moles of each constituent present in the exhaust then will be its coefficient from Equation (2-2) multiplied by $\frac{n_5}{n_{t0t}}$. In the case of oxygen, the absolute value is taken to avoid a negative value for rich mixtures. Calculated in this manner, the exhaust composition is only approximate.

This initial approximation is not repeated for subsequent cycles. At the end of the complete cycle, a new exhaust temperature and composition is computed from equilibrium considerations. The cycle is then repeated until there is an insignificant change in these quantities from cycle to cycle. To facilitate the computation of the first cycle, the allowable

relative error ϵ_1 , discussed later, is increased by a factor of 100.

2. Mixing of the Fresh Charge and Exhaust Residual

The inlet conditions, Point 0, being known, the thermodynamic properties of the fresh charge can be calculated per mole. After induction, the mixture will be contained in a volume of $1 \cdot r$ liters, where 'r' is the compression ratio. The number of moles of fresh charge and the final temperature of the mixture are to be found.

If an initial guess T'_1 is made for the mixture temperature, the number of moles of fresh charge is

$$n'_0 = \frac{P_i r}{RT'_1} - n_5 \quad (\text{F-3})$$

The prime denotes a temporary or approximate value. In a manner similar to that of Part 1, the individual number of moles of fuel, oxygen, and nitrogen can be calculated. If this entering mass is lumped with the residual exhaust gas and the enthalpy of the mixture is calculated, at T'_1 a value H'_1 is found.

For the ideal adiabatic inlet process envisioned here and described in detail in Reference 45, Chapter 4, the enthalpy of the mixture is

$$H_1 = H'_0 + U_5 + P_i r J \quad (\text{F-4})$$

Whenever

$$\left| \frac{H'_1 - H_1}{H_1} \right| > \epsilon_1 \quad (\text{F-5})$$

where ϵ_1 is defined as the relative error, a corrected value for T'_1 , T''_1 , must be found. This corrected value can be found by Newton's method, a

derivation for which is found in Appendix G. The new Value is

$$T_1''' = T_1' - \frac{H_1' - H_1}{\left(\frac{dH_1}{dT}\right)'} \quad (F-6)$$

where, in light of Equations F-3 and F-4,

$$\left(\frac{dH_1}{dT}\right)' = \frac{H_0'}{T_1'} \left(1 - \frac{n_5}{n_0'}\right) + \frac{dH_0'}{dT} \quad (F-7)$$

This process is repeated until Equation(F-5) is not satisfied. The conditions at Point 1 are now known.

3. Isentropic Compression

At Point 2, the volume is one liter and $S_2 = S_1$. The composition remains unchanged during the compression. The temperature can be found by the following trial and error method. Assume a temperature T_2' . Then

$$P_2' = n_2 R T_2' \quad (F-8)$$

where $n_2 = n_1$. The entropy S_2' then can be calculated. Whenever

$$\left| \frac{S_2' - S_2}{S_2} \right| > \epsilon_1 \quad (F-9)$$

a new value T_2'' must be found.

$$T_2''' = T_2' - \frac{S_2' - S_2}{\left(\frac{\partial S_2}{\partial T}\right)'_V} \quad (F-10)$$

where

$$\left(\frac{\partial S_2}{\partial T}\right)'_V = \left(\frac{\partial S_2}{\partial T}\right)'_P = \frac{n_2 R}{T_2'} \quad (F-11)$$

The process is repeated until Equation(F-9) is not satisfied. The conditions at Point 2 are now known.

4. Constant Volume Adiabatic Combustion

This process occurs at constant internal energy and constant volume, that is $U_3 = U_2$ and $V_3 = 1$. Moreover the products are in a state of chemical equilibrium. The problem is to find the adiabatic flame temperature and pressure at Point 3. If a temperature T_3' is assumed, the composition can be calculated according to the methods of Chapter II. The internal energy U_3' can then be calculated. Whenever

$$\left| \frac{U_3' - U_3}{U_3} \right| > \epsilon_1 \quad (\text{F-12})$$

a new value T_3'' must be found. One method for finding this is to use Newton's method,

$$T_3'' = T_3' - \frac{U_3' - U_3}{\left(\frac{dU_3}{dT} \right)}, \quad (\text{F-13})$$

wherein it is assumed that the difference between T_3'' and T_3' is so small that the effects of composition change are negligible. Because this is not always true, convergence may be slow or may not be obtained at all. However a second method can be employed, the Half-Interval convergence method. This method, which will always converge on the correct answer, is described in Appendix G. In this analysis both methods are employed. If a satisfactory solution is not found after using Newton's convergence method to generate five successive approximations to T_3 , the Half-Interval method is used. If at the end of this fifth trial, the difference between T_3'' and T_3' is ΔT , the next trial for T_3 will be

$$T_3''' = T_3' - \Delta T \cdot \frac{U_3' - U_3}{|U_3' - U_3|} \quad (\text{F-14})$$

Then whenever the quantity $(U_3^1 - U_3)$ changes sign, ΔT is halved. This half-interval method is continued until Equation F-12 is not satisfied. The conditions at Point 3 are then known.

5. Isentropic Expansion

At Point 4, the volume is 'r' liters and $S_4 = S_3$. The products are still in a state of chemical equilibrium. The temperature T_4 can be found by a method similar to that described in Section 3 for the compression. A temperature T_4' is assumed, the composition calculated, and the entropy S_4' determined. Whenever

$$\frac{S_4' - S_4}{S_4} > \epsilon_1 \quad (\text{F-15})$$

a new value T_4'' must be found. Newton's method can be successfully used to generate T_4' . Whenever Equation F-15 is not satisfied, the conditions at Point 4 can be considered to be known.

6. Expansion to Exhaust Pressure

When the exhaust valve opens the contents of the cylinder expand to the exhaust pressure, the overall process being irreversible. Nevertheless, those gases remaining in the cylinder have undergone an isentropic expansion. At Point 4' $V_4' = r$, $\bar{S}_4' = \bar{S}_4$ and the pressure is P_e . For the purposes of calculating T_4' , let $S_4'' = S_4$, that is, assume all the gas has expanded isentropically. The temperature T_4' can be found by a method identical to that of Section 5. The equilibrium composition is calculated for the case where the pressure and temperature, rather than the volume and temperature are known.

7. Exhaust Residual

At Point 5, the piston has exhausted all the products except those remaining in the clearance volume. The molar thermodynamic properties are identical to those at Point 4'. The fraction of total mass retained in the cylinder is $1/V_4''$. Thus the temperature and composition of the exhaust residual are completely determined.

Whenever

$$\left| \frac{T_{4'} - T_5}{T_5} \right| > \epsilon_1 \quad (\text{F-16})$$

where T_5 is the exhaust gas temperature at the beginning of the cycle and $T_{4'}$ is that at the end, the entire cycle is repeated starting with Section 2. Prior to the second time the cycle is executed the relative error, ϵ_1 , is reduced to its initial value by dividing it by a factor of 100. The cycle is always repeated at least once. When $\epsilon_1 = .00001$, about four complete cycles must be exhausted before Equation (F-16) is not satisfied.

8. Net Work

The net work is the difference between the expansion and compression work plus any pumping work. This quantity is

$$W_t = (U_3 - U_4) - (U_2 - U_1) + (r-1)(P_1 - P_e) J \quad (\text{F-17})$$

9. Thermal Efficiency

The indicated thermal efficiency, defined as the net cycle work divided by the heat of combustion of the fuel consumed, is

$$\eta = \frac{W_t \cdot 100}{\Delta H_c^o \cdot F_o} \quad (\text{F-18})$$

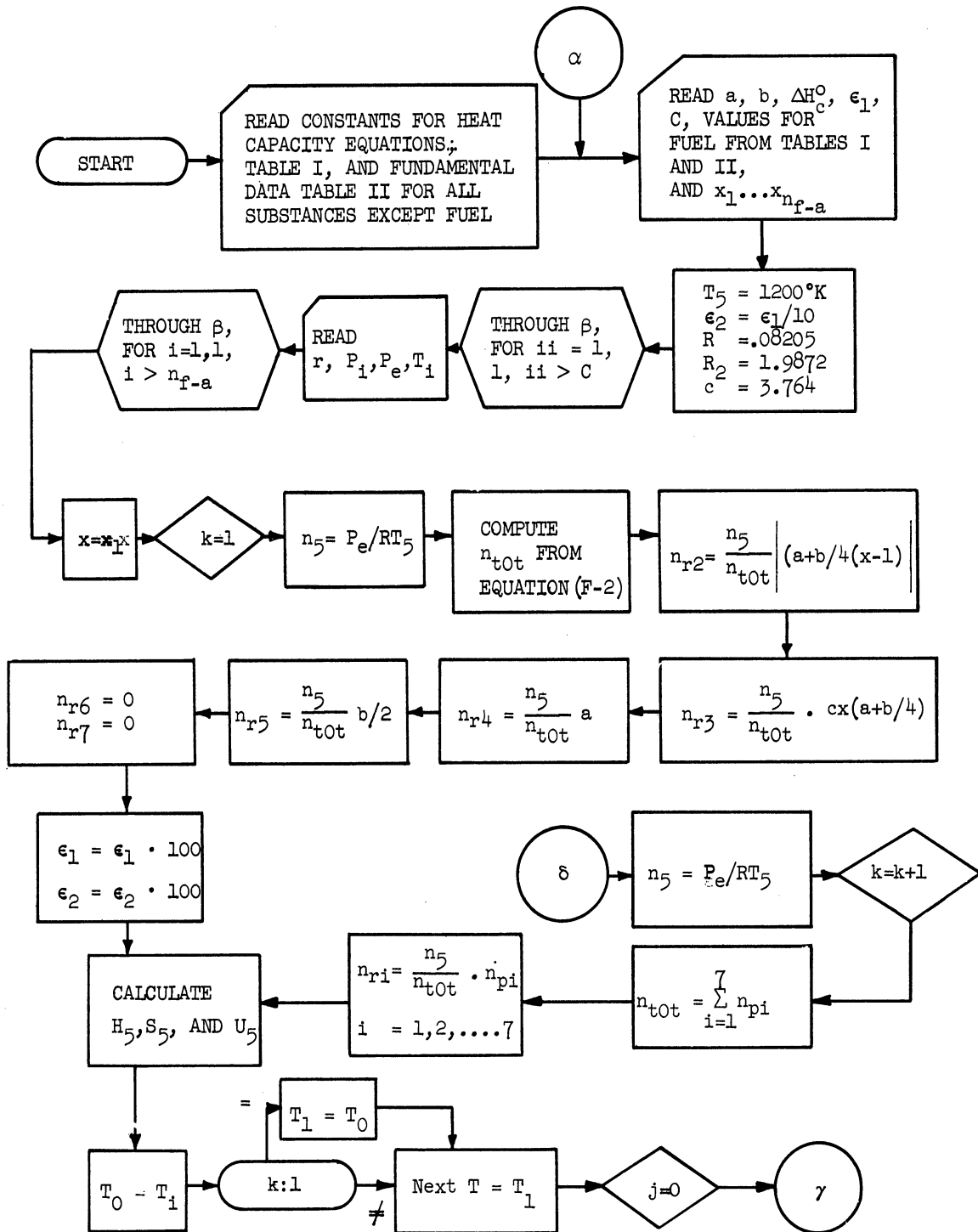
where F_0 are the moles of fuel consumed, calculated in Section 2, and ΔH_c^0 is the higher heating value of the fuel per mole at 298.16°K and one atmosphere pressure.

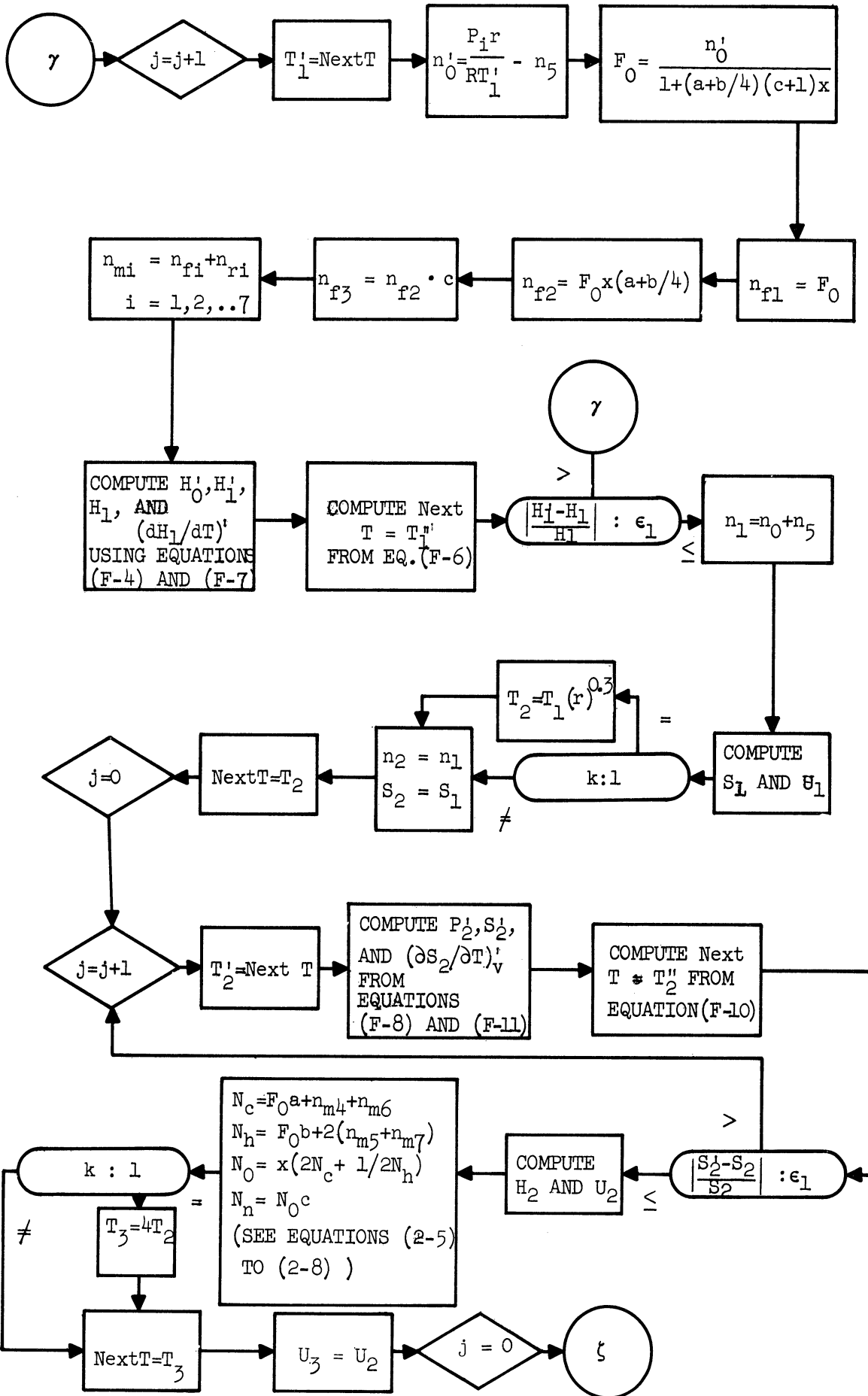
10. Mean Effective Pressure

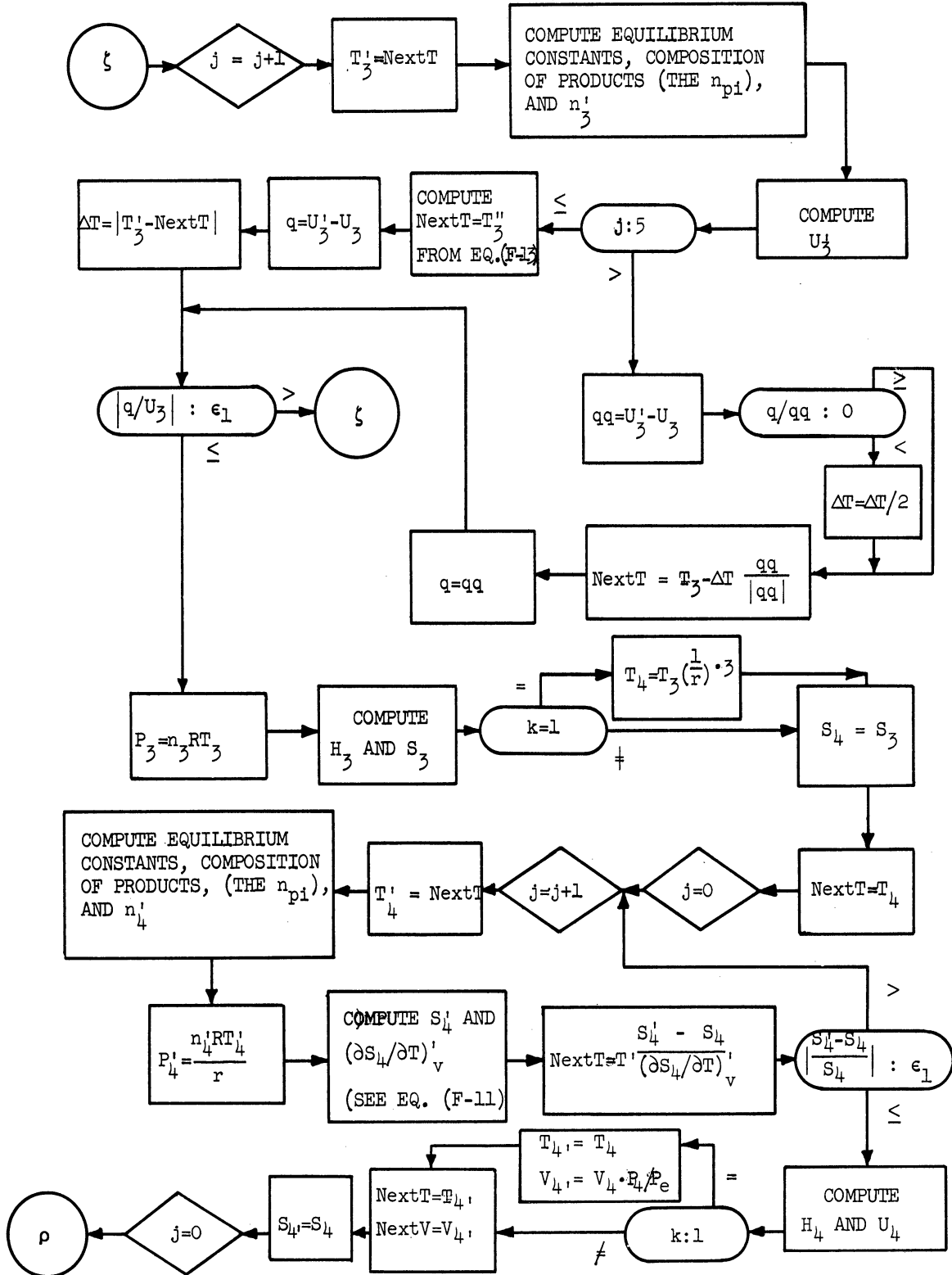
The mean effective pressure is defined as the net work divided by the displacement of the engine. This quantity is

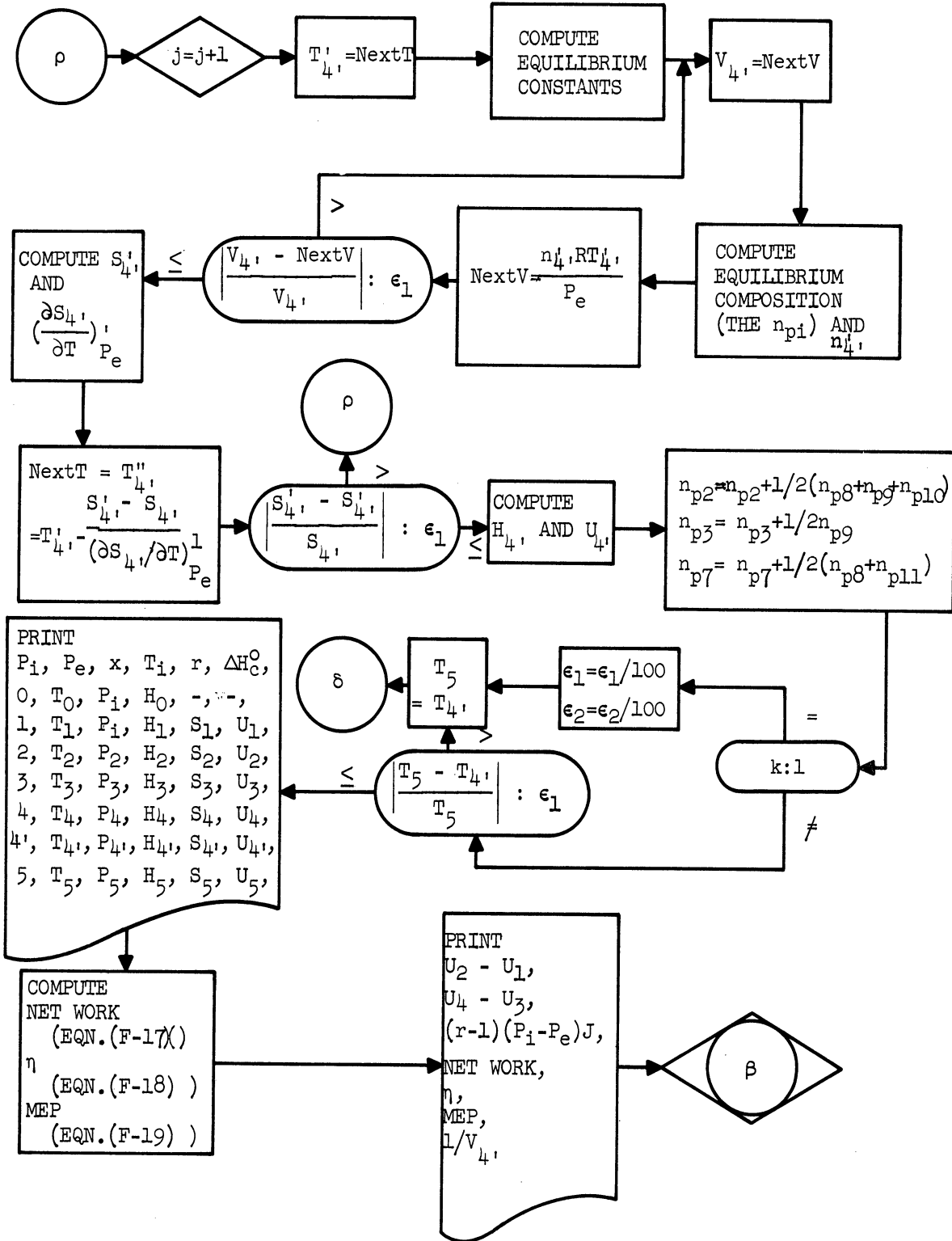
$$\text{mep} = \frac{W_t}{(r - 1)J} \quad (\text{F-19})$$

C. Flow Diagram









D. Computer Program for the Fuel-Air Cycle

The following pages contain a listing of the computer program for the fuel-air cycle engine written in the MAD language for an IBM 704 computer. The external functions ENTHAL., ENTROP., HPRIME., and SPRIME. which compute the enthalpy and entropy as well as their temperature derivatives at constant pressure; EQUIL., the function which calculates the six equilibrium constants; and AMOUNT., the function which calculates the equilibrium composition, are listed in Appendix I and are not repeated here.

The program contains a number of variable names most of which are explained in Appendix I. *Those names pertaining to this program only are:

CARDS number of combinations of compression ratio, manifold pressure and temperature, and exhaust pressure for which computations are to be made: C

NAME whenever NAME is not blank, additional intermediate output is provided.

PH_i an initial estimate of the value of $X = \sqrt{p_{O_2}}$ at Point 'i' (This is used in the computation of the equilibrium composition. In Appendix E it corresponds to X₁ and in the external function AMOUNT. it corresponds to PH.)

R2 gas constant in heat units

VB clearance plus swept volumes, equal to r

VT clearance volume, equal to l

* Symbol after colon is an alternate designation.


```

-----
$COMPILE MAD,PRINT OBJECT,PUNCH OBJECT,EXECUTE,DUMP                                HOTEL001
  DIMENSION AL(87,ALDIM),AH(87,ALDIM),N(11),RE(12),K(11),P(11)
  1,X(50),SPACE(41)
  INTEGER I,I1,J1,J2,RE,CARDS,NFA,NAME
  VECTOR VALUES ALDIM = 2,0,8
  PROGRAM COMMON AL,AH,P,K,R2,NC,NH,NO,NN,R,J1,SPACE,ERROR2
  READ FORMAT SPHT1,NAME,AL(2,1)...AL(11,8),AH(2,1)...AH(11,8)
  VECTOR VALUES SPHT1 = $C6/(4E15.8)*$
START  READ FORMAT FUEL,RE(1)...RE(12),A,B,HCOMB,ERROR1,CARDS,AL(1,1
  1)...AL(1,8),AH(1,1)...AH(1,8)
  VECTOR VALUES FUEL = $12C6/4F10.5,I5/(4E15.8)*$
  PRINT FORMAT DUMP1,AL(1,1)...AL(7,8)
  VECTOR VALUES DUMP1 = $1H1,4HAL =/(8E15.8)*$
  PH3 = 1.
  PH4 = 1.
  T5 = 1200.
  ERROR2 = ERROR1/10.
  R = .08205
  R2 = 1.9872
  C = 3.764
  PRINT FORMAT DUMP2,AH(1,1)...AH(11,8)
  VECTOR VALUES DUMP2 = $1H0,4HAH =/(8E15.8)*$
  READ FORMAT FULAIR,NFA,X(1)...X(NFA)
  VECTOR VALUES FULAIR = $I10/(14F5.4)*$
  THROUGH LOOP1,FOR I1 = 1,1,I1.G.CARDS
  READ FORMAT ENGINE,CR,PMAN,PEXH,TMAN
  VECTOR VALUES ENGINE = $ 6F10.5*$
  PMAN = PMAN/14.696
  PEXH = PEXH/14.696
  TMAN = TMAN/1.8
  VT = 1./61.03
  VB = CR*VT
  THROUGH LOOP1,FOR I=1,1,I.G.NFA
  X = X(I)
  WHENEVER NAME.NE.$          $
  PRINT FORMAT HEAD1,RE(1)...RE(12),PMAN*14.696,PEXH*14.696,X,T
  1MAN*1.8,CR,HCOMB,VT*61.03,(VB-VT)*61.03
  END OF CONDITIONAL
  VECTOR VALUES HEAD1 = $1H1 /1H0,$28,64HIDEALIZED THERMODYN
  1AMIC CYCLE FOR THE INTERNAL COMBUSTION ENGINE/1H0,10HTHIS IS
  2A C6,15H RUN EMPLOYING 2C6,37H AS A FUEL. ADDITIONAL COMMENT
  3S ARE./1H0,9C6/                                1H0,19HMANI
  4FOLD PRESSURE = F10.5,4H PSIS7,18HEXHAUST PRESSURE = F10.5,
  54H PSIS8,16HAIR-FUEL RATIO = F10.5/1H0,22HMANIFOLD TEMPERATUR
  6E = F10.5,2H KS6,19HCOMPRESSION RATIO = F10.5,S11,      17HH
  7 OF COMBUSTION = F10.2,9H CAL/MOLE/1H0,18HCLEARANCE VOLUME =
  8F10.5,7H CU.IN.,S5,14HSWEPT VOLUME = F10.5,7H CU.IN.*$
  T = T5
  J2 = 1
  N5 = PEXH*VT/R/T5
  F5 = N5/(A+B/2.+(A+B/4.)*(C*X+X-1.))
  N(2) = .ABS.(F5*(A+B/4.)*(X-1.))
  N2 = N(2)
  N(3) = F5*(A+B/4.)*C*X
  N3 = N(3)
  N(4) = F5*A
  N(5) = F5*B/2.
  N(6) = 0.
  N(7) = 0.
-----

```

```

-----
ERROR1 = ERROR1*100.
ERROR2 = ERROR2*100.
-----
LOOP  TRANSFER TO LOC1
      J2 = J2 + 1
      T5 = T4P
      T = T5
      N5 = PEXH*VT/R/T
      F5 = N5/(N(2)+N(3)+N(4)+N(5)+N(6)+N(7))
      N(2) = F5*N(2)
      N2 = N(2)
      N(3) = F5*N(3)
      N3 = N(3)
      N(4) = F5*N(4)
      N(5) = F5*N(5)
      N(6) = F5*N(6)
      N(7) = F5*N(7)
-----
LOC1  EXECUTE ENTROP.(2,7,S5,T,N5,PEXH,N)
      EXECUTE ENTHAL.(2,7,H5,T,N)
      U5 = H5 - N5*R2*T
      WHENEVER NAME,NE,$      $
      PRINT FORMAT HEAD2
      PRINT FORMAT OUT,5,T5,PEXH,H5,S5,U5,N5,F5,N(2)...N(7)
      END OF CONDITIONAL
      VECTOR VALUES HEAD2 =$1H0/8HOSTATION,S5,6HT - *K,S5,8HP - ATM
      1.,S5,8HH - CAL.,S7,11HS - CAL./*KS8,8HU - CAL.,S7,11HTOTAL MOL
      2ES,S7,1HF*$
      T0 = TMAN
      J = 0.
      WHENEVER J2.E.1,T1=T0
      NEXTT = T1
-----
LOOP4 T = NEXTT
      J = J + 1.
      N0 = PMAN*VB/R/T-N5
      F0 = N0/(1. +X*(A+B/4.)+C*X*(A+B/4.))
      N(1) = F0
      N(2) = F0*X*(A+B/4.)
      N(3) = N(2)*C
      EXECUTE ENTHAL.(1,3,H0,TMAN,N)
      EXECUTE HPRIME.(1,3,HPRIM,T,N)
      HPRIM = HPRIM - H0/T*(1. + N5/N0)
      H1 = H0 + U5 + PMAN*VT*24.2179
      N(2) = N(2) + N2
      N(3) = N(3) + N3
      EXECUTE ENTHAL.(1,7,HMIX,1,N)
      NEXTT = T - (HMIX-H1)/HPRIM
      WHENEVER .ABS.((HMIX-H1)/H1).G.ERROR1,TRANSFER TO LOOP4
      T1 = T
      N1 = N0 + N5
      EXECUTE ENTROP.(1,7,S1,T,N1,PMAN,N)
      U1 = H1-N1*R2*T
      WHENEVER NAME,NE,$      $
      PRINT FORMAT OUT, 0,T0,PMAN,H0,0,0,N0,F0,F0,F0*X*(A+B/4.),F0*
      1X*C*(A+B/4.),J
      PRINT FORMAT OUT,1,T1,PMAN,H1,S1,U1,N1,0,N(1)...N(7),J
      END OF CONDITIONAL
      WHENEVER J2.E.1, T2 = T1*CR,P..3
      N2 = N1
      J = 0.
-----

```

```

NEXTT = T2
LOOP5  T = NEXTT
      J = J + 1.
      P2 = T*CR*PMAN/T1
      EXECUTE ENTROP.(1,7,S2,T,N2,P2,N)
      EXECUTE SPRIME.(1,7,SPRIM,T,N)
      SPRIM = SPRIM - N2*R2/T
      NEXTT = T - (S2 - S1)/SPRIM
      WHENEVER .ABS.((S2-S1)/S1).G.ERROR1,TRANSFER TO LOOP5
      EXECUTE ENTHAL.(1,7,H2,T,N)
      T2 = T
      U2 = H2 - N2*R2*T
      WHENEVER NAME.NE.$ $
      PRINT FORMAT OUT, 2,T2,P2,H2,S2,U2,N2,0.,N(1)...N(7),J
      END OF CONDITIONAL
      NC = F0*A + N(4) + N(6)
      NH = F0*B + (N(5) + N(7))*2.
      NO = (2.*NC+.5*NH)*X
      NN = NO*C
      WHENEVER J2.E.1,T3 = 4.*T2
      NEXTT = T3
      J = 0.
LOOP6  T = NEXTT
      J = J + 1.
      EXECUTE EQUIL.(T)
      EXECUTE AMOUNT.(T,VT,N3,PH3,N)
      EXECUTE ENTHAL.(2,11,H3,T,N)
      U3 = H3 - N3*R2*T
      WHENEVER J.G.5.,TRANSFER TO HALFIN
      EXECUTE HPRIME.(2,11,HPRIM,T,N)
      UPRIM = HPRIM - N3*R2
      Q = (U2-U3)/T/UPRIM
      WHENEVER .ABS.(Q).L..1
      NEXTT = T*EXP.(Q)
      OTHERWISE
      NEXTT = T + 100.*Q/.ABS.(Q)
      END OF CONDITIONAL
      DIF = .ABS.(T - NEXTT)
      SAVE = DIF
      WHENEVER .ABS.((U3-U2)/U2).G.ERROR1,TRANSFER TO LOOP6
      TRANSFER TO END
HALFIN QQ = U2 - U3
      WHENEVER Q/QQ .L.0.,DIF = DIF/2.
      NEXTT = T + DIF*QQ/.ABS.(QQ)
      Q = QQ
      WHENEVER .ABS.((U3-U2)/U2).G.ERROR1,TRANSFER TO LOOP6
END    P3 = N3*R*T/VT
      T3 = T
      EXECUTE ENTROP.(2,11,S3,T,N3,P3,N)
      WHENEVER NAME.NE.$ $
      PRINT FORMAT OUT,3,T3,P3,H3,S3,U3,N3,0.,N(1)...N(11),J,P(1)..
      1.P(11),SAVE
      END OF CONDITIONAL
      WHENEVER J2.E.1,T4 = T3*(1./CR).P..3
      NEXTT = T4
      J = 0.
LOOP7  T = NEXTT
      J = J + 1.
```

```
EXECUTE EQUIL.(T)
EXECUTE AMOUNT.(T,VB,N4,PH4,N)
P4 = N4*R*T/VB
EXECUTE ENTROP.(2,11,S4,T,N4,P4,N)
EXECUTE SPRIME.(2,11,SPRIM,T,N)
SPRIM = SPRIM - N4*R2/T
NEXTT = T - (S4-S3)/SPRIM
WHENEVER .ABS.((S4-S3)/S3).G.ERROR1,TRANSFER TO LOOP7
T4 = T
EXECUTE ENTHAL.(2,11,H4,T,N)
U4 = H4 - N4*R2*T
WHENEVER NAME.NE.$ $
PRINT FORMAT OUT,4,T4,P4,H4,S4,U4,N4,0.,N(1)...N(11),J,P(1)..
1.P(11)
END OF CONDITIONAL
WHENEVER J2 .E.1
T4P = T4
PH4P = PH4
V4P = N4*R*T4P/PEXH
END OF CONDITIONAL
NEXTT = T4P
J = 0.
NEXTV = V4P
LOOP8 T = NEXTT
J = J + 1.
EXECUTE EQUIL.(T)
J3 = 0.
BACK V = NEXTV
J3 =J3 + 1.
EXECUTE AMOUNT.(T,V,N4P,PH4P,N)
NEXTV = N4P*R*T/PEXH
WHENEVER .ABS.((NEXTV-V )/V ).G.ERROR1,TRANSFER TO BACK
EXECUTE ENTROP.(2,11,S4P,T,N4P,PEXH,N)
EXECUTE SPRIME.(2,11,SPRIM,T,N)
NEXTT = T - (S4P-S4)/SPRIM
WHENEVER .ABS.((S4P-S4)/S4).G.ERROR1,TRANSFER TO LOOP8
T4P = T
V4P = V
P4P = N4P*R*T/V
EXECUTE ENTHAL.(2,11,H4P,T,N)
U4P = H4P - N4P*R2*T
WHENEVER NAME.NE.$ $
PRINT FORMAT OUT,45, T4P,P4P, H4P,S4P,U4P,N4P,0.,N(1)...N(11)
1,J,P(1)...P(11),J3
PRINT FORMAT NOTICE ,J2,.ABS.(T5-T4P),NC,NH,NO,NN,VT,VB
END OF CONDITIONAL
VECTOR VALUES NOTICE = $1H0,23HAFTER ITERATION NUMBER I2,51H
1 THE ABSOLUTE DIFFERENCE BETWEEN T5 AND T4PRIME IS F10.5/1H0,
246HTHE MOLE ATOMS OF THE FOLLOWING SUBSTANCES ARE/9H CARBON =
3E15.8,11H HYDROGEN = E15.8,9H OXYGEN = E15.8,11H NITROGEN = E
415.8/5H VT = E15.8,12H L. AND VB = E15.8,3H L.*$
N(2) = N(2)+.5*(N(8)+N(9)+N(10))
N(3) = N(3) + .5*N(9)
N(7) = N(7) + .5*(N(8)+N(11))
WHENEVER J2 .E. 1
ERROR1 = ERROR1/100.
ERROR2 = ERROR2/100.
TRANSFER TO LOOP
```

```
END OF CONDITIONAL
WHENEVER .ABS.((T5-T4P)/T5).G.ERROR1,TRANSFER TO LOOP
PRINT FORMAT HEAD1,RE(1)...RE(12),PMAN*14.696,PEXH*14.696,X,T
1MAN*1.8,CR,HCOMB,VT*61.03,(VB-VT)*61.03
PRINT FORMAT HEAD3
VECTOR VALUES HEAD3 = $1H0,S18,36H SUMMARY OF RESULTS IN ENGLI
ISH SYSTEM/1H0, 7HSTATION,S5,6HT - *R,S5,8HP - PSI.,S5,
2      8HH - BTU.,S5,11HS - BTU./*R,S6,8HU - CAL.*$
PRINT FORMAT SUMMRY,0,T0*1.8,PMAN*14.696,H0/252.161,0..0..1,T
11*1.8,PMAN*14.696,H1/252.161,S1/252.161/1.8,U1/252.161,2,T2*1
2.8,P2*14.696,H2/252.161,S2/252.161/1.8,U2/252.161,3,T3*1.8,P3
3*14.696,H3/252.161,S3/252.161/1.8,U3/252.161,4,T4*1.8,P4*14.6
496,H4/252.161,S4/252.161/1.8,U4/252.161,45,T4P*1.8,P4P*14.696
5,H4P/252.161,S4P/252.161/1.8,U4P/252.161,
65,T5*1.8,PEXH*14.696,H5/252.161,S5/252.161/1.8,U5/252.161
VECTOR VALUES SUMMRY = $1H ,I4,S6,2F10.3,3F15.8*$
VECTOR VALUES OUT = $1H0,I4,S6,2F10.5,5E17.8/(6E20.8)*$
WORK12 = (U2-U1)/252.161
WORK43 = (U3-U4)/252.161
WPUMP = ((VB-VT)*(PMAN-PEXH))*24.2179/252.161
WNET = WORK43-WORK12+WPUMP
PRINT FORMAT RESULT,WORK12,WORK43,WPUMP,WNET,WNET/HCOMB/FO*25
1216.1,WNET*14.696/(VB-VT)/24.2179*252.161
VECTOR VALUES RESULT = $1H0/1H0,S10,7HRESULTS/1H+,S10,7HRESUL
1TS/1H0,31HUNDER THE FOLLOWING ASSUMPTIONS/40H 1) ISENTROPIC C
2OMPRESSION AND EXPANSION/30H 2) CONSTANT VOLUME COMBUSTION/20
3H 3) NO HEAT TRANSFER/1H0/1H0/
4 S6,30HCOMPRESSION WORK = E15.8,4H BTU/
5 S6,30HEXPANSION WORK = E15.8,4H BTU/
6 S6,30HPUMPING WORK = E15.8,4H BTU/
7 S6,30HNET CYCLE WORK = E15.8,4H BTU/
8 S6,30HINDICATED THERMAL EFFICIENCY= E15.8,8H PERCENT/
9 S6,30HMEAN EFFECTIVE PRESSURE = E15.8,4H PSI*$
LOOP1 PRINT FORMAT RESULL,VT/V4P,FO*(A+B/4.)*(C*X+X)/(VB-VT)*24.464
1028
VECTOR VALUES RESULT = $
1 S6,30HRESIDUAL WEIGHT FRACTION = E15.8,4H = F/
2 S6,30HVOLUMETRIC EFFICIENCY = E15.8,25H BASED ON 1 ATM
3. AND 25*K *$
TRANSFER TO START
END OF PROGRAM
```

APPENDIX G

ITERATIVE METHODS FOR SOLUTION OF IMPLICIT FUNCTIONS OF A SINGLE VARIABLE

A. General

Consider a function of one variable,

$$f(x) = 0 \tag{G-1}$$

It is desired to find the value of x which will satisfy (G-1).

Assuming an initial value, x_1 , in general the function $f(x_1)$ will not be equal to zero and must be corrected by some amount, $\Delta f(x_1)$, in order to equate it to zero. That is

$$f(x_1) + \Delta f(x_1) = 0 \tag{G-2}$$

It remains to find a correction to x_1 , Δx_1 , giving a new value x_2 which is closer to the correct answer.

B. Methods of Solution

1. Newton's Method

Newton's method proposes the following relation for determining x_2 :

$$\Delta f(x_1) \approx \left(\frac{df(x)}{dx}\right)_1 \cdot \Delta x_1 = f'(x_1) \cdot \Delta x_1 \tag{G-3}$$

Substituting (G-3) in (G-2),

$$x_2 = x_1 + \Delta x_1 = x_1 - \frac{f(x_1)}{f'(x_1)} \tag{G-4}$$

In general, for the i-th trial,

$$x_{i+1} = x_i - \frac{f(x_i)}{f'(x_i)}$$

The iteration is repeated until Δx_i is less than some prescribed value.

2. Half-Interval Method

The Half-Interval method proposes that an arbitrary initial step size, Δx_1 , be used between successive trials for x . This step size is then reduced as the correct answer is approached. The following relation is used to find x_2 :

$$x_2 = x_1 + \Delta x_1 \cdot \frac{-f(x_1)}{|f(x_1)|} \quad (G-5)$$

If $f(x_2)$ is different in sign than $f(x_1)$, $\Delta x_2 = \frac{\Delta x_1}{2}$. The negative sign is used in (G-5) since $f(x_2) > f(x_1)$ whenever $x_2 > x_1$, for all the functions considered.

In general, for the i-th trial,

$$x_{i+1} = x_i - \Delta x_i \cdot \frac{f(x_i)}{|f(x_i)|} \quad (G-6)$$

where $\Delta x_i = \frac{\Delta x_{i-1}}{2}$ whenever the ratio $\frac{f(x_{i-1})}{f(x_i)}$ is negative.

The iteration is repeated until Δx_i is less than some prescribed value.

APPENDIX H

CALCULATION OF THE RATE OF RADIANT HEAT TRANSFER

The rate of radiant heat transfer in an internal combustion engine will be calculated by the method of Hottel⁽⁸⁷⁾ and McAdams⁽⁶⁸⁾, Chapter IV. A sample calculation will be made to estimate the magnitude of the radiant heat transfer in a 10 to 1 compression ratio engine which will be taken to be representative of a present day high compression engine. This engine is assumed to run on the fuel-air-cycle (see Chapter III) which postulates an adiabatic combustion process at top center. If the radiation is calculated based on the state of the working fluid after this adiabatic combustion, the amount of radiant heat transfer found is expected to be considerably higher than that found in an actual engine where the maximum temperatures and pressures are appreciably lower. A normal combustion process is assumed for which the principal emitters are CO₂ and H₂O.

The following notation is used in connection with the calculations; the figure numbers refer to figures in McAdams:

- ϵ_{gc} emissivity of CO₂ from Figure 4-13
- ϵ_{gw} emissivity of H₂O from Figure 4-15
- $\Delta\epsilon$ correction for overlapping emission spectra from Figure 4-17
- C_c pressure correction factor for CO₂ from Figure 4-16
- C_w pressure correction factor for H₂O from Figure 4-14
- ϵ_g emissivity of gas mixture
- a absorbtivity of wall
- ϵ' factor to account for the non-black body character of the enclosure

For the transfer of heat by radiation from hot gases whose principal emitters are CO₂ and H₂O, the following formula applies:

$$q/A = 1723 \epsilon' \left[\epsilon_g \left(\frac{T_g}{100} \right)^4 - \left(\frac{T_w}{100} \right)^4 \right] \text{ Btu/ft}^2\text{-sec} \quad (\text{H-1})$$

where

$$\begin{aligned} \epsilon_g &= \epsilon_{gc} C_c + \epsilon_{gw} C_w - \Delta\epsilon_g \\ a &= \epsilon_{sc} C_c + \epsilon_{sw} C_w - \Delta\epsilon_s \end{aligned}$$

For the 10 to 1 compression ratio engine running on a stoichiometric mixture of iso-octane and air the following values were obtained from a fuel-air cycle analysis:

maximum cycle gas temperature	5000°R
maximum cycle gas pressure	100 atm
mole fraction of CO ₂	0.1
mole fraction of H ₂ O	0.13

It is recognized that in an actual engine during normal combustion the pressures are never as high as those predicted by the fuel-air cycle analysis. A figure of 67 atm or 1000 psia seemed to be a more reasonable value for the pressure than 100 atm and thus was used in the calculations.

An average hydraulic radius is required for the calculations for which Hottel suggests

$$L = 3.4 \frac{\text{volume of gas}}{\text{area of boundary surface}} \text{ ft}$$

For an engine with a compact combustion chamber such as that used for the calculations reported in Chapter VII, 0.075 ft is an appropriate value.

Following the method of McAdams,

$$p_c L = 0.5 \text{ atm-ft}$$

$$p_w L = 0.65 \text{ atm-ft}$$

where p_c and p_w are the partial pressures of CO_2 and H_2O respectively. From the appropriate figures in Reference 68,

$$\epsilon_{gw} = 0.05$$

$$\epsilon_{gc} = 0.037$$

$$\Delta\epsilon_g = 0.046$$

$$a \approx 1.0$$

$$\left. \begin{array}{l} C_c = 1.75 \\ C_w = 3.75 \end{array} \right\} \text{ by extrapolation}$$

Then

$$\epsilon_g = 0.21$$

for $\epsilon_{\text{surface}} = 0.9$, $\epsilon' = 0.94$ (from Reference 87)

Substitution these values in (H-1)

$$q/A = 70 \text{ Btu/ft}^2\text{-sec}$$

The convective component of heat transfer can be calculated from Equation (5-18). If the speed is 3000 rpm and the stroke is 4 in., the average piston speed, w , is approximately 10 m/sec. For a wall temperature of 300°F,

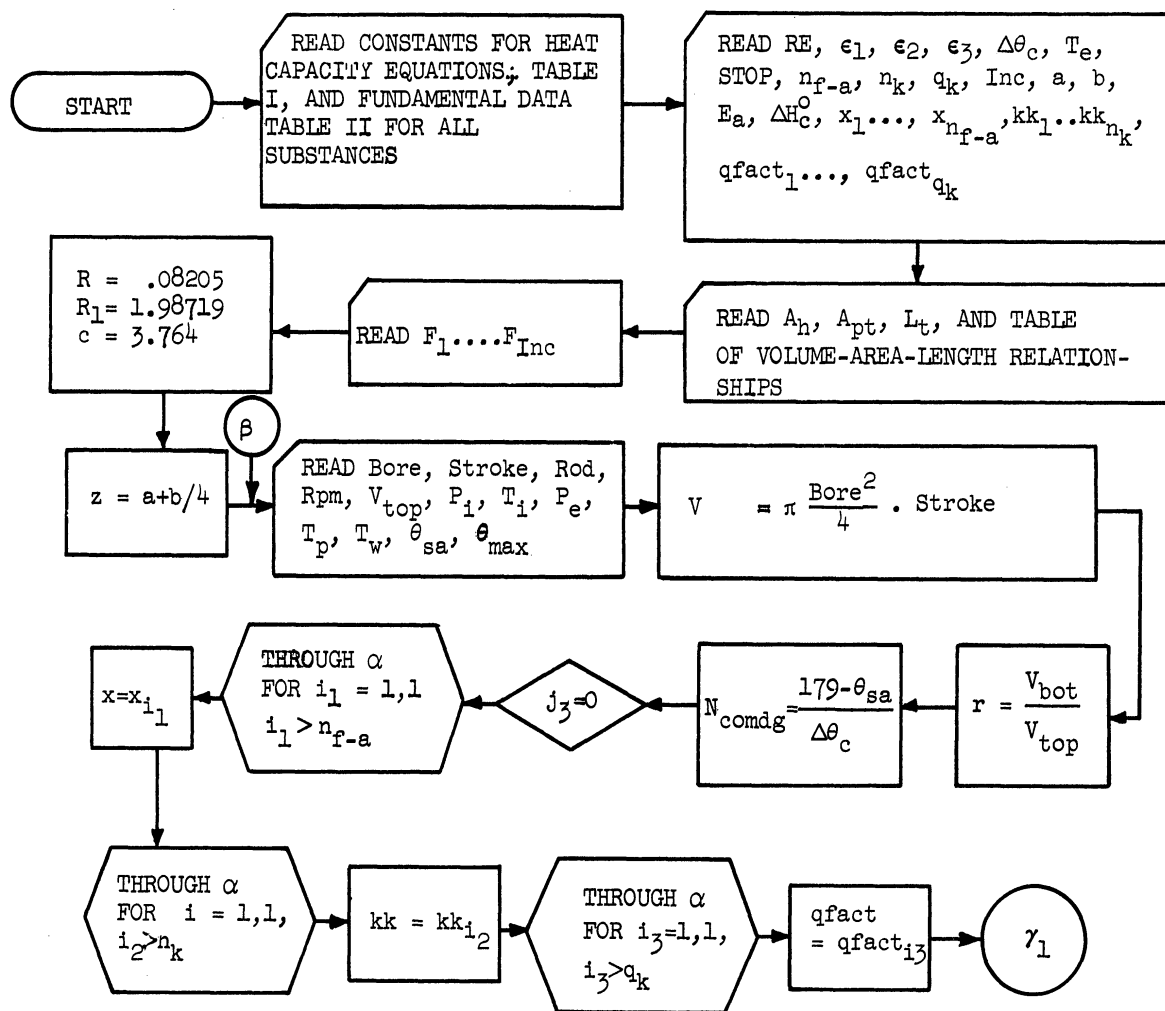
$$q/A = 490 \text{ Btu/ft}^2\text{-sec}$$

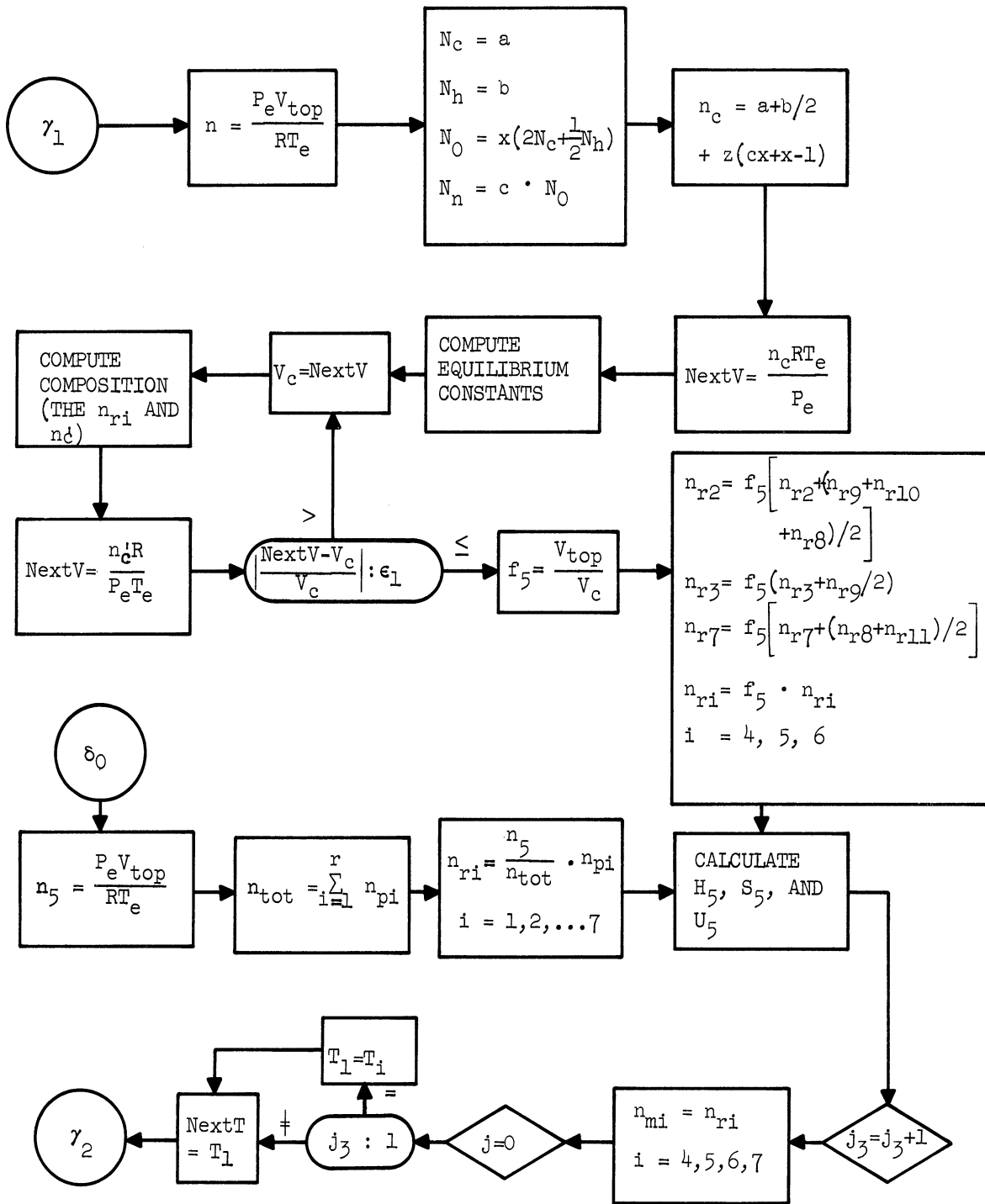
The total heat transfer is 560 Btu/ft²-sec of which that due to radiation is about 12 percent. Because the assumptions and extrapolations used for this calculation always tended to overestimate the amount of radiant heat transfer it is felt that the percentage found is probably too high.

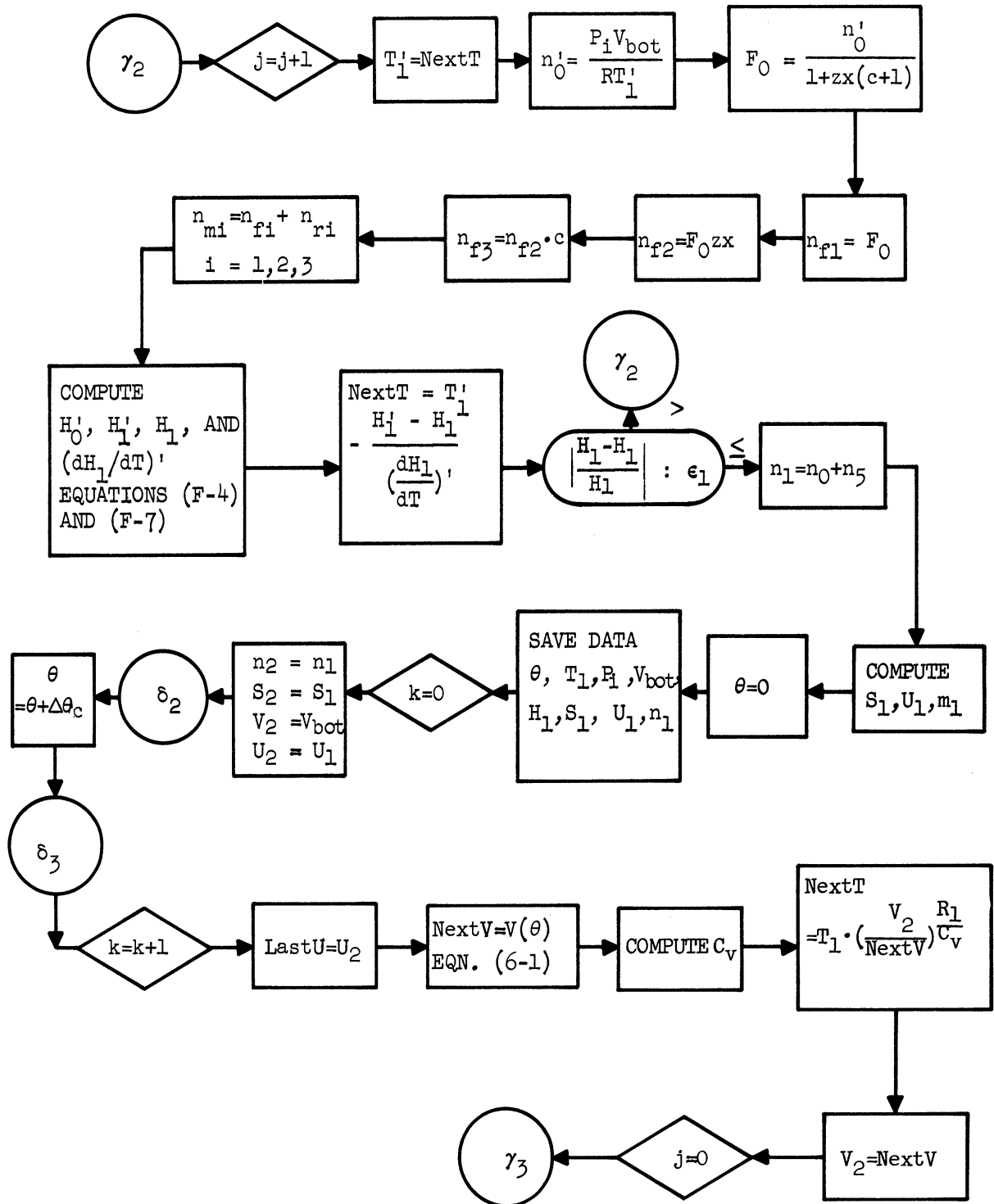
APPENDIX I

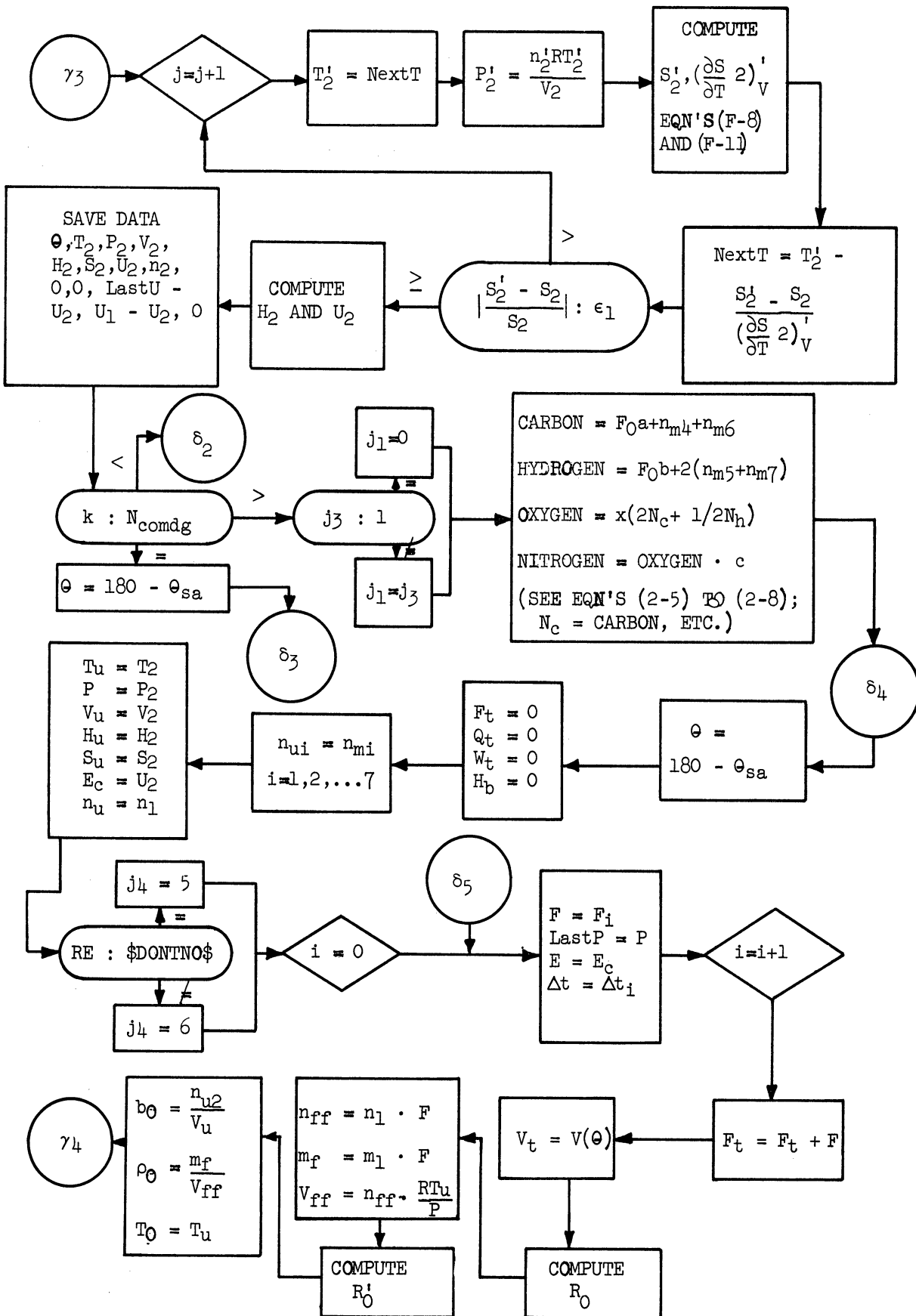
FLOW DIAGRAM AND PROGRAM LISTING FOR THE ANALYTICAL MODEL

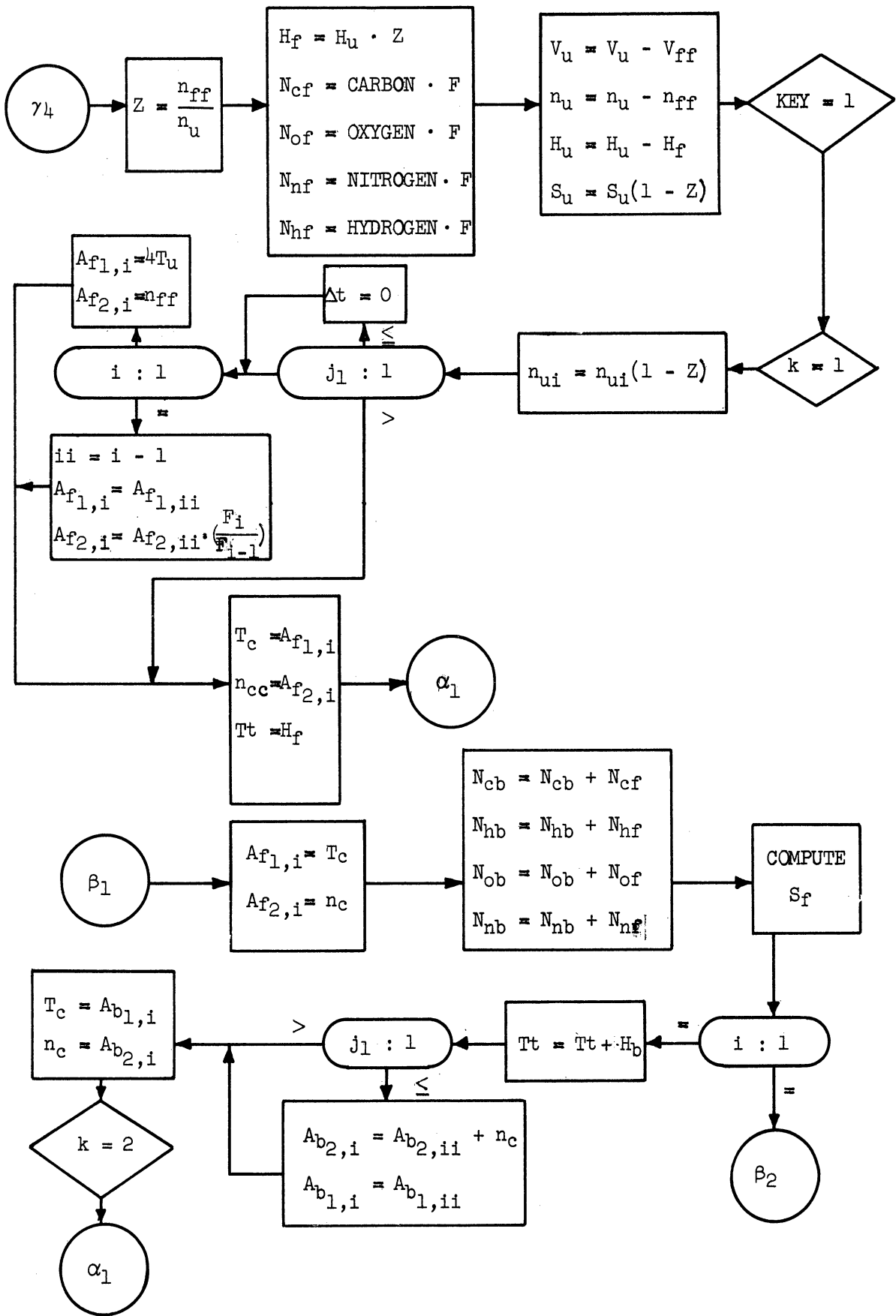
A. Flow Diagram

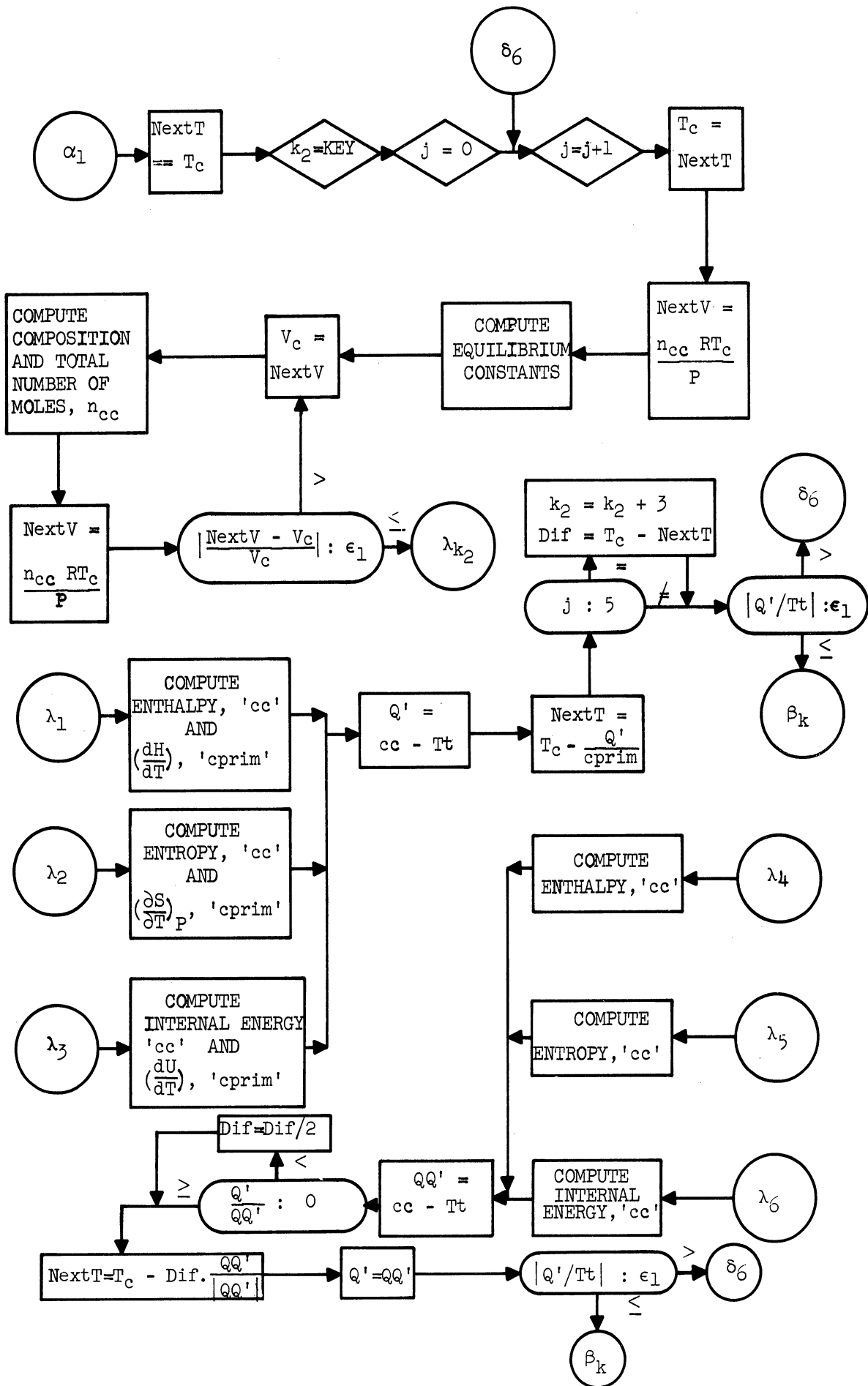


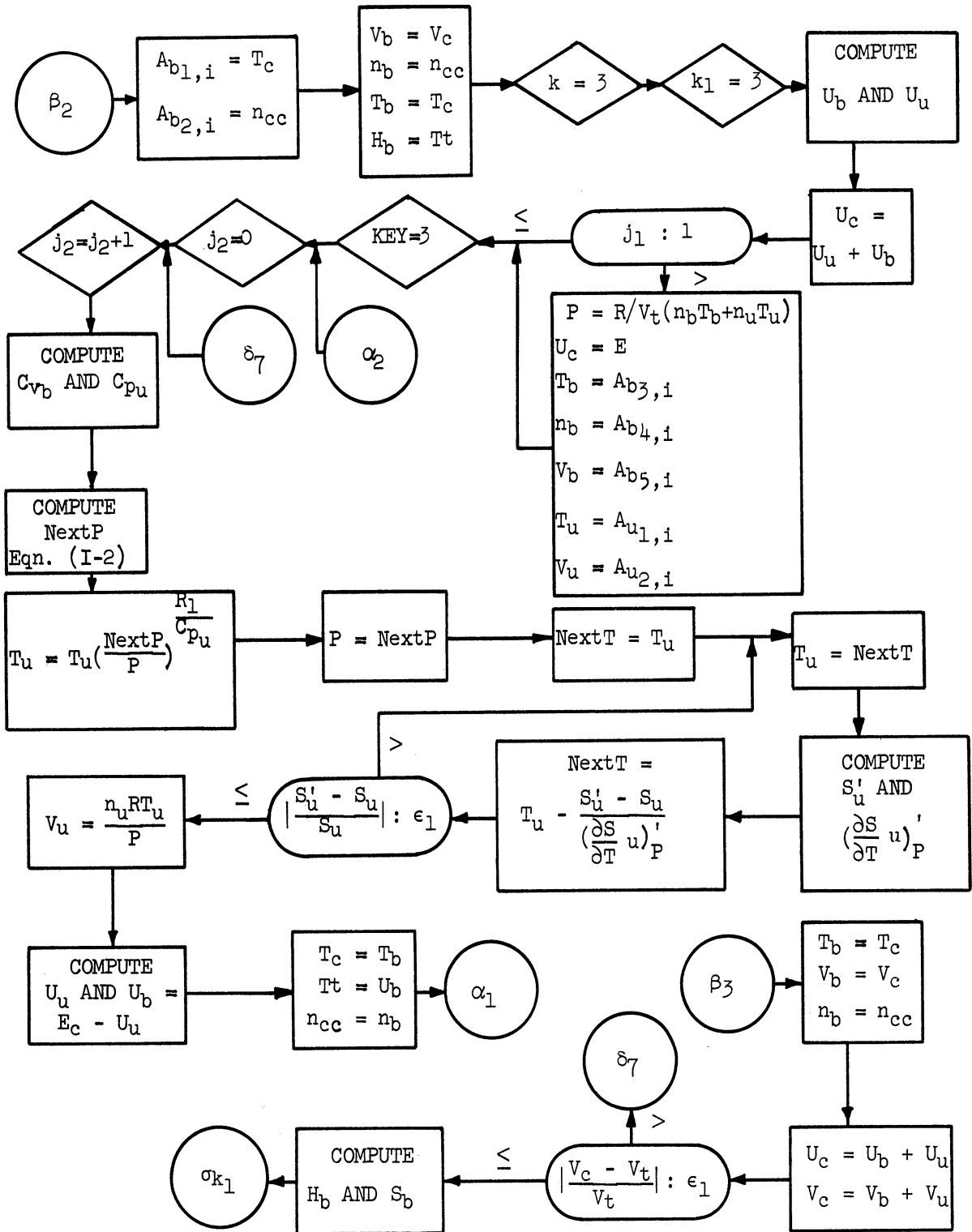


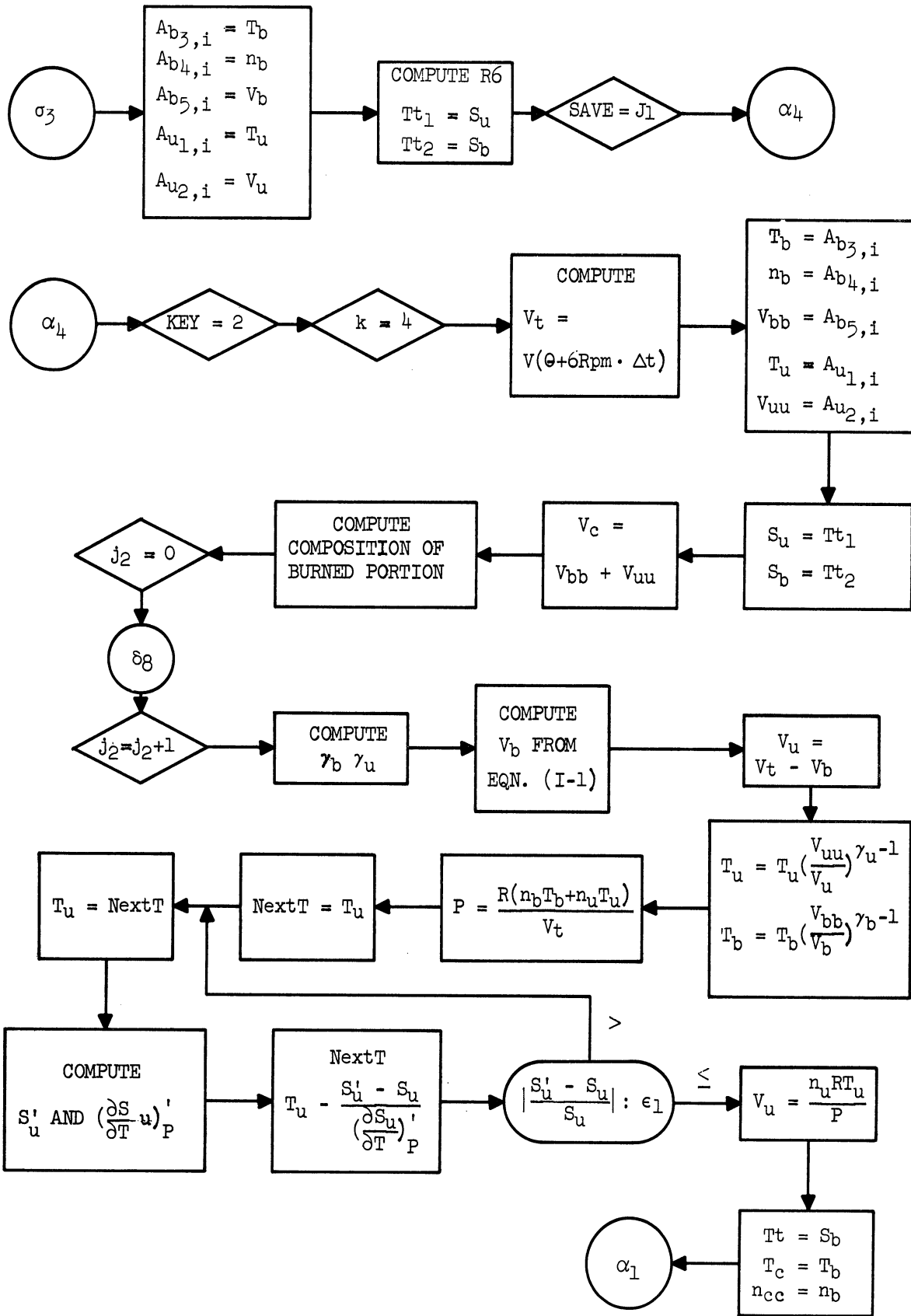


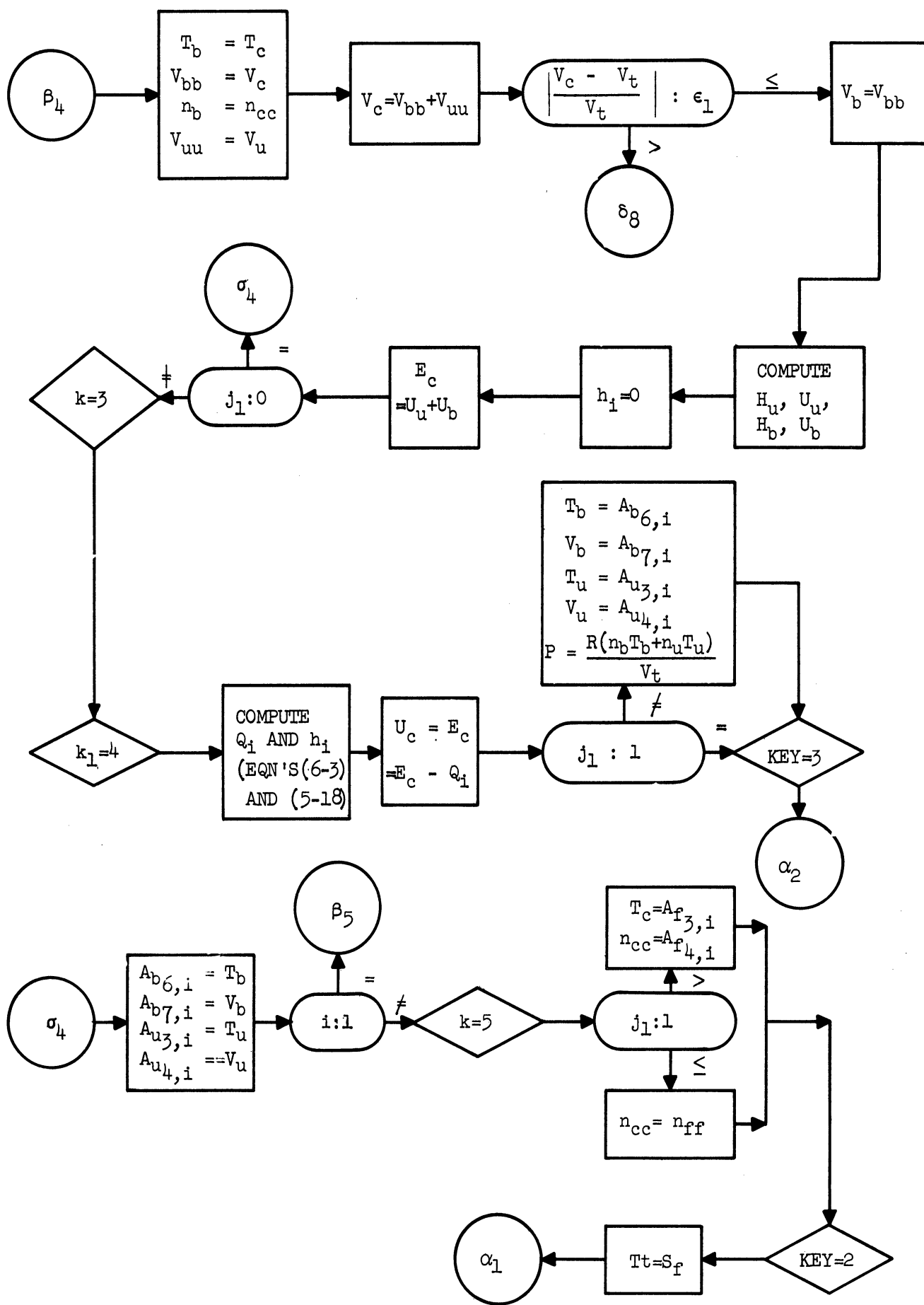


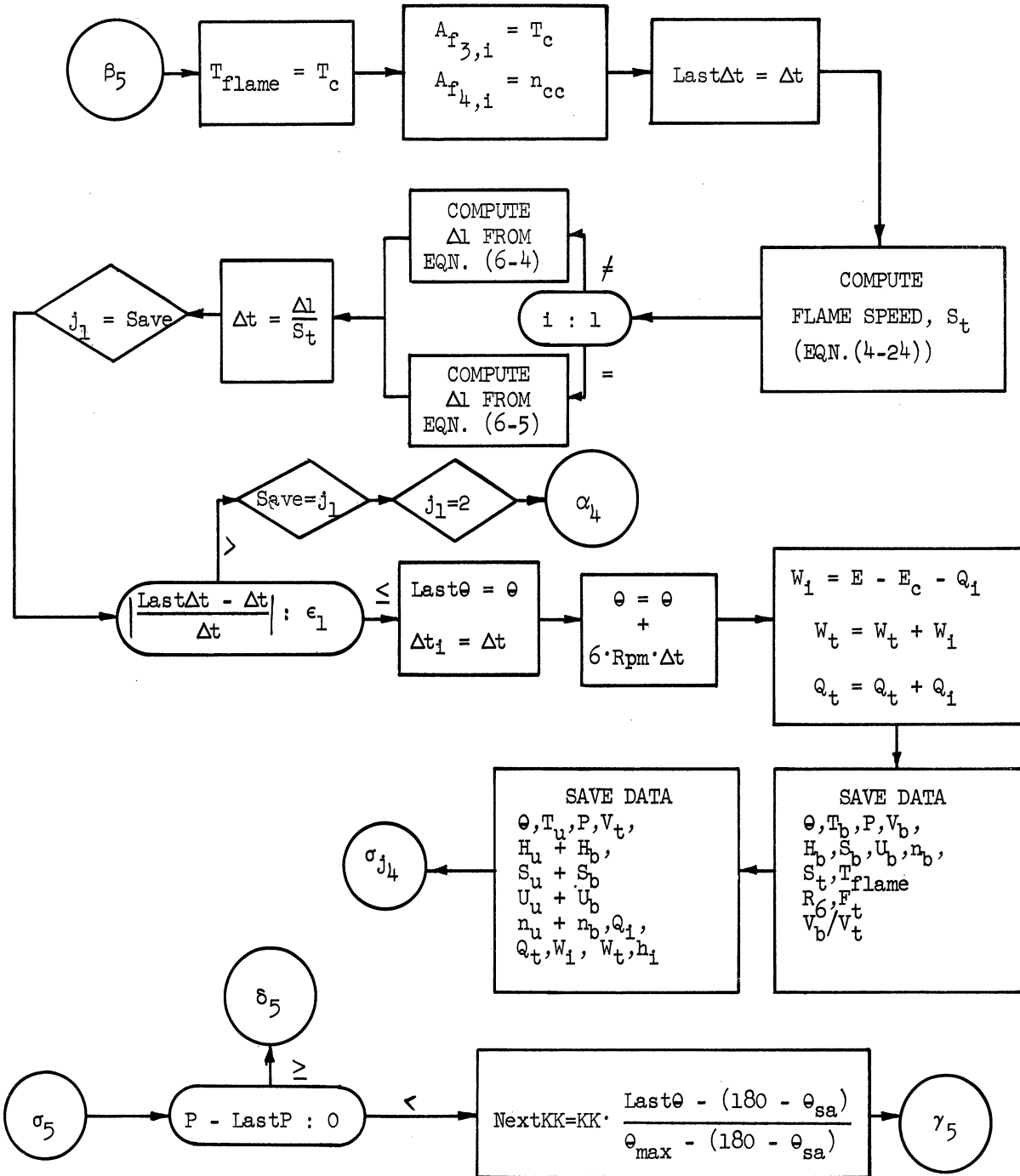


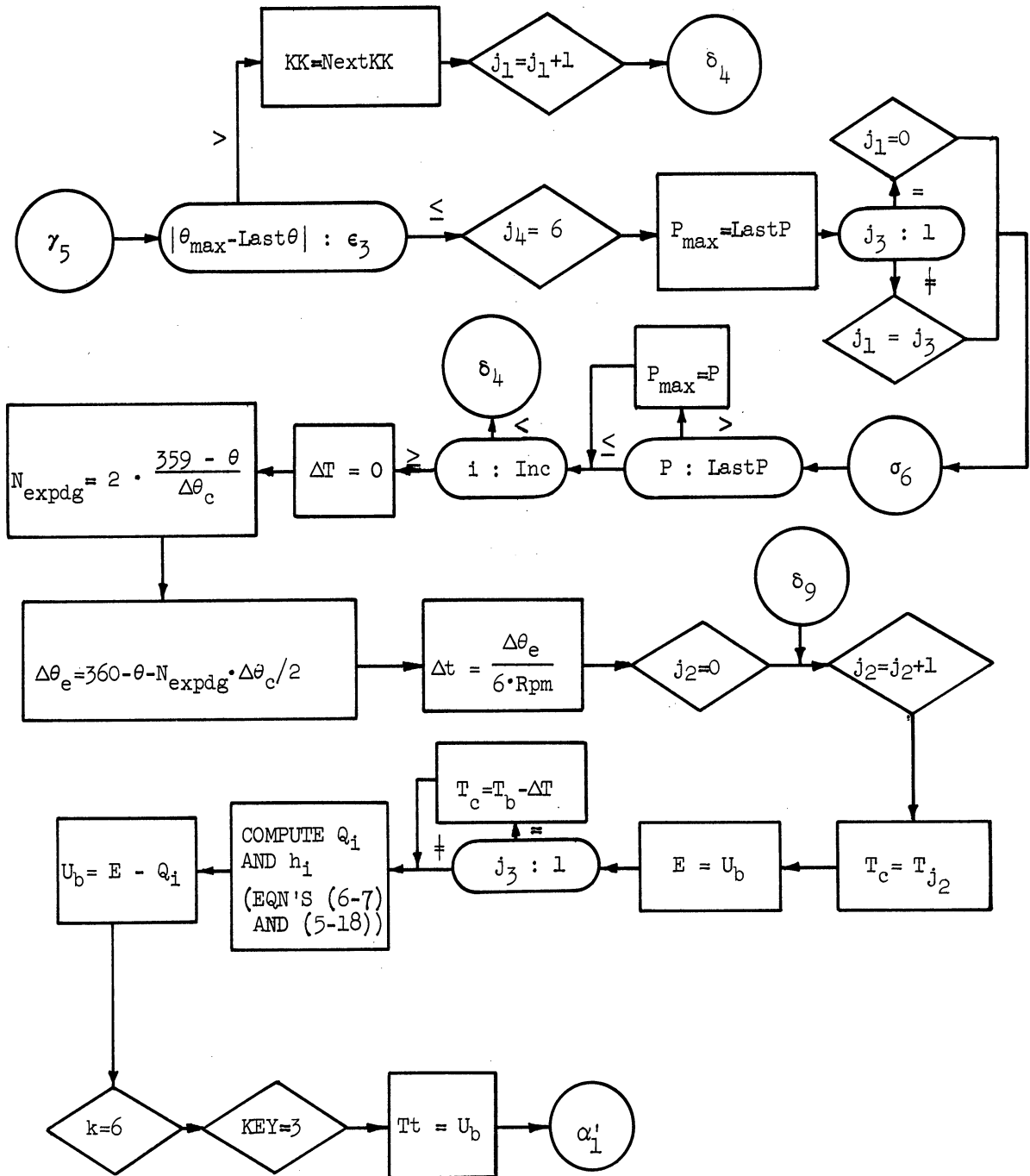


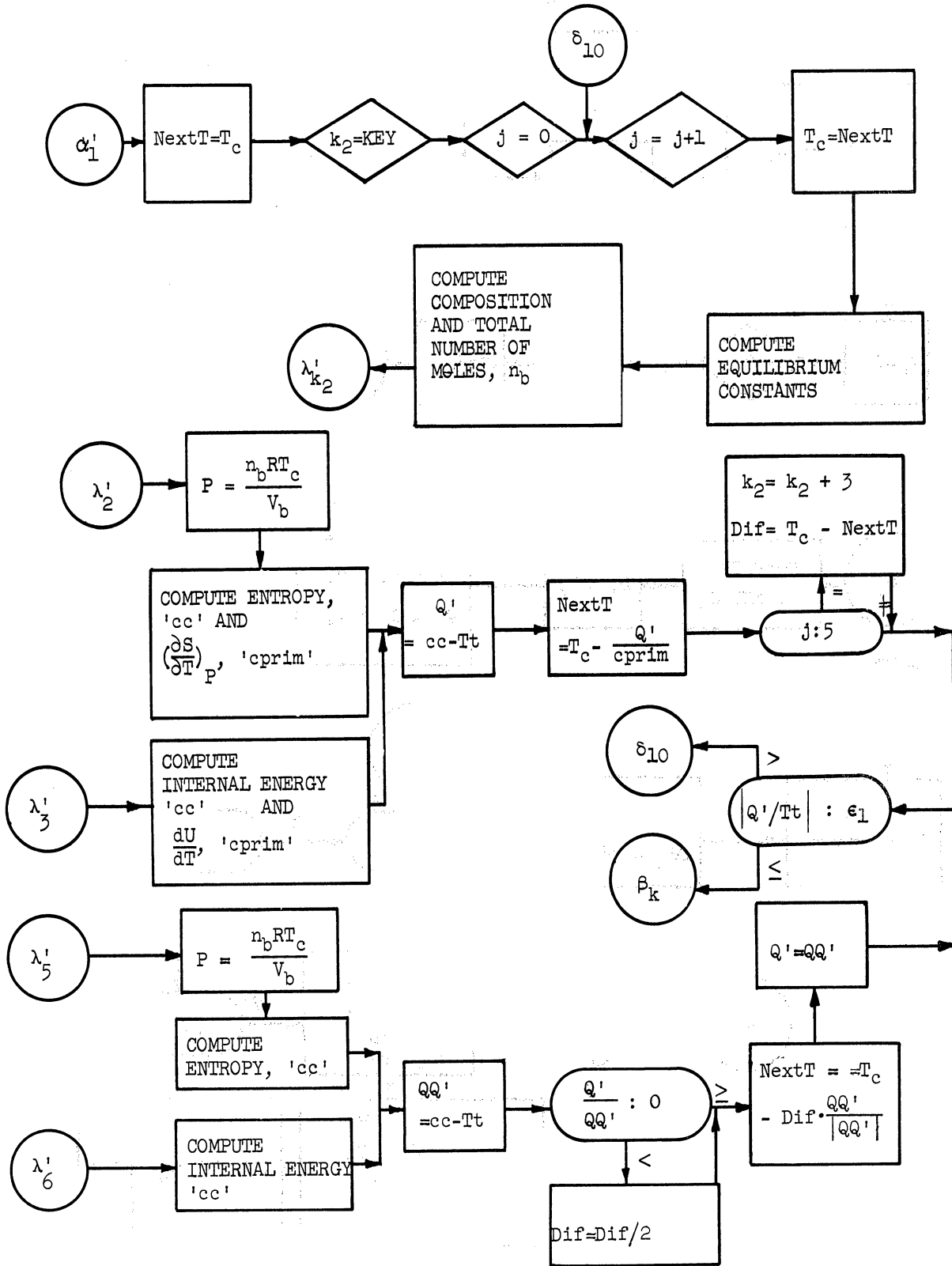


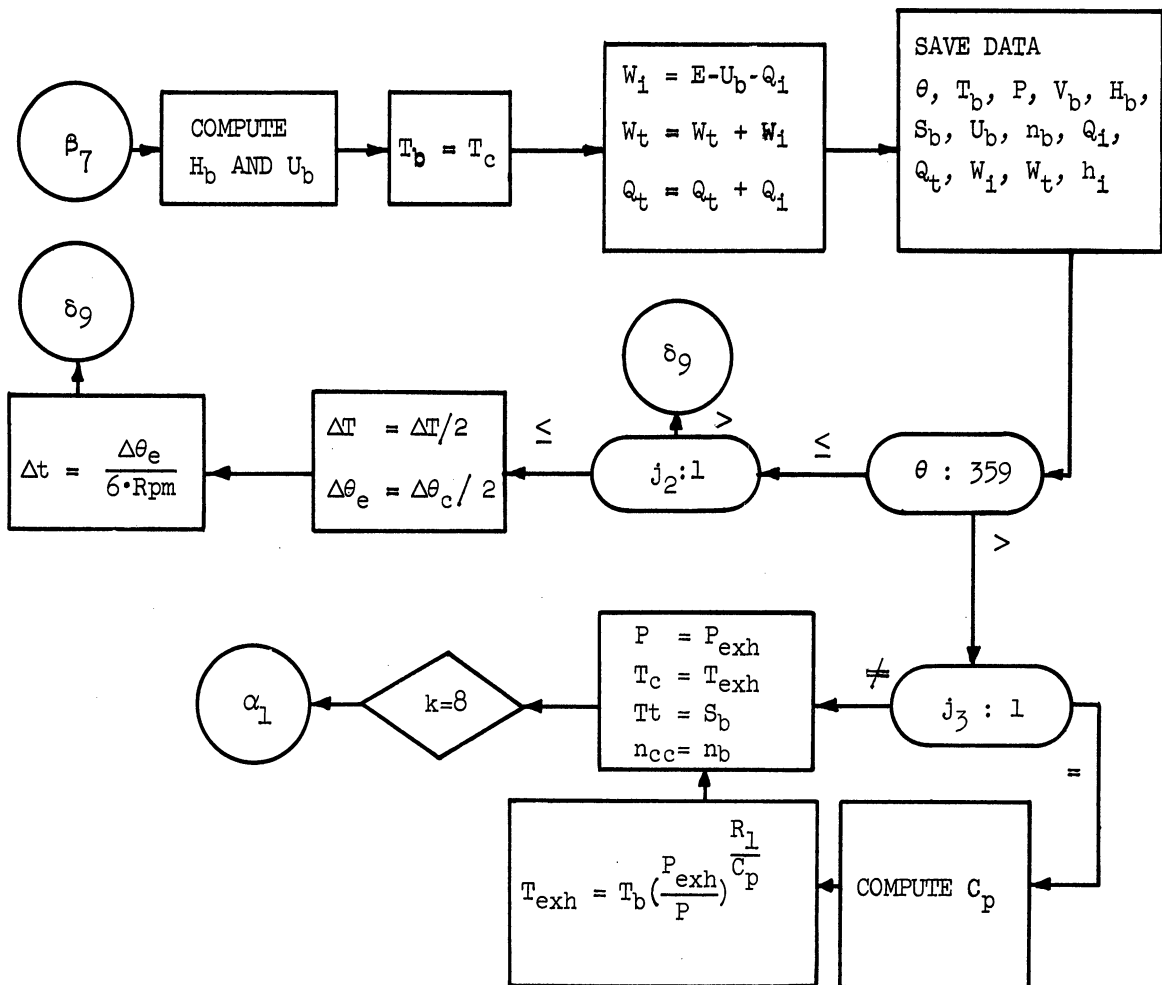
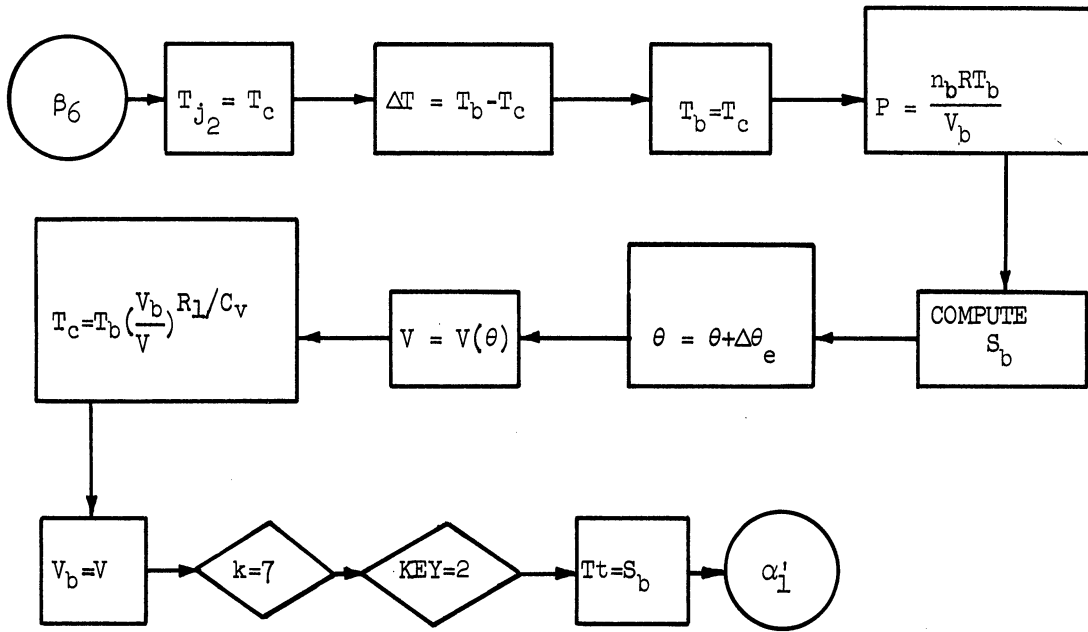


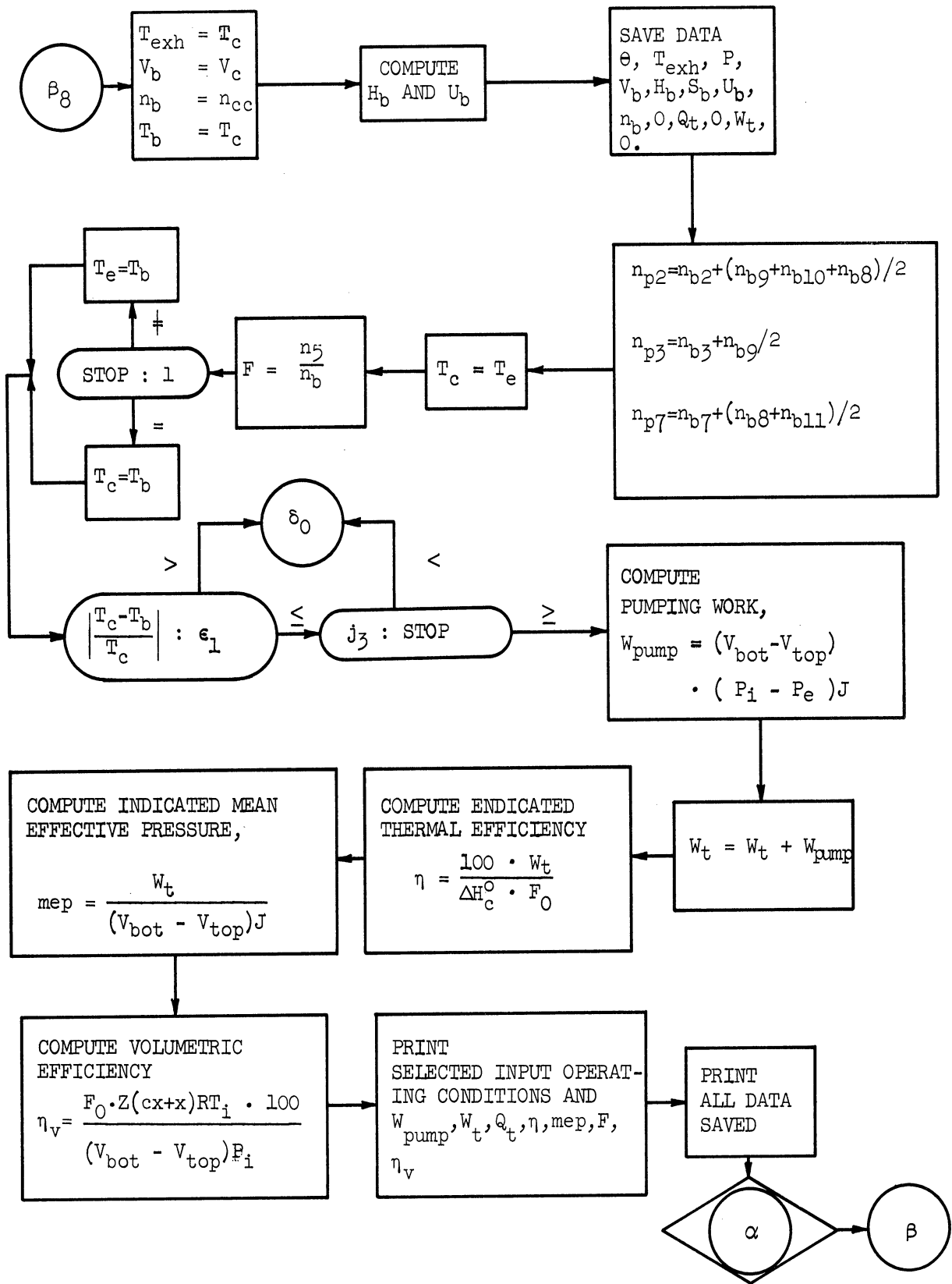












B. Explanation of Symbols Used in Flow Diagram and Computer Program

Symbol used in Flow Diagram	Meaning	Symbol used in Computer Program
A_L	matrix of constants for empirical heat capacity equations in temperature range 298.16 - 1500°K and fundamental data at 298.16°K (see Tables I and II)	AL
A_H	matrix of constants for empirical heat capacity equations in temperature range 1500 - 3500°K and fundamental data at 1500°K (see Tables I and II)	AH
A_h	surface area of combustion chamber head - cm^2	ATOT (1 or 2)
A_{pb}	piston head surface exposed to burned gas - cm^2	ANS (3)
A_{pu}	piston head surface exposed to unburned gas - cm^2	ANS (4)
A_{wb}	wall surface exposed to burned gas - cm^2	ANS (1)
A_{wu}	wall surface exposed to unburned gas - cm^2	ANS (2)
A_{pt}	total piston head surface area - cm^2	ATOT (3 or 4)
A_{wt}	total exposed wall area - cm^2	ARAA
$A_{F,j,i}, A_{b,j,i}, A_{u,j,i}$	three matrices in which are stored values of temperature, volume, and number of moles at selected points in each of the 'i' combustion increments	AF(j,i) AB(j,i) AU(j,i)
a	number of carbon atoms per molecule of fuel	A
Bore	engine bore - in	BORE
b	number of hydrogen atoms per molecule of fuel	B
b_0	density of oxygen molecules in flame element prior to combustion	BO

C_p	constant pressure heat capacity - cal/mole-°K	CP
C_v	constant volume heat capacity - cal mole-°K	CV
c	ratio of nitrogen to oxygen atoms in atmospheric air = 3.764	C
cc	trial value of a thermodynamic property	CC
cprime	trial value of the derivative of a thermo- dynamic property	CPRIM
Dif	an incremental change in temperature - °K	DIF
E	internal energy - cal	E
E_a	activation energy - cal/mole of fuel	EA
F	mass fraction of gas remaining in clearance volume - residual mass fraction	F
F_0	moles of fuel inducted	FO
F_t	total fraction of mass burned	FTOT
F_1-F_{Inc}	the Inc fractions into which the combustion process is divided	FR(1)...FR(INC)
H	enthalpy - cal	H
$\frac{dH}{dT}$	$\frac{dH}{dT} = C_p \cdot n$	HPRIM
h	convective heat transfer coefficient in portion - cal/cm ² -°K	COEF
Inc	number of increments into which the combus- tion process is divided	INC
j_3	number of times complete cycle has been executed	J3
KEY	type of process undergone during a given step by the burned portion or flame element Whenever KEY = 1, process is isenthalpic KEY = 2, process is isentropic and KEY = 3, process occurs at constant internal energy	KEY

kk_1-kk_{nk}	the n_k flame speed constants	KK(1)...KK(NK)
L_t	maximum extent of flame travel if combustion occurs at top center - cm	ATOT(5)
m_l	mass of charge in cylinder - gm	ML
m_f	mass of fresh charge inflamed - gm	MF
mep	indicated mean effective pressure - atm	IMEP
N_c, N_h, N_o, N_n or Carbon, Hydrogen, Oxygen, Nitrogen	mole atoms of carbon, hydrogen, oxygen, and nitrogen (see Equations (2-5) to (2-8))	NC,NH,NO,NN or CARBON,HYDRGN OXYGEN,NITRGN
N_{comdg}	number of compression increments	NCOMDG
N_{expdg}	number of expansion increments	NEXPDG
n	number of moles	N
n_b	total moles of burned gas	NB
n_u	total moles of unburned gas	NU
n_k	number of flame speed constants (K_1 Equation (4-24)) for which computa- tions are to be made	NK
n_{bi}	moles of i-th component of burned mixture	N(i)
n_{fi}	moles of i-th component of flame element	NUN(i)
n_{mi}	moles of i-th component of unburned mixture prior to combustion	NUN(i)
n_{pi}	moles of i-th component of burned products during exhaust process	N(i)
n_{ri}	moles of i-th component of exhaust residual	N(i)
n_{ui}	moles of i-th component of unburned gas	NU(i)
n_{cc}	temporary designation of number of moles	NCC
n_{ff}	moles of fresh charge inflamed	NFF
n_{f-a}	number of air-fuel ratios for which computations are to be made	NFA
P	pressure - atm	P

P_e	exhaust pressure - atm	PEXH
P_i	intake manifold pressure - atm	PMAN
PH	an initial estimate of the value of $X = \sqrt{P_{O_2}}$ (This is used in the computation of the equilibrium composition)	PH
Q	heat transferred - cal	QB
Q_t	total heat transferred - cal	QTOT
q_k	number of heat transfer constants (by which the heat transfer coefficient, Equation (5-18), and subsequently the computed amount of heat transferred, Equation (5-19) are multiplied) for which computations are to be made	QK
$q_{fact_1} - q_{fact_{qk}}$	the q_k heat transfer constants	QFACT(1)... QFACT(QK)
R	gas constant in work units = 0.08205 l-atm/mole-°K	R
R_l	gas constant in heat units = 1.98719 cal/mole-°K	RL
R_0	flame front radius at end of (i-1)-st combustion increment	LASTL
R'_0	outer radius of reaction zone prior to combustion of i-th combustion increment	NEWL
R_6	flame front radius at end of i-th combustion increment	ANS(5)
Rod	connecting rod length - in	ROD
RE	remark - If RE is DONTINO it is assumed that the flame speed constant K_1 , Equation (4-24), is not known	RE
r	compression ratio	CR
rpm	engine speed-rev/min	RPM
S	entropy - cal/°K	S
S_t	turbulent flame speed - cm/sec	ST

STOP	the maximum allowable number of times the complete cycle can be executed	STOP
Stroke	piston stroke - cm	STROKE
$\left(\frac{\partial S}{\partial T}\right)_V$	$\left(\frac{\partial S}{\partial T}\right)_V$	SPRIM
T	temperature - °K	T
T _e	exhaust temperature - °K	T5
T _{exh}	computed exhaust temperature - °K	TEXH
T _i	intake manifold temperature - °K	TMAN
T _{j₂}	temperature of the gas after the heat transfer step of the j ₂ -th expansion increment - °K	T(J1)
T _p	piston head surface temperature - °K	TPISTN
T _w	wall temperature - °K	TW
Tt	alternate designation of a thermodynamic property	TT
U	internal energy - cal	U
$\frac{dU}{dT}$	$\frac{dU}{dT} = C_v \cdot n$	HPRIM
V	volume - liters	V
V _t	total system volume - liters	VTOT
V _{ff}	volume of fresh charge inflamed - liters	VFF
V _{top}	clearance volume - liters	VTOP
V _{bot}	clearance plus swept volumes - liters	VBOT
W	work - cal	WORK
W _{pump}	pumping work - cal	WPUMP
W _t	total work performed - cal	WTOT
x ₁ - x _{n_{f-a}}	the n _{f-a} air-fuel ratios (fraction chemically correct air)	X(1)...X(NFA)
Z	Z = a + b/4	Z

γ	ratio of heat capacity at constant pressure to that at constant volume	G
ΔH_c^0	enthalpy of combustion - cal/mole of fuel	HCOMB
Δl	extension of flame front due to chemical reaction	DS
Δt_i	time required for the combustion of the i-th increment	DT(i)
$\Delta \theta_c$	crank angle increments during the compression process	COMDG
$\Delta \theta_e$	crank angle increments during the expansion process	EXPDG
ϵ_1	allowable relative error in the thermodynamic properties	ERROR1
ϵ_2	allowable relative error in the composition of the products	ERROR2
ϵ_3	allowable absolute error between the calculated and experimental crank angles where maximum pressure occurs	ERROR3
η	indicated thermal efficiency - percent	EFF
η_v	volumetric efficiency - percent	VOLEFF
θ_{max}	crank angle where maximum pressure occurs ($\theta = 0^\circ$ at bottom center and 180° at top center)	PIMAX
θ_{sa}	spark advance - degrees	PI
ρ_0	density of flame element prior to combustion - gr/liter	RH 0

SUPERSCRIPTS

'	designates a trial value of a property	c
---	--	---

SUBSCRIPTS

b	burned	B
u	unburned	U

f	flame element	F
i	intake manifold	MAN
e	exhaust manifold	EXH

STATEMENT LABELS

α	end of computation for one data set	END
α_1	entry to section which finds $T = T(H \text{ or } S \text{ or } U \text{ and } P)$	ENTRY1
α_2	section which finds the pressure of the system with known volume and internal energy and known entropy in the burned portion	ENTRY2
α_4	entry point for change of volume due to piston motion	ENTRY4
α'_1	entry to section which finds $T = T(H \text{ or } S \text{ or } U \text{ and } V)$	ENTCY1
β	entry point for consideration of a second data set	BEGIN
β_1	end of constant pressure combustion of i-th flame element - Step 1	LOC(1)
β_2	end of constant pressure mixing	LOC(2)
β_3	end of recompression to original volume	LOC(3)
β_4	end of change of volume due to piston motion	LOC(4)
β_5	end of recompression of flame element to final pressure - Step 7	LOC(5)
β_6	end of constant volume heat transfer of j_2 -th expansion increment	LCC(6)
β_7	end of isentropic change of volume due to piston motion	LCC(7)
γ_2	beginning of induction and mixing process	LOOP3

δ_0	entry point for recomputation of cycle using a newly computed exhaust temperature	LOOOP1
δ_2	end of induction and beginning of step-wise compression process	LOOOP5
δ_4	end of compression process and entry to combustion process	RETURN
δ_5	beginning of combustion	ANOTHR
δ_9	beginning of j_2 -th expansion increment	LCC1
λ_{1-3}	finds T using Newton's method	ENTRY(1-3)
λ'_{2-3}	finds T using Newton's method	ENTCY(2-3)
λ_{3-6}	finds T using half interval convergence method	ENTRY(4-6)
λ'_{5-6}	finds T using half interval convergence	ENTCY(5-6)
σ_4	end of heat transfer and constant volume readjustment of the i-th combustion increment - Step 5 and 6	SPT(4)
σ_5	entry to section which computes the flame speed constant	SPT(5)
σ_6	completion of the i-th combustion increment	SPT(6)

```
SCOMPILE MAD,PRINT OBJECT,PUNCH OBJECT,EXECUTE,DUMP 3-ICE001
PROGRAM COMMON AL,AH,P,K,R1,NC,NH,NO,NN,R,J2,RE,KK,X,TW,
1QFACT,EA,RATIO,B0,RHO,SPEED,ERROR2,CPF,TP1STN,ATOT,XX,XNUM,
2BORE,STROKE,ROD,VTOP,VBOT,PMAN,PEXH,TMAN,RPM,N,THETA,ERROR1,
3J3,STOP,T5,H5,S5,U5,N5,QTOT,WTOT,F0,NEXTV,HCOMB,INC,Z,C,
4 TZ,TX,COMDG, PMAX,NAME,CR,PI
INTEGER RE,I,I1,I2,I3,J,J1,J2,J3,J4,L,L1,L2,JC,NK,NFA,INC,QK,
1NAME,XNUM,NCOMDG, I,SAVE,KEY,STOP,TZSAVE,TZZERO,TXZERO
DIMENSION AL(87,ALDIM),AH(87,ALDIM),RE(12),X(10),KK(5),QFACT
1(5),NUN(7),P(11),N(11),K(11),XX(105,XDIM),NAME(2),
2FR( 60),ANS(5),NU(7),ATOT(5),TZ(1980), TX( 780),
3 AB(420,DIM),AU(240,DIM),AF(240,DIM),PHF(120,DIM),
4PHB(120,DIM),DT( 60),TT(2)
EQUIVALENCE (CPRIM,SPRIM),(HPRIM,CV),(HMIX,TT),(TZ,TZZERO),
1(TX,TXZERO)
VECTOR VALUES DIM = 2,0,60
VECTOR VALUES ALDIM = 2,0,8
VECTOR VALUES XDIM = 2,0,21
READ FORMAT SPHT,AL(1,1)...AL(11,8),AH(1,1)...AH(11,8)
VECTOR VALUES SPHT = $4E15.8*$
READ FORMAT REMARK,RE...RE(12),ERROR1,ERROR2,ERROR3,COMDG,T5,
1STOP,NFA,NK,QK,INC,A,B,EA,HCOMB,SPEED,NAME...NAME(2),X(1)...X
2(NFA),KK(1)...KK(NK),QFACT(1)...QFACT(QK)
PRINT FORMAT DUMP1,$IAL = $,AL(1,1)...AL( 7,8)
Z = A+B/4.
PRINT FORMAT DUMP1,$ AH = $,AH(1,1)...AH(11,8)
VECTOR VALUES DUMP1 = $C6/(8E15.8)*$
VECTOR VALUES REMARK = $13C6/5F10.5,5I2/5F10.1,3C6/(7F10.2)*$
READ FORMAT FRACTN,XDIM(2),ATOT(1)...ATOT(5),FR(1)...FR(INC)
VECTOR VALUES FRACTN = $I10,5F10.5/(14F5.5)*$
R = .08205
R1 = 1.98719
C = 3.764
READ FORMAT FRAT,XX(1,1)...XX(5,XDIM(2))
VECTOR VALUES FRAT = $14F5.5*$
XNUM = XDIM(2)
DIM(2) = INC
PRINT FORMAT REMARK,RE...RE(12),ERROR1,ERROR2,ERROR3,COMDG,T5
1,STOP,NFA,NK,QK,INC,A,B,EA,HCOMB,SPEED,NAME...NAME(2),X(1)...
2X(NFA),KK(1)...KK(NK),QFACT(1)...QFACT(QK)
PRINT FORMATFRACFB,XDIM(2),ATOT(1)...ATOT(5),FR(1)...FR(INC),
1XX(1,1)...XX(5,XDIM(2))
VECTOR VALUES FRACFB = $ I10,5F10.5/(14F8.5)*$
PH51 = 1.
BEGIN READ FORMAT ENGINE,BORE,STROKE,ROD,RPM,VTOP,PMAN,PEXH,TMAN,TP
1ISTN,TW,PI,PIMAX,D1,D2
VECTOR VALUES ENGINE = $7F10.5*$
BORE = BORE *2.54
STROKE=STROKE*2.54
ROD = ROD*2.54
VTOP = VTOP/61.03
VBOT = VTOP+3.1416*BORE/4000.*BORE*STROKE
CR = VBOT/VTOP
NCOMDG = (179.-PI)/COMDG
J3 = 0
THROUGH END, FOR I1 = 1,1,I1.G.NFA
X = X(I1)
THROUGH END, FOR I2 = 1,1,I2.G.NK
KK = KK(I2)
```

```
THROUGH END, FOR I3 = 1,1,I3.G.QK
QFACT = QFACT(I3)
PRINT FORMAT INTRO, NAME...NAME(2),X,BORE/2.54,EA,TPISTN,STRO
1KE/2.54,TW,ROD/2.54,HCOMB,TMAN,VTOP*61.03,PMAN,61.03*(VBOT-VT
2OP),PEXH,CR,PI
N5 = PEXH/R*VTOP/T5
NC = A
NH = B
NO = X*(2.*NC+.5*NH)
NN = C*NO
NCC = A+B/2.+Z*(C*X+X-1.)
NEXTV = NCC*R/PEXH*T5
EXECUTE EQUIL.(T5)
PRINTFORMATINTOR1,RPM,INC,ERROR1*100.,KK,QFACT,RE(4)...RE(12)
BACK VC = NEXTV
EXECUTE AMOUNT.(T5,VC,NCC,PH51,N)
NEXTV = NCC*R/PEXH*T5
WHENEVER .ABS.((NEXTV-VC)/VC).G.ERROR1,TRANSFER TO BACK
F5 = VTOP/VC
N(2) = F5*(N(2)+(N(9)+N(10)+N(8))/2.)
N(3) = F5*(N(3)+N(9)/2.)
N(4) = F5*N(4)
N(5) = F5*N(5)
N(6) = F5*N(6)
N(7) = F5*(N(7)+(N(8)+N(11))/2.)
TRANSFER TO LOOC1
LOOP1 N5 = PEXH/R*VTOP/T5
F5 = N5/(N(2)+N(3)+N(4)+N(5)+N(6)+N(7))
THROUGH LOOP2,FORL1 = 2,1,L1.G.7
LOOP2 N(L1) = F5*N(L1)
LOOC1 EXECUTE ENTHAL.(2,7,H5,T5,N)
J3 = J3 + 1
U5 = H5 - N5*R1*T5
PRINT FORMAT HEAD3
VECTOR VALUES HEAD3 = $4H0LOC,S2,1HJS2,3HINCS4,5HTHETAS4, 6HT
1 - *K,S4,8HP - ATM.S7,6HV - L.S4,8HH - CAL.S6,11HS - CAL./*KS
26,8HU - CAL.S5,11HTOTAL MOLES*$
EXECUTE ENTROP.(2,7.S5,T5,N5,PEXH,N)
PRINTFORMAT OUT,5,0,1,540.,T5,PEXH,VTOP,H5,S5,U5,N5,N(2)...N(
17)
NUN(4) = N(4)
NUN(5) = N(5)
NUN(6) = N(6)
NUN(7) = N(7)
J = 0
WHENEVER J3.E.1,T1 = TMAN
NEXTT = T1
T = NEXTT
LOOP3 J = J + 1
NO = PMAN/R*VBOT/T - N5
FO = NO/(1. + Z*X*(C+1.))
NUN(1) = FO
NUN(2) = FO*Z*X
NUN(3) = NUN(2)*C
EXECUTE ENTHAL.(1,3,H0,TMAN,NUN)
EXECUTE HPRIME.(1,3,HPRIM,T,NUN)
HPRIM = HPRIM - H0/T*(1. + N5/NO)
H = H0 + U5 + PMAN*VTOP*24.2179
```

```

-----
NUN(2) = NUN(2) + N(2)
NUN(3) = NUN(3) + N(3)
-----
EXECUTE ENTHAL.(1,7,HMIX,T,NUN)
NEXTT = T - (HMIX-H)/HPRIM
-----
WHENEVER .ABS.((HMIX-H)/H).G.ERROR1,TRANSFER TO LOOP3
N1 = NO + N5
-----
EXECUTE ENTROP.(1,7,S1,T,N1,PMAN,NUN)
U = H - N1*R1*T
-----
U1 = U
M1 = NUN(1)*(12.01*A+1.008*B)+NUN(2)*32.+NUN(3)*28.02+NUN(4)*
144.01+NUN(5)*18.016+NUN(6)*28.01+NUN(7)*2.016
T1 = T
PRINT FORMATOUT,0,J,1,0.,TMAN,PMAN,VBOT,H0,0.,0.,NO,F0*X*Z,F0
1*C*X*Z
-----
THETA = 0.
JC = 0
TZZERO = 1
TZ(TZZERO ) = 0.
TZ(TZZERO + 1 ) = T
TZ(TZZERO + 2 ) = PMAN
-----
TZ(TZZERO + 3 ) = VBOT
TZ(TZZERO + 4 ) = H
TZ(TZZERO + 5 ) = S1
TZ(TZZERO + 6 ) = U
TZ(TZZERO + 7 ) = N1
TZZERO = TZZERO + 13
-----
PRINT FORMAT OUT,1,J,1,0.,T,PMAN,VBOT,H,S1,U,N1,NUN(1)...
1NUN(7),M1
-----
LOOP5
THETA = THETA + COMDG
-----
LOOP2
JC = JC + 1
LASTU = U
NEXTV = VOLUME.(THETA)
EXECUTE HPRIME.(1,7,CV,T,NUN)
CV = CV/N1 - R1
NEXTT = T*(V/NEXTV).P.(R1/CV)
V = NEXTV
J = 0
-----
LOOP4
J = J + 1
T = NEXTT
P = T*R/V*N1
EXECUTE ENTROP.(1,7,S,T,N1,P,NUN)
EXECUTE SPRIME.(1,7,SPRIM,T,NUN)
SPRIM = SPRIM - N1/T*R1
NEXTT = T - (S-S1)/SPRIM
WHENEVER .ABS.((NEXTT-T)/T).G.ERROR1,TRANSFER TO LOOP4
EXECUTE ENTHAL.(1,7,H,T,NUN)
U = H - N1*R1*T
-----
TZ(TZZERO ) = THETA
TZ(TZZERO + 1 ) = T
TZ(TZZERO + 2 ) = P
TZ(TZZERO + 3 ) = V
TZ(TZZERO + 4 ) = H
TZ(TZZERO + 5 ) = S
TZ(TZZERO + 6 ) = U
TZ(TZZERO + 7 ) = N1
-----
TZ(TZZERO + 8 ) = 0.
TZ(TZZERO + 9 ) = 0.
-----

```

```
-----
TZ(TZZERO + 10) = LASTU - U
TZ(TZZERO + 11) = U1 - U
-----
TZ(TZZERO + 12) = 0.
TZZERO = TZZERO + 13
-----
PRINT FORMAT OUT,2,J,JC,THETA,T,P,V,H,S,U
WHENEVER JC.L.NCOMDG, TRANSFER TO LOOOP5
WHENEVER JC.E.NCOMDG
THETA = 180.-PI
-----
TRANSFER TO LOOC2
END OF CONDITIONAL
-----
V2 = V
T2 = T
-----
H2 = H
U2 = U
-----
P2 = P
THROUGH END1, FOR L1=1,1,L1.G.7
END1  NU(L1) = NUN(L1)
      WHENEVER J3.E.1
      J1 = 0
      OTHERWISE
      J1 = J3
      END OF CONDITIONAL
      CARBON = F0*A +NU(4) +NU(6)
      HYDRGN = F0*B + 2.*(NU(5) +NU(7))
      OXYGEN = X*(2.*CARBON+.5*HYDRGN)
      NITRGN = C*OXYGEN
-----
RETURN TZSAVE = TZZERO
      THETA = 180.-PI
      TZZERO = TZSAVE
      TXZERO = 1
      QB = .
      LASTL = 0.
      FTOT = 0.
      QTOT = 0.
      WTOT = U1-U2
      SEC = 0.
-----
HERE  THROUGH HERE, FOR L1 = 1,1,L1.G.7
      NUN(L1) = NU(L1)
      TU = T2
      P = P2
      VU = V2
      HU = H2
      SU = S1
      EC = U2
-----
      NU = N1
      HB = 0.
      VB = 0.
      WHENEVER RE(0).E.$DONTNOS
      J4 = 5
      OTHERWISE
      J4 = 6
      END OF CONDITIONAL
      I = 0
ANOTHR LASTP = P
      I = I + 1
      E = EC
-----
      DT = DT(I)
      FR = FR(I)
-----
```

```
-----
FTOT = FTOT + FR
VTOT = VOLUME.(THETA)
-----
NFF = N1*FR
MF = M1*FR
-----
VFF = NFF*R/P*TU
EXECUTE AREA.(VB,VTOT,ANS,D1,D2)
-----
LASTL = ANS(5)
EXECUTE AREA.(VB+VFF,VTOT,ANS,D1,D2)
-----
NEWL = ANS(5)
DS = NEWL - LASTL
-----
B0 = NUN(2)/VU
RHO = MF/VFF
-----
T0 = TU
Z1 = NFF/NU
-----
HF = HU*Z1
VU = VU-VFF
-----
NU = NU-NFF
HU = HU-HF
-----
SU = SU*(1.-Z1)
NFC = CARBON*FR
-----
NFH = HYDRGN*FR
NFO = OXYGEN*FR
-----
NFN = NITRGN*FR
KEY = 1
-----
L = 1
THROUGH LOC, FOR L1 = 1,1,L1.G.7
-----
LOC  NUN(L1) = NUN(L1)*(1.-Z1)
-----
WHENEVER J1.G.1, TRANSFER TO SKIP
-----
DT(I) = 0.
WHENEVER I.G.1
-----
II = I-1
MQ = FR(I)/FR(II)
-----
AF(1,I) = AF(1,II)
AF(2,I) = AF(2,II)*MQ
-----
PHF(I,I) = PHF(I,II)*MQ
OTHERWISE
-----
PHF(1,I) = 1.
AF(1,I) = 4.*TU
-----
AF(2,I) = NFF
END OF CONDITIONAL
-----
SKIP TC = AF(1,I)
-----
NCC = AF(2,I)
PH = PHF(1,I)
-----
TT = HF
NC = NFC
-----
NO = NFO
NH = NFH
-----
NN = NFN
ENTRY1 J = 0
-----
L2 = KEY
NEXTT = TC
-----
LOC1 TC = NEXTT
-----
J = J+1
NEXTV = NCC*R/P*TC
EXECUTE EQUIL.(TC)
-----
LOC2 VC = NEXTV
-----
EXECUTE AMOUNT.(TC,VC,NCC,PH,N)
NEXTV = NCC*R/P*TC
-----
```

	WHENEVER $\cdot \text{ABS} \cdot ((\text{NEXTV} - \text{VC}) / \text{VC}) \cdot \text{G} \cdot \text{ERROR1}$, TRANSFER TO LOC2 TRANSFER TO ENTRY(L2)
ENTRY(1)	EXECUTE ENTHAL $\cdot (2, 11, \text{CC}, \text{TC}, \text{N})$ EXECUTE HPRIME $\cdot (2, 11, \text{CPRIM}, \text{TC}, \text{N})$ TRANSFER TO LESS
ENTRY(2)	EXECUTE ENTROP $\cdot (2, 11, \text{CC}, \text{TC}, \text{NCC}, \text{P}, \text{N})$ EXECUTE SPRIME $\cdot (2, 11, \text{CPRIM}, \text{TC}, \text{N})$ TRANSFER TO LESS
ENTRY(3)	EXECUTE ENTHAL $\cdot (2, 11, \text{CC}, \text{TC}, \text{N})$ $\text{CC} = \text{CC} - \text{NCC} * \text{R1} * \text{TC}$ EXECUTE HPRIME $\cdot (2, 11, \text{CPRIM}, \text{TC}, \text{N})$ $\text{CPRIM} = \text{CPRIM} - \text{NCC} * \text{R1}$
LESS	$\text{Q} = \text{CC} - \text{TT}$ $\text{NEXTT} = \text{TC} - \text{Q} / \text{CPRIM}$ WHENEVER $\text{J} \cdot \text{E} \cdot 5$ $\text{L2} = \text{L2} + 3$ $\text{DIF} = \cdot \text{ABS} \cdot (\text{TC} - \text{NEXTT})$ END OF CONDITIONAL WHENEVER $\cdot \text{ABS} \cdot (\text{Q} / \text{TT}) \cdot \text{G} \cdot \text{ERROR1}$, TRANSFER TO LOC1 TRANSFER TO LOC(L)
ENTRY(4)	EXECUTE ENTHAL $\cdot (2, 11, \text{CC}, \text{TC}, \text{N})$ TRANSFER TO MORE
ENTRY(5)	EXECUTE ENTROP $\cdot (2, 11, \text{CC}, \text{TC}, \text{NCC}, \text{P}, \text{N})$ TRANSFER TO MORE
ENTRY(6)	EXECUTE ENTHAL $\cdot (2, 11, \text{CC}, \text{TC}, \text{N})$ $\text{CC} = \text{CC} - \text{NCC} * \text{R1} * \text{TC}$
MORE	$\text{QQ} = \text{CC} - \text{TT}$ WHENEVER $\text{Q} / \text{QQ} \cdot \text{L} \cdot 0 \cdot \cdot \text{DIF} = \text{DIF} / 2 \cdot$ $\text{NEXTT} = \text{TC} - \text{DIF} * \text{QQ} / \cdot \text{ABS} \cdot (\text{QQ})$ $\text{Q} = \text{QQ}$ WHENEVER $\cdot \text{ABS} \cdot (\text{Q} / \text{TT}) \cdot \text{G} \cdot \text{ERROR1}$, TRANSFER TO LOC1 TRANSFER TO LOC(L)
LOC(1)	$\text{AF}(1, \text{I}) = \text{TC}$ $\text{AF}(2, \text{I}) = \text{NCC}$ $\text{PHF}(1, \text{I}) = \text{PH}$ EXECUTE ENTROP $\cdot (2, 11, \text{SF}, \text{TC}, \text{NCC}, \text{P}, \text{N})$ WHENEVER $\text{I} \cdot \text{E} \cdot 1$ TRANSFER TO LOC(2) END OF CONDITIONAL $\text{TT} = \text{HB} + \text{TT}$ $\text{NC} = \text{NBC} + \text{NC}$ $\text{NO} = \text{NBO} + \text{NO}$ $\text{NH} = \text{NBH} + \text{NH}$ $\text{NN} = \text{NBN} + \text{NN}$ WHENEVER $\text{J1} \cdot \text{L} \cdot 2$ $\text{AB}(1, \text{I}) = \text{AB}(1, \text{II})$ $\text{AB}(2, \text{I}) = \text{AB}(2, \text{II}) + \text{NCC}$ $\text{PHB}(1, \text{I}) = 1 \cdot$ END OF CONDITIONAL $\text{TC} = \text{AB}(1, \text{I})$ $\text{NCC} = \text{AB}(2, \text{I})$ $\text{PH} = \text{PHB}(1, \text{I})$ $\text{L} = 2$ TRANSFER TO ENTRY1
LOC(2)	$\text{AB}(1, \text{I}) = \text{TC}$ $\text{AB}(2, \text{I}) = \text{NCC}$ $\text{PHB}(1, \text{I}) = \text{PH}$ $\text{VB} = \text{VC}$

NB = NCC
TB = TC
HB = TT
NBC = NC
NBH = NH
NBO = NO
NBN = NN
L = 3
L1 = 3
UB = HB - NB*R1*TB
UU = HU - NU*R1*TU
UC = UB + UU
WHENEVER J1.G.1
TB = AB(3,I)
NB = AB(4,I)
VB = AB(5,I)
TU = AU(1,I)
VU = AU(2,I)
PH = PHB(2,I)
UC = E
P = R/VTOT*(NB*TB+NU*TU)
END OF CONDITIONAL

ENTRY2
LOC3

KEY = 3
JC = 0
JC = JC + 1
EXECUTE HPRIME.(2,11,CP,TB,N)
CP = CP/NB-R1
NEXTP = R/VTOT*((EC-UC)/CP + NB*TB + NU*TU)
EXECUTE HPRIME.(1,7,CP,TU,NUN)
TU = TU*(NEXTP/P).P.(R1/CP*NU)
P = NEXTP
NEXTT = TU

LOC4

TU = NEXTT
EXECUTE ENTROP.(1,7,TT,TU,NU,P,NUN)
EXECUTE SPRIME.(1,7,CPRIM,TU,NUN)
NEXTT = TU-((TT-SU)/CPRIM)
WHENEVER .ABS.((NEXTT-TU)/TU).G.ERROR1,TRANSFER TO LOC4
VU = NU*R/P*TU
EXECUTE ENTHAL.(1,7,HU,TU,NUN)
UU = HU - NU*R1*TU
UB = EC - UU
TC = TB
TT = UB
NCC = NB

LOC(3)

TRANSFER TO ENTRY1
TB = TC
VB = VC
NB = NCC
UC = UB + UU
VC = VU + VB
WHENEVER .ABS.((VC-VTOT)/VTOT).G.ERROR1,TRANSFER TO LOC3
EXECUTE ENTHAL.(2,11,HB,TB,N)
PRINT FORMAT OUT,L1,J,I,THETA,TU,P,VU,HU,SU,UU,NU,NUN(1)...NU
1N(7)
EXECUTE ENTROP.(2,11,SB,TB,NB,P,N)
PRINT FORMAT OUT,L1,JC,I,THETA,TB,P,VB,HB,SB,UB,NB,N(2)...N(1
11)
TRANSFER TO SPT(L1)


```
SPT(3)  AB(3,I) = TB
        AB(4,I) = NB
        AB(5,I) = VB
        AU(1,I) = TU
        AU(2,I) = VU
        PHB(2,I) = PH
        TT(1) = SU
        TT(2) = SB
        PRINT FORMAT HEAD3
        SAVE = J1
        EXECUTE AREA.(VB,VTOT,ANS,D1,D2)
ENTRY4  KEY = 2
        PHETA = 6.*RPM*DT+THETA
        VTOT = VOLUME.(PHETA)
        L = 4
        TB = AB(3,I)
        NB = AB(4,I)
        VBB = AB(5,I)
        TU = AU(1,I)
        VUU = AU(2,I)
        PH = PHB(2,I)
        SU = TT(1)
        SB = TT(2)
        VC = VBB + VUU
        EXECUTE EQUIL.(TB)
        EXECUTE AMOUNT.(TB,VBB,NB,PH,N)
        JC = 0
           3  + +
        GB = CP/(CP-R1*NB)
        WHENEVER I.E.INC
        VB = VTOT
        TB = TB*(VBB/VB).P.(GB-1.)
        P = P*(VBB/VB).P.GB
        J4 = 6
        TRANSFER TO SPT1
        END OF CONDITIONAL
        EXECUTE HPRIME.(1,7,CP,TU,NUN)
        GU = CP/(CP-R1*NU)
        VB = VTOT/(1.+VUU/VBB*(VC/VTOT).P.((GU-GB)/GB))
        VU = VTOT - VB
        TU = TU*(VUU/VU).P.(GU-1.)
        TB = TB*(VBB/VB).P.(GB-1.)
        P = R/VTOT*(NB*TB+NU*TU)
        NEXTT = TU
LOC6    TU = NEXTT
        EXECUTE ENTROP.(1,7,SC,TU,NU,P,NUN)
        EXECUTE SPRIME.(1,7,SPRIM,TU,NUN)
        NEXTT = TU-((SC-SU)/SPRIM)
        WHENEVER .ABS.((NEXTT-TU)/TU).G.ERROR1,TRANSFER TO LOC6
        VU = NU*R/P*TU
SPT1    TT = SB
        TC = TB
        NCC = NB
        TRANSFER TO ENTRY1
LOC(4)  TB = TC
        VBB = VC
        NB = NCC
        VUU = VU
```

```
VC = VBB + VUU
WHENEVER .ABS.((VC-VTOT)/VTOT).G.ERROR1,TRANSFER TO LOC5
VB = VBB
EXECUTE ENTHAL.(1,7,HU,TU,NUN)
EXECUTE ENTHAL.(2,11,HB,TB,N)
UU = HU-NU*R1*TU
PRINT FORMAT OUT,L,JC,I,PHETA,TU,P,VU,HU,SU,UU,NU,NUN(1)...NU
1N(7)
UB = HB-NB*R1*TB
COEF = 0.
EC = UU + UB
PRINT FORMAT OUT,L,J,I,PHETA,TB,P,VB,HB,SB,UB,NB,N(2)...N(11)
WHENEVER J1.E.0,TRANSFER TO SPT(4)
WHENEVER FTOT .LE. .02, TRANSFER TO SPT(4)
L = 3
L1 = 4
QB = QTRAN2.(TB,TU,P,DT,ANS,COEF)
EC = EC-QB
UC = EC
WHENEVER J1.NE.1
TB = AB(6,I)
VB = AB(7,I)
TU = AU(3,I)
VU = AU(4,I)
P = R/VTCT*(NB*TB+NU*TU)
END OF CONDITIONAL
KEY = 3
```

SPT(4)

```
TRANSFER TO ENTRY2
AB(6,I) = TB
AB(7,I) = VB
AU(3,I) = TU
AU(4,I) = VU
WHENEVER I.E.1,TRANSFER TO LOC(5)
L = 5
WHENEVER J1.G.1
PH = PHF(2,I)
TC = AF(3,I)
NCC= AF(4,I)
OTHERWISE
PH = 1.
NCC= NFF
END OF CONDITIONAL
TT = SF
NC = NFC
NH = NFH
NO = NFO
NN = NFN
KEY = 2
TRANSFER TO ENTRY1
```

LOC(5)

```
TFLAME = TC
PHF(2,I) = PH
AF(3,I) = TC
AF(4,I) = NCC
RATIO = NFF/NCC
EXECUTE HPRIME.(2,11,CPF,TC,N)
CPF = CPF/MF
R2 = R1*NCC/MF
EXECUTE STURB.(T0,TFLAME,ST,R2)
DT = DS/ST
```

```
NC = NBC
NH = NBH
NO = NBO
NN = NBN
J1 = SAVE
WHENEVER .ABS.((PHETA-THETA-6.*RPM*DT)/PHETA).G.ERROR1
SAVE = J1
J1 = 2
TRANSFER TO ENTRY4
END OF CONDITIONAL
PRINT FORMAT VOLU,ST,TFLAME,COEF,FTOT
VECTOR VALUES VOLU =$14HQVOLUMES AGREE,S5,13HFLAME SPEED =
1F10.5,S5,8HTFLAME = F10.5,S5,20HHEAT TRANSFER COEF = E15.8/
219H FRACTION BURNED = F10.8*$
WORK = E-EC-QB
WTOT = WTOT+WORK
QTOT = QTOT+QB
SEC = SEC + DT
LSTTH = THETA
THETA = PHETA
DT(I) = DT
PRINT FORMAT INTRI,THETA,TU,P,VTOT,HU+HB,SU+SB,UU+UB,NU+NB,QB
1,QTOT,WORK,WTOT, SEC,TB,P,VB,HB,SB,UB,NB,ANS...ANS(5),NEWL,
2LASTL
VECTOR VALUES INTRI=$S6,4HTIMES5,1HTS9,1HPS9,1HVS9,1HHS9,1HS
1S9,1HUS9,1HNS9,2HQBS8,4HQTOTS6,4HWORKS6,4HWTOT/(12F10.5)*$
TX(TXZERO ) = THETA
TX(TXZERO + 1 ) = TB
TX(TXZERO + 2 ) = P
TX(TXZERO + 3 ) = VB
TX(TXZERO + 4 ) = HB
TX(TXZERO + 5 ) = SB
TX(TXZERO + 6 ) = UB
TX(TXZERO + 7 ) = NB
TX(TXZERO + 8 ) = ST
TX(TXZERO + 9 ) = TFLAME
TX(TXZERO + 10) = ANS(5)
TX(TXZERO + 11) = FTOT
TX(TXZERO + 12) = VB/VTOT
TXZERO = TXZERO + 13
TZ(TZZERO ) = THETA
TZ(TZZERO + 1 ) = TU
TZ(TZZERO + 2 ) = P
TZ(TZZERO + 3 ) = VTOT
TZ(TZZERO + 4 ) = HU+HB
TZ(TZZERO + 5 ) = SU+SB
TZ(TZZERO + 6 ) = UU+UB
TZ(TZZERO + 7 ) = NU+NB
TZ(TZZERO + 8 ) = QB
TZ(TZZERO + 9 ) = QTOT
TZ(TZZERO + 10) = WORK
TZ(TZZERO + 11) = WTOT
TZ(TZZERO + 12) = COEF
TZZERO = TZZERO + 13
PRINT FORMAT HEAD3
TRANSFER TO SPT(J4)
SPT(5) WHENEVER (P-LASTP).GE.0.,TRANSFER TO ANOTHR
NEXTK = KK*(LSTTH-(180.-PI))/(PIMAX-(180.-PI))
```

```
WHENEVER .ABS.(PIMAX-LSTTH).G.ERROR3
J1 = J1 + 1
KK = NEXTK
PRINT FORMAT OUTPUT, KK
VECTOR VALUES OUTPUT = $5HOKK = E15.8#3
TRANSFER TO RETURN
END OF CONDITIONAL
J4 = 6
PMAX = LASTP
WHENEVER J3.E.1
J1 = 0
OTHERWISE
J1 = J3
END OF CONDITIONAL
SPT(6) WHENEVER P.G.LASTP, PMAX = P
WHENEVER I.L.INC, TRANSFER TO ANOTHR
EXECUTE FINISH.(EC, TB, VB, PH, BACK, LOOOP1, F, D1)
EXECUTE OUTPUT.(F)
END CONTINUE
TRANSFER TO BEGIN
VECTOR VALUES INTRO = $1H1, $35, 49HA MODEL SIMULATING THE INTE
1RNAL-COMBUSTION ENGINE/7HOENGINES39, 4HFUELS36, 20HOPERATING CO
2NDITIONS/28HOFOUR STROKE AUTOMOTIVE TYPESI273C6, S22, I4HAIR-FU
3EL RATIO F9.2/5H BORES8, F10.2, 4H IN.S13, 17HACTIVATION ENERGY
4F10.2, 24H CAL/MOLE PISTON TEMP F12.2, 2H K/7H STROKES6, F10.
52, 4H IN.S13, 11HENTHALPY OFS29, 9HWALL TEMPF14.2, 2H K/4H RODS9,
6F10.2, 4H IN.S13, 10HCOMBUSTIONF17.2, 22H CAL/MOLE MAN. TEMP
7F14.2, 2H K/12H CLEAR. VOL.F11.2, 7H CU.IN.S50, 8HMAN. PR.F15.2,
84H ATM/11H SWEPT VOL.F12.2, 7H CU.IN.S90, 8HEXH. PR.F15.2, 4H AT
9M/18H COMPRESSION RATIOF5.2, 7H TO ONES50, 10HSPARK ADV.F13.2#5
VECTOR VALUES INTORI = $580, 5HSPEEDS10, F8.2, 4H RPM/ 21HCOMBU
1STION OCCURS IN 13, 34H INCREMENTS. ERRORS ARE LESS THAN F10.
25, 33H PERCENT. FLAME SPEED FACTOR IS E15.8/24H HEAT TRANSFER
3 FACTOR IS F10.5, S3, 10C6#5
VECTOR VALUES OUT = $1H0, 3I3, S2, 3F10, 3, F15, 8, 4E15.8/(6E20.8)*5
INTERNAL FUNCTION VOLUME.(ZETA) = .0007854*BORE*BORE*(STROKE/
12.*(1.+COS.(ZETA*.01745))+ROD*(1.-SQRT.(1.-TSTROKE/2.*SIN#T
2ZETA*.01745)/(ROD).P.2)))+ VTOP
END OF PROGRAM
```

SCOMPILE MAD, PRINT OBJECT,PUNCH OBJECT 4FNSH001

EXTERNAL FUNCTION(ENERGY,TEMPT,VOL,PHP,BACK,LOOOP1,F,D1)
PROGRAM COMMON AL,AH,P,K,R1,NC,NH,NO,NN,R,J2,RE,KK,X,TW,
1QFACT,EA,RATIO,B0,RHO,SPEED,ERROR2,CPF,TPISTN,ATOT,XX,XNUM,
2BORE,STROKE,ROD,VTOP,VBOT,PMAN,PEXH,TMAN,RPM,N,THETA,ERROR1,
3J3,STOP,T5,H5,S5,U5,N5,QTOT,WTOT,F0,NEXTV,HCOMB,INC,Z,C,
4 TZ,TX,COMDG, PMAX,NAME,CR,PI

STATEMENT LABEL BACK,LOOOP1
INTEGER RE,I,II,I2,I3,J,J1,J2,J3,G,L2,N1,N2,INC,NAME,XNUM,
2NEXPDG, KEY,STOP,TZZERO
DIMENSION AL(87,ALDIM),AH(87,ALDIM),RE(12),X(10),KK(5),QFACT
1(5),P(11),N(11),K(11),XX(105,XDIM),NAME(2),RT(40),ATOT(5),B(7
2),TZ(1980),TX(780)

EQUIVALENCE (TZ,TZZERO)
VECTOR VALUES XDIM = 2,0,21
VECTOR VALUES ALDIM = 2,0,8
ENTRY TO FINISH.

UB = ENERGY
TB = TEMPT
VB = VOL
PH = PHP
DELTAT = 0.
NEXPDG = (359.-THETA)/COMDG*2.
DTHETA = 360.-THETA-NEXPDG*COMDG/2.
DT = DTHETA/6./RPM

LCC1

J1 = 0
J1 = J1 + 1
TC =RT(J1)
E = UB
WHENEVER J3.E.1,TC=TB-DELTAT
ARAA = ARREA.(THETA)
QB = QTRANS.(P,ARAA,COEF)
QTOT = QTOT + QB
UB = E-QB
G = 6
KEY = 3
TT = UB

ENTCY1

J = 0
L2 = KEY
NEXTT = TC
LCC(2) TC = NEXTT
J = J + 1
EXECUTE EQUIL.(TC)
EXECUTE AMOUNT.(TC,VB,NB,PH,N)
TRANSFER TO ENTCY(L2)

ENTCY(2)

P = NB*R/VB*TC
EXECUTE ENTROP.(2,11,CC,TC,NB ,P,N)
EXECUTE SPRIME.(2,11,CPRIM,TC,N)
CPRIM = CPRIM - NB*R1/TC
TRANSFER TO LESSC

ENTCY(3)

EXECUTE ENTHAL.(2,11,CC,TC,N)
CC = CC- NB*R1*TC
EXECUTE HPRIME.(2,11,CPRIM,TC,N)
CPRIM = CPRIM- NB*R1

LESSC

Q = CC-TT
NEXTT = TC-Q/CPRIM
WHENEVER J.E.5
L2 = L2+3
DIF = .ABS.(TC - NEXTT)

```
END OF CONDITIONAL
WHENEVER .ABS.(Q/TT).G.ERROR1,TRANSFER TO LCC(2)
TRANSFER TO COMMOC
ENTCY(5) P = NB*R/VB*TC
EXECUTE ENTROP.(2,11,CC,TC,NB ,P,N)
TRANSFER TO MOREC
ENTCY(6) EXECUTE ENTHAL.(2,11,CC,TC,N)
CC = CC- NB*R1*TC
MOREC QQ = CC-TT
WHENEVER Q/QQ.L.0.,DIF = DIF/2.
NEXTT = TC - DIF*QQ/.ABS.(QQ)
Q = QQ
WHENEVER .ABS.(Q/TT).G.ERROR1,TRANSFER TO LCC(2)
COMMOC PRINTFORMATOUT,G,J,I,THETA,TC,P,VB,TT,0.,CC,NB
TRANSFER TO LCC(G)
LCC(6) RT(J1)= TC
DELTAT = TB-TC
TB = TC
P = NB*R/VB*TB
EXECUTE ENTROP.(2,11,SB,TB,NB,P,N)
THETA = THETA+DTHETA
V = VOLUME.(THETA)
TC = TB*(VB/V).P.(NB*R1/CPRIM)
VB = V
G = 7
KEY = 2
TT = SB
TRANSFER TO ENTCY1
LCC(7) EXECUTE ENTHAL.(2,11,HB,TC,N)
TB = TC
UB = HB-NB*R1*TB
WORK = E-UB-QB
WTOT = WTOT+WORK
PRINT FORMAT OUT,G,J,J1,THETA,TB,P,VB,HB,SB,UB,NB,N(2)...N(11
1),0.,QB,QTOT,WORK,WTOT,COEF,ARAA
TZ(TZZERO ) = THETA
TZ(TZZERO + 1 ) = TB
TZ(TZZERO + 2 ) = P
TZ(TZZERO + 3 ) = VB
TZ(TZZERO + 4 ) = HB
TZ(TZZERO + 5 ) = SB
TZ(TZZERO + 6 ) = UB
TZ(TZZERO + 7 ) = NB
TZ(TZZERO + 8 ) = QB
TZ(TZZERO + 9 ) = QTOT
TZ(TZZERO + 10) = WORK
TZ(TZZERO + 11) = WTOT
TZ(TZZERO + 12) = COEF
TZZERO = TZZERO + 13
WHENEVER THETA.G.359.,TRANSFER TO LCC3
WHENEVER J1.G.1,TRANSFER TO LCC1
DELTAT = DELTAT/2.
DTHETA = COMDG/2.
DT = DTHETA/6./RPM
TRANSFER TO LCC1
LCC3 WHENEVER J3.E.1
EXECUTE HPRIME.(2,11,CPRIM,TC,N)
TEXH = TB*(PEXH/P).P.(R1*NB/CPRIM)
```

```
END OF CONDITIONAL
NEXTT = TEXH
P = PEXH
NEXTV = NB*R/PEXH*TEXH
J = 0
LCC(3) TC = NEXTT
J = J + 1
LCC5 EXECUTE EQUIL.(TC)
VB = NEXTV
EXECUTE AMOUNT.(TC,VB,NB,PH,N)
NEXTV = NB*R/PEXH*TC
WHENEVER .ABS.((NEXTV-VB)/VB).G.ERROR1,TRANSFER TO LCC5
EXECUTE ENTROP.(2,11,CC,TC,NB,P,N)
Q = CC - TT
EXECUTE SPRIME.(2,11,CPRIM,TC,N)
NEXTT = TC-Q/CPRIM
WHENEVER .ABS.(Q/TT).G.ERROR1,TRANSFER TO LCC(3)
TB = TC
TEXH = TC
EXECUTE ENTHAL.(2,11,HB,TB,N)
UB = HB-NB*R1*TB
TZ(TZZERO ) = THETA
TZ(TZZERO + 1 ) = TB
TZ(TZZERO + 2 ) = PEXH
TZ(TZZERO + 3 ) = VB
TZ(TZZERO + 4 ) = HB
TZ(TZZERO + 5 ) = SB
TZ(TZZERO + 6 ) = UB
TZ(TZZERO + 7 ) = NB
TZ(TZZERO + 8 ) = 0.
TZ(TZZERO + 9 ) = QTOT
TZ(TZZERO + 10) = 0.
TZ(TZZERO + 11) = WTOT
TZ(TZZERO + 12) = 0.
TZZERO = TZZERO + 13
PRINT FORMAT OUT,9,J1,J,360.,TB,P,VB,HB,SB,UB,NB,N(2)...N(11)
N(2) = N(2) +(N(9)+N(10)+N(8))/2.
N(3) = N(3) + N(9)/2.
N(7) = N(7)+(N(8)+N(11))/2.
TC = T5
F = N5/NB
TZ(TZZERO ) = 540.
TZ(TZZERO + 1 ) = T5
TZ(TZZERO + 2 ) = PEXH
TZ(TZZERO + 3 ) = VTOP
TZ(TZZERO + 4 ) = H5
TZ(TZZERO + 5 ) = S5
TZ(TZZERO + 6 ) = U5
TZ(TZZERO + 7 ) = N5
TZ(TZZERO + 8 ) = 0.
TZ(TZZERO + 9 ) = QTOT
TZ(TZZERO + 10) = 0.
TZ(TZZERO + 11) = WTOT
TZ(TZZERO + 12) = 0.
TZZERO = TZZERO + 12
WHENEVER STOP.G.1,T5=TB
WHENEVER .ABS.((TC-TB)/TC).G.ERROR1.AND.J3.L.STOP,TRANSFER TO
1LOOP1
```

```
FUNCTION RETURN
INTERNAL FUNCTION ARREA.(ZETA) = 3.1416*BORE*(STROKE/2.*(1.+
1COS.(ZETA*.01745))+ROD*(1.-SQRT.(1.-(SIN.(ZETA*.01745)*STROKE
2/2./ROD).P.2))+D1)+ATOT(1)
INTERNAL FUNCTION VOLUME.(ZETA) = .0007854*BORE*BORE*(STROKE/
12.*(1.+COS.(ZETA*.01745))+ROD*(1.-SQRT.(1.-(STROKE/2.*SIN.(
2ZETA*.01745)/ROD).P.2)))+ VTOP
VECTOR VALUES OUT =$1H0,3I3,S2,3F10,3,F15.8,4E15.8/(6E20.8)*$
INTERNAL FUNCTION(PR,AREA,S)
ENTRY TO QTRANS.
S = RPM/3000.*STROKE
S = 2.1*SQRT.(PR*TB)*S.P..33/36000.
FUNCTION RETURN DT*S*(AREA*(TB-TW)+ATOT(3))*(TB-TPISTN))*QFACT
END OF FUNCTION
INTERNAL FUNCTION(TEMP)
ENTRY TO EQUIL.
WHENEVER TEMP.LE. 1500.,FUNCTION RETURN
TR = TEMP/1500.
RT = 1./TR
B(1) = (ELOG.(TR)+RT-1.)/1.9872
B(2) = 750.00*(TR+RT-2.)/1.9872
B(3) = 375000.00*(TR*TR+2.*RT-3.)/1.9872
B(4) = 281250000.*(TR*TR*TR+3.*RT-4.)/1.9872
B(7) = (1.-RT)/2980.785
K(4) = 0.
THROUGH LOOP7, FOR VALUES OF I = 1,2,3,4,7
LOOP7 K(4) = K(4)+(AH(6,I)+.5*AH(2,I)-AH(4,I) )*B(I)
K(4) = EXP.(AH(4,8) +K(4))
K(5) = 0.
THROUGH LOOP8, FOR VALUES OF I = 1,2,3,4,7
LOOP8 K(5) = K(5)+(AH(7,I)+.5*AH(2,I)-AH(5,I))*B(I)
K(5) = EXP.(AH(5,8) +K(5))
K(8) = 0.
THROUGH LOOP9, FOR VALUES OF I = 1,2,3,4,7
LOOP9 K(8) = K(8)+(.5*(AH(2,I)+AH(7,I))-AH(8,I))*B(I)
K(8) = EXP.(AH(8,8) +K(8))
K(7) = 0.
THROUGH LOOP10, FOR VALUES OF I = 1,2,3,4,7
LOOP10 K(7) = K(7)+(.5*AH(7,I)-AH(11,I))*B(I)
K(7) = EXP.(AH(7,8) +K(7))
K(9) = 0.
THROUGH LOOP11, FOR VALUES OF I = 1,2,3,4,7
LOOP11 K(9) = K(9)+(.5*(AH(2,I)+AH(3,I))-AH(9,I))*B(I)
K(9) = EXP.(AH(9,8) +K(9))
K(2) = 0.
THROUGH LOOP12, FOR VALUES OF I = 1,2,3,4,7
LOOP12 K(2) = K(2)+(.5*AH(2,I)-AH(10,I))*B(I)
K(2) = EXP.(AH(2,8) +K(2))
FUNCTION RETURN
END OF FUNCTION
INTERNAL FUNCTION(N1,N2,PROPTY,TEMP,BLANK)
ENTRY TO SPRIME.
B(1) = 1./TEMP
B(2) = 1.
B(3) = TEMP
B(4) = TEMP*TEMP
TRANSFER TO MATRIX.
ENTRY TO HPRIME.
```



```
B(1) = 1.
B(2) = TEMP
B(3) = TEMP*TEMP
B(4) = TEMP*TEMP*TEMP
TRANSFER TO MATRIX
ENTRY TO ENTHAL.
I = 5
PROPTY = 0.
WHENEVER TEMP.G.1500., TRANSFER TO HIGHT
TR = TEMP/298.16
B(1) = 298.16*(TR - 1.)
B(2) = 44449.693*(TR*TR - 1.)
B(3) = 8835413.6*(TR*TR*TR - 1.)
B(4) = 1975775200.*(TR*TR*TR*TR - 1.)
B(5) = 1.
TRANSFER TO LOW
HIGHT TR = TEMP/1500.
B(1) = 1500.*(TR - 1.)
B(2) = 1125000.*(TR*TR - 1.)
B(3) = 1125000000.*(TR*TR*TR - 1.)
B(4) = .1265625E13*(TR*TR*TR*TR - 1.)
B(5) = 1.
TRANSFER TO HIGH
MATRIX I = 4
PROPTY = 0.
WHENEVER TEMP .G. 1500., TRANSFER TO HIGH
LOW THROUGH SUMB, FOR I2 = N1,1,I2.G.N2
TA = 0.
THROUGH SUMA, FOR I3 = 1,1,I3.G.I
SUMA TA = TA + B(I3)*AL(I2,I3)
SUMB PROPTY = PROPTY + N(I2)*TA
FUNCTION RETURN
HIGH THROUGH ADDA, FOR I2 = N1,1,I2.G.N2
TA = 0.
THROUGH ADDB, FOR I3 = 1,1,I3.G.I
ADDB TA = TA + B(I3)*AH(I2,I3)
ADDA PROPTY = PROPTY + N(I2)*TA
FUNCTION RETURN
END OF FUNCTION
INTERNAL FUNCTION (N1,N2,PROPTY,TEMP,MOLES,PTOTAL,BLANK)
ENTRY TO ENTROP.
PROPTY = 0.
WHENEVER TEMP.G.1500., TRANSFER TO BIGHT
TR = TEMP/298.16
B(1) = ELOG.(TR)
B(2) = 298.16*(TR - 1.)
B(3) = 44449.693*(TR*TR - 1.)
B(4) = 8835413.6*(TR*TR*TR - 1.)
B(6) = 1.
THROUGH BUMB, FOR I2 = N1,1,I2.G.N2
WHENEVER N(I2) .E. 0.,TRANSFER TO BUMB
TA = 0.
THROUGH BUMA, FOR VALUES OF I3 = 1,2,3,4,6
BUMA TA = TA + B(I3)*AL(I2,I3)
PROPTY = PROPTY + N(I2)*(TA - 1.9872*ELOG.(N(I2)*PTOTAL/MOL
1ES))
BUMB CONTINUE
FUNCTION RETURN
```

```

BIGHT TR = TEMP/1500.
      B(1) = ELOG.(TR)
      B(2) = 1500.*(TR - 1.)
      B(3) = 1125000.*(TR*TR - 1.)
      B(4) = 112500000.*(TR*TR*TR - 1.)
      B(6) = 1.
      THROUGH BDDA, FOR I2 = N1,1,I2.G.N2
      WHENEVER N(I2) .E. 0.,TRANSFER TO BDDA
      TA = 0.
      THROUGH BDDB, FOR VALUES OF I3 = 1,2,3,4,6
BDDB  TA = TA + B(I3)*AH(I2,I3)
      PROPTY = PROPTY + N(I2)*(TA - 1.9872*ELOG.(N(I2)*PTOTAL/MOL
1ES))
BDDA  CONTINUE
      FUNCTION RETURN
      END OF FUNCTION
      INTERNAL FUNCTION(TEMP,VOL,MOLES,PHH,BLANK)
      ENTRY TO AMOUNT.
      P(1) = 0.
      FA = TEMP*R/VOL
      WHENEVER TEMP .LE.1500.,TRANSFER TO H2OGAS
      WHENEVER PHH.LE.0.,PHH = 1.
      NEXTL = PHH
LOOP1  L = NEXTL
      S1 = L/K(8) + 1./K(7)
      S2 = 4.*(1. + L/K(5))
      S3 = SQRT.(L*L/K(7)*K(9)+8.*FA*NN)
      S4 = SQRT.(S1*S1 + 2.*S2*FA*NH)
      M = (S4 - S1)/S2
      S = S3/4. - L/4./K(9)
      DNWRL = (L/K(9)/S3-1.)/4./K(9)
      DMWRL = -(S2*(1./K(8)-(S1/K(8)+4.*FA*NH/K(5))/S4)+(S4-S1)*4./
1K(5))/S2/S2
      P(2) = L*L
      P(3) = S*S
      P(4) = NC*FA/(K(4)/L + 1.)
      P(5) = L*M*M/K(5)
      P(6) = P(4)*K(4)/L
      P(7) = M*M
      P(8) = L*M/K(8)
      P(9) = S*L/K(9)
      P(10) = L/K(2)
      P(11) = M/K(7)
      FL = 2.*(P(2) + P(4))+P(5) + P(6)+P(9)+P(8)+P(10) - FA*NO
      FPRIML = FA*NC*K(4)/(L+K(4))/(L+K(4))+4.*L*M*M/K(5)+ M/K(8)+
1S/K(9)+1./K(2)+DMWRL*(L/K(8)+2.*L*M/K(5) +DNWRL/K(9)*L
      M = -FL/L/FPRIML
      WHENEVER .ABS.(M) .L.16.
      NEXTL = L*EXP.(M)
      OTHERWISE
      NEXTL = L*EXP.(M/.ABS.(M))
      END OF CONDITIONAL
      WHENEVER .ABS.((NEXTL-L)/L).G.ERROR2,TRANSFER TO LOOP1
      PHH = L
      TRANSFER TO LOC2
H2OGAS  WHENEVER (NC+NH/4.)/NO.LE.0.5,TRANSFER TO SIMPLE
      S1 = NO/NC-1.
      S3 = (1.+NH/NC/.7494)/S1-1.6688
      L = (S3+SQRT.(S3*S3+6.6752/S1))/2.
      P(2) = 0.
      P(4) = FA*NC/L
      P(5) = FA*(NO-NC)-P(4)
      P(6) = .ABS.(FA*NC-P(4))
      P(7) = .ABS.(FA*NH/2. -P(5))
      TRANSFER TO HERR
SIMPLE  P(2) = .ABS.(FA*(NO/2.-NC-NH/4.))
      P(4) = FA*NC
      P(5) = FA*NH/2.
      P(6) = 0.
      P(7) = 0.
HERR    P(8) = 0.
      P(9) = 0.
      P(10) = 0.
      P(11) = 0.
      P(3) = FA*NN/2.
      PHH = 1.
LOC2    MOLES = 0.
      THROUGH LOOP2, FOR I = 1,1,I.G.11
      N(I) = P(I)/FA
LOOP2   MOLES = MOLES + N(I)
      FUNCTION RETURN
      END OF FUNCTION
      END OF FUNCTION

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SCOMPILE MAD,PRINT OBJECT ,PUNCH OBJECT

30UTP201

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EXTERNAL FUNCTION (F)
PROGRAM COMMON AL, AH, P, K, R1, NC, NH, NO, NN, R, J2, RE, KK, X, TW,
1QFACT, EA, RATIO, BO, RHO, SPEED, ERROR2, CPF, TPISTN, ATOT, XX, XNUM,
2BORE, STROKE, ROD, VTOP, VBOT, PMAN, PEXH, TMAN, RPM, N, THETA, ERROR1,
3J3, STOP, T5, H5, S5, U5, N5, QTOT, WTOT, F0, NEXTV, HCOMB, INC, Z, C,
4 TZ, TX, COMDG, PMAX, NAME, CR, PI
INTEGER NUM2, INC, RE, NAME, TZZERO
DIMENSION AL(87,ALDIM), AH(87,ALDIM), RE(12), X(10), KK(5), QFACT
1(5), P(11), N(11), K(11), XX(105, XDIM), NAME(2), ATOT(5),
2 TZ(1980, TDIM), TX( 780, TDIM)
EQUIVALANCE (TZZERO, TZ)
VECTOR VALUES TDIM = 2, 1, 13
VECTOR VALUES ALDIM = 2, 0, 8
VECTOR VALUES XDIM = 2, 0, 21
ENTRY TO OUTPUT.
PRINT FORMAT INTRO, NAME...NAME(2), X, BORE/2.54, EA, TPISTN, STRO
1KE/2.54, TW, ROD/2.54, HCOMB, TMAN, VTOP*61.03, PMAN, 61.03*(VBOT-VT
2OP), PEXH, CR, PI
WPUMP = (VBOT-VTOP)*(PMAN-PEXH)*24.2179
WTOT = WTOT + WPUMP
EFF = WTOT/HCOMB*100./F0
PRINT FORMAT INTOR1, RPM, INC, ERROR1*100., KK, QFACT, RE(4)...RE(
112)
IMEP = WTOT/(VBOT-VTOP)/24.2179
VOLEFF = F0*(C*X+X)/(VBOT-VTOP)*R/PMAN*TMAN
PRINT FORMAT RESULT, WPUMP, WTOT, QTOT, EFF, IMEP, F, VOLEFF
VECTOR VALUES RESULT = $1H2, $20, 18HSUMMARY OF RESULTS/1H0/S6,
112HPUMPING WORKS18, E15.8, 4H CAL/S6, 14HNET CYCLE WORKS16, E15.8
2, 4H CAL/S6, 19HNET HEAT TRANSFEREDS11, E15.8, 4H CAL/S6, 28HINDIC
3ATED THERMAL EFFICIENCY2, E15.8, 8H PERCENT/S6, 23HMEAN EFFECTI
4VE PRESSURE5, E15.8, 4H ATM/S6, 24HRESIDUAL WEIGHT FRACTION, S6,
5E15.8/S6, 21HVOLUMETRIC EFFICIENCY9, E15.8, 8H PERCENT*S
NUM2 = TZZERO/13
PRINT FORMAT HEADER
VECTOR VALUES HEADER = $1H1/6HOTHETAS5, 1HTS6, 1HPS5, 1HVS9, 1HH,
1 S9, 1HSS9, 1HUS9, 1HNS9, 1HQS8, 4HQOTS7, 1HWS8, 4HWTOTS7, 2HHS*5
PRINT FORMAT TABLE1, TZ(1,1)...TZ(NUM2,13)
VECTOR VALUES TABLE1 = $1H F6.2, F8.2, F7.2, F8.5, 9F10.5*S
PRINT FORMAT HEADR1
VECTOR VALUES HEADR1 = $1H1/6HOTHETAS5, 1HTS6, 1HPS5, 1HVS9, 1HH,
1 S9, 1HSS9, 1HUS9, 1HNS5, 50HFLAME SPEED, TEMP., AND EXTENT MASS-
2FR. BURN.-VOL.*5
PRINT FORMAT TABLE1, TX(1,1)...TX(INC,13)
FUNCTION RETURN
VECTOR VALUES INTRO = $1H1, $35, 49HA MODEL SIMULATING THE INTE
1RNAL-COMBUSTION ENGINE/7HOENGINES33, 4HFUELS36, 20HOPERATING CO
2NDITIONS/28HOFUR STROKE AUTOMOTIVE TYPES12, 3C6, S22, 14HAIR-FU
3EL RATIO F9.2/5H BORES8, F10.2, 4H IN. S13, 17HACTIVATION ENERGY
4F10.2, 24H CAL./MOLE PISTON TEMP F12.2, 2H K/7H STROKES6, F10.
52, 4H IN. S13, 11HENTHALPY OFS29, 9HWALL TEMPF14.2, 2H K/4H RODS9,
6F10.2, 4H IN. S13, 10HCOMBUSTIONF17.2, 22H CAL/MOLE MAN. TEMP
7F14.2, 2H K/12H CLEAR. VOL. F11.2, 7H CU. IN. S50, 8HMAN. PR. F15.2,
84H ATM/11H SWEPT VOL. F12.2, 7H CU. IN. S50, 8HEXH. PR. F15.2, 4H AT
9M/18H COMPRESSION RATIOF5.2, 7H TO ONES0, 10HSPARK ADV. F13.2*S
VECTOR VALUES INTOR1 = $580, 5HSPEEDS10, F8.2, 4H RPM/ 2IHOCOMBU
1STION OCCURS IN 13, 34H INCREMENTS. ERRORS ARE LESS THAN F10.
25, 33H PERCENT. FLAME SPEED FACTOR IS E15.8/24H HEAT TRANSFER
3 FACTOR IS F10.5, S3, 10C6*S

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SCOMPILE MAD,PRINT OBJECT,PUNCH OBJECT

4QTRAN01

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EXTERNAL FUNCTION(TB, TU, PR, DT, ANS, HB)
DIMENSION AL(87,ALDIM), AH(87,ALDIM), RE(12), X(10), KK(5), QFACT
1(5), P(11), K(11), XX(105, XDIM), A(5)
PROGRAM COMMON AL, AH, P, K, R1, NC, NH, NO, NN, R, J2, RE, KK, X, TW, QFAC
1T, EA, RATIO, BO, RHO, SPEED, ERROR2, CPF, TPISTN, A, XX, XNUM, BORE, STRO
2KE, ROD, VTOP, VBOT, PMAN, PEXH, TMAN, RPM
VECTOR VALUES ALDIM = 2, 0, 8
VECTOR VALUES XDIM = 2, 0, 21
ENTRY TO QTRAN2.
S = RPM/3000.*STROKE
HB = 2.1*SQRT.(PR*TB)*S.P..33/36000.
HU = 2.1*SQRT.(PR*TU)*S.P..33/36000.
FUNCTION RETURN DT*(HB*(ANS(1)*(TB-TW)+ANS(3)*(TB-TPISTN)) +
1HU*(ANS(2)*(TU-TW)+ANS(4)*(TU-TPISTN)))*QFACT
END OF FUNCTION

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\$COMPILE MAD,PRINT OBJECT,PUNCH OBJECT

6ANS0001

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EXTERNAL FUNCTION (VB,VTOT,ANS,D1,D2)
PROGRAM COMMON AL, AH,P,K,R1,NC,NH,NO,NN,R,J2,RE,KK,X,TW,QFAC
1T,EA,RATIO,BO,RHO,SPEED,ERROR2,CPF,TPISTN,A,XX,XNUM,BORE,
2STROKE,ROD,VTOP,VBOT,PMAN,PEXH,TMAN,RPM,N,THETA,ERROR1
ERASABLE LOP,L1P,L2P,DER,DVWRF,NEXTF,I0,XMUM,F,DL,I1,I2,I3,
1L0,L1,L2,AW,AWB,AWU,S,Q,QQ,DIF
INTEGER I,I0,I1,I2,XNUM,I3,J2
DIMENSION AL(87,ALDIM),AH(87,ALDIM),RE(12),X(10),KK(5),QFACT
1(5),P(11),K(11),XX(105,XDIM),A(5),N(11)
VECTOR VALUES ALDIM = 2,0,8
VECTOR VALUES XDIM = 2,0,21
ENTRY TO AREA.
J2 = 0
XDIM(2) = XNUM
XMUM = XNUM - 1
DL = STROKE/2.*(1.+COS.(THETA*.01745)) + ROD*(1.-SQRT.(1.-
1(STROKE/2.*SIN.(THETA*.01745)/ROD),P,2))
WHENEVER THETA .G. 180.,TRANSFER TO DOWN
F = VB/VTOT*XMUM
WHENEVER VB/VTOT .L..03
ANS(5) = (VB/.0041888).P..33
OTHERWISE
I1 = F + .5
WHENEVER I1.LE.1,I1 = 2
WHENEVER I1.GE.XNUM-1,I1 = I1 - 1
I2 = I1 + 1
I3 = I2 + 1
I0 = I1 - 1
L0 = (F-I1)*(F-I2)/2.
L1 = (I0-F)*(F-I2)
L2 = (F-I0)*(F-I1)/2.
ANS(5) = A(5)*(L0*XX(5,I1) + L1*XX(5,I2) + L2*XX(5,I3))
END OF CONDITIONAL
ANS = VB
F = ANS(5)/A(5)*XMUM
TRANSFER TO FINISH
NEXTF = ANS(5)/A(5)*XMUM
F = NEXTF
J2 = J2 + 1
ANS(5) = F*A(5)/XMUM
I1 = F + .5
WHENEVER I1.LE.0,I1 = 1
WHENEVER I1.GE.XNUM-1,I1 = XNUM-2
I2 = I1 + 1
I3 = I2 + 1
I0 = I1 - 1
L0 = (F-I1)*(F-I2)/2.
L1 = (I0-F)*(F-I2)
L2 = (F-I0)*(F-I1)/2.
WHENEVER J2.LE.5
LOP = F-(I1+I2)/2.
L1P = I0+I2-2.*F
L2P = F-(I0+I1)/2.
ANS = VTOP*(L0*XX(4,I1) + L1*XX(4,I2) + L2*XX(4,I3))
WHENEVER ANS(5).LE.D2
ANS = ANS + .0031416*ANS(5)*ANS(5)*DL
DER = .0062832*F*DL*(A(5)/XMUM).P.2
OTHERWISE
```

DOWN
BACK

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ANS = ANS + DL*ABS(A(1)*(LO*XX(1,I1)+L1*XX(1,I2)+L2*XX(1,I3
1))) / 1000.
DER = DL*A(1)*(LOP*XX(1,I1)+L1P*XX(1,I2)+L2P*XX(1,I3)) / 1000.
END OF CONDITIONAL
DVWRF = VTOP*(LOP*XX(4,I1)+L1P*XX(4,I2)+L2P*XX(4,I3))
Q = VB-ANS
NEXTF = F + Q / (DVWRF+DER)
DIF = ABS(NEXTF - F)
OTHERWISE
ANS = VTOP*(LO*XX(4,I1) + L1*XX(4,I2) + L2*XX(4,I3))
WHENEVER ANS(5) LE D2
ANS = ANS + .0021416*ANS(5)*ANS(5)*DL
OTHERWISE
ANS = ANS + DL*ABS(A(1)*(LO*XX(1,I1)+L1*XX(1,I2)+L2*XX(1,I3
1))) / 1000.
END OF CONDITIONAL
QQ = VB-ANS
WHENEVER Q/QQ LE .0. DIF = DIF/2.
NEXTF = F+DIF*Q/ABS(QQ)
Q = QQ
END OF CONDITIONAL
WHENEVER ABS((NEXTF-F)/F) GE .0001,TRANSFER TO BACK
FINISH
I1 = F + 1.
I2 = I1 + 1
I0 = I1 - 1
LO = I1 - F
LI = F - I0
AW = 3.1416*BORE*(DL + D1)
WHENEVER ANS(5) GE .99*A(5)
AWB = AW
AWU = 0.
TRANSFER TO END
END OF CONDITIONAL
WHENEVER ANS(5) GE D2
S = (BORE - D2 + ANS(5))/2.
AWB = AW*2./3.1416*ATAN(SQRT((S-BORE/2.)*(S-BORE/2.+D2
1)/S/(S-ANS(5))))
AWU = AW - AWB
OTHERWISE
AWU = AW
AWB = 0.
END OF CONDITIONAL
END
ANS(1) = A(1)*(LO*XX(1,I1) + L1*XX(1,I2))
ANS(2) = A(1) - ANS(1)
ANS(1) = ANS(1) + AWB
ANS(2) = ANS(2) + AWU
ANS(3) = A(3)*(LO*XX(3,I1) + L1*XX(3,I2))
ANS(4) = A(3) - ANS(3)
FUNCTION RETURN
END OF FUNCTION
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SCOMPILE MAD,PRINT OBJECT,PUNCH OBJECT                                4ENTHL01  
EXTERNAL FUNCTION(N1,N2,PROPT,TEMP,N)  
DIMENSION AL(87,ALDIM),AH(87,ALDIM),X(10),K(11),P(11),B(5),  
1RE(12),KK(5),QFACT(5)  
ERASABLE B,I,I2,I3,TR,TA,PROPTY  
PROGRAM COMMON AL,AH,P,K,R1,NC,NH,NO,NN,R,J2,RE,KK,X,TW,QFACT  
1,EA,RATIO,BO,RHO,SPEED,ERROR2  
VECTOR VALUES ALDIM = 2,0,8  
ENTRY TO SPRIME,  
B(1) = 1./TEMP  
B(2) = 1.  
B(3) = TEMP  
B(4) = TEMP*TEMP  
TRANSFER TO MATRIX  
ENTRY TO HPRIME,  
B(1) = 1.  
B(2) = TEMP  
B(3) = TEMP*TEMP  
B(4) = TEMP*TEMP*TEMP  
TRANSFER TO MATRIX  
ENTRY TO ENTHAL,  
I = 5  
PROPTY = 0.  
WHENEVER TEMP.G.1500., TRANSFER TO HIGHT  
TR = TEMP/298.16  
B(1) = 298.16*(TR - 1.)  
B(2) = 44449.693*(TR*TR - 1.)  
B(3) = 8834413.6*(TR*TR*TR - 1.)  
B(4) = 1975775200.*(TR*TR*TR*TR - 1.)  
B(5) = 1.  
TRANSFER TO LOW  
HIGHT TR = TEMP/1500.  
B(1) = 1500.*(TR - 1.)  
B(2) = 1125000.*(TR*TR - 1.)  
B(3) = 1125000000.*(TR*TR*TR - 1.)  
B(4) = .1265625E13*(TR*TR*TR*TR - 1.)  
B(5) = 1.  
TRANSFER TO HIGH  
MATRIX I = 4  
PROPTY = 0.  
WHENEVER TEMP .G. 1500., TRANSFER TO HIGH  
LOW THROUGH SUMB, FOR I2 = N1,1,I2.G.N2  
TA = 0.  
THROUGH SUMA, FOR I3 = 1,1,I3.G.I  
SUMA TA = TA + B(I3)*AL(I2,I3)  
SUMB PROPTY = PROPTY + N(I2)*TA  
PROPT = PROPTY  
FUNCTION RETURN  
HIGH THROUGH ADDA, FOR I2 = N1,1,I2.G.N2  
TA = 0.  
THROUGH ADDB, FOR I3 = 1,1,I3.G.I  
ADDB TA = TA + B(I3)*AH(I2,I3)  
ADDA PROPTY = PROPTY + N(I2)*TA  
PROPT = PROPTY  
FUNCTION RETURN  
INTEGER I2,I3,N1,N2,I  
END OF FUNCTION  
-----
```

```
SCOMPILE MAD,PRINT OBJECT, PUNCH OBJECT                                4ENTRP01
EXTERNAL FUNCTION (N1,N2,PROPTY,TEMP,MOLES,PTOTAL,N)
DIMENSION AL(87,ALDIM),AH(87,ALDIM),X(10),B(6),K(11),P(11),
1RE(12),KK(5),QFACT(5),SPACE(5)
PROGRAM COMMON AL,AH,P,K,R1,NC,NH,NO,NN,R,J2,RE,KK,X,TW,QFACT
1,EA,RATIO,B0,RHO,SPEED,ERROR1
ERASABLE SPACE,B,I2,I3,TR,TA,PROPTY
VECTOR VALUES ALDIM = 2,0,8
ENTRY TO ENTROP,
PROPTY = 0.
WHENEVER TEMP.G.1500., TRANSFER TO BIGHT
TR = TEMP/298.16
B(1) = ELOG.(TR)
B(2) = 298.16*(TR - 1.)
B(3) = 44449.693*(TR*TR - 1.)
B(4) = 8835413.6*(TR*TR*TR - 1.)
B(6) = 1.
THROUGH BUMB, FOR I2 = N1,1,I2.G.N2
WHENEVER N(I2) .E. 0.,TRANSFER TO BUMB
TA = 0.
THROUGH BUMA, FOR VALUES OF I3 = 1,2,3,4,6
BUMA TA = TA + B(I3)*AL(I2,I3)
PROPTY = PROPTY + N(I2)*(TA -1.9872*ELOG.(N(I2)*PTOTAL/MOL
1ES))
BUMB CONTINUE
PROPTT = PROPTY
FUNCTION RETURN
BIGHT TR = TEMP/1500.
B(1) = ELOG.(TR)
B(2) = 1500.*(TR - 1.)
B(3) = 1125000.*(TR*TR - 1.)
B(4) = 1125000000.*(TR*TR*TR - 1.)
B(6) = 1.
THROUGH BDDA, FOR I2 = N1,1,I2.G.N2
WHENEVER N(I2) .E. 0.,TRANSFER TO BDDA
TA = 0.
THROUGH BDDB, FOR VALUES OF I3 = 1,2,3,4,6
BDDB TA = TA + B(I3)*AH(I2,I3)
PROPTY = PROPTY + N(I2)*(TA -1.9872*ELOG.(N(I2)*PTOTAL/MOL
1ES))
BDDA CONTINUE
PROPTT = PROPTY
FUNCTION RETURN
INTEGER I2,I3,N1,N2
END OF FUNCTION
```

```
SCOMPILE MAD,PRINT OBJECT,PUNCH OBJECT                                4EQUIL01
EXTERNAL FUNCTION(TEMP)
DIMENSION AL(87,ALDIM),AH(87,ALDIM),X(10),K(11),P(11),B(7),
IRE(12),KK(5),QFACT(5),SPACE(5)
PROGRAM COMMON AL,AH,P,K,R1,NC,NH,NO,NN,R,J2,RE,KK,X,TW,QFACT
I,EA,RATIO,B0,RHO,SPEED,ERROR1
VECTOR VALUES ALDIM = 2,0,8
ERASABLE SPACE,B,TR,RT,I
ENTRY TO EQUIL.
WHENEVER TEMP.LE. 1500.,FUNCTION RETURN
TR = TEMP/1500.
RT = 1./TR
B(1) = (ELOG.(TR)+RT-1.)/1.9872
B(2) = 750.00*(TR+RT-2.)/1.9872
B(3) = 375000.00*(TR*TR+2.*RT-3.)/1.9872
B(4) =281250000.*(TR*TR*TR+3.*RT-4.)/1.9872
B(7) = (1.-RT)/2980.785
K(4) = 0.
THROUGH LOOP7,FOR VALUES OF I = 1,2,3,4,7
LOOP7  K(4) = K(4)+(AH(6,I)+.5*AH(2,I)-AH(4,I))*B(I)
      K(4) = EXP.(AH(4,8) +K(4))
      K(5) = 0.
THROUGH LOOP8,FOR VALUES OF I = 1,2,3,4,7
LOOP8  K(5) = K(5)+(AH(7,I)+.5*AH(2,I)-AH(5,I))*B(I)
      K(5) = EXP.(AH(5,8) +K(5))
      K(8) = 0.
THROUGH LOOP9,FOR VALUES OF I = 1,2,3,4,7
LOOP9  K(8) = K(8)+(.5*(AH(2,I)+AH(7,I))-AH(8,I))*B(I)
      K(8) = EXP.(AH(8,8) +K(8))
      K(7) = 0.
THROUGH LOOP10,FOR VALUES OF I = 1,2,3,4,7
LOOP10 K(7) = K(7)+(.5*AH(7,I)-AH(11,I))*B(I)
      K(7) = EXP.(AH(7,8) +K(7))
      K(9) = 0.
THROUGH LOOP11,FOR VALUES OF I = 1,2,3,4,7
LOOP11 K(9) = K(9)+(.5*(AH(2,I)+AH(3,I))-AH(9,I))*B(I)
      K(9) = EXP.(AH(9,8) +K(9))
      K(2) = 0.
THROUGH LOOP12,FOR VALUES OF I = 1,2,3,4,7
LOOP12 K(2) = K(2)+(.5*AH(2,I)-AH(10,I))*B(I)
      K(2) = EXP.(AH(2,8) +K(2))
FUNCTION RETURN
INTEGER I
END OF FUNCTION
```



```

$COMPILE MAD,PUNCH OBJECT,PRINT OBJECT
                                                    3AMONT01
EXTERNAL FUNCTION(TEMP,VOL,MOLES,PHH,N)
DIMENSION AL(87,ALDIM),AH(87,ALDIM),X(10),K(11),P(11),
1RE(12),KK(5),QFACT(5),SPACE(5)
PROGRAM COMMON AL,AH,P,K,R1,NC,NH,NO,NN,R,J1,RE,KK,X,TW,QFACT
1,EA,RATIO,BO,RHO,SPEED,ERROR1
ERASABLE SPACE,I,M,S,DNWRL,DMWRL,FL,Q,FPRIML,S4,S3,S2,S1,L,
1NEXTL,FA
VECTOR VALUES ALDIM = 2,0,8
ENTRY TO AMOUNT.
P(1) = 0.
FA = TEMP*R/VOL
WHENEVER TEMP.LE.1500.,TRANSFER TO H2OGAS
WHENEVER PHH.LE.0.,PHH = 1.
NEXTL = PHH
J1 = 0
LOOP1 L = NEXTL
J1 = J1 + 1
S1 = L/K(8) + 1./K(7)
S2 = 4.*(1. + L/K(5))
S3 = SQRT.(L*L/K(9)/K(9)+8.*FA*NN)
S4 = SQRT.(S1*S1 + 2.*S2*FA*NH)
M = (S4 - S1)/S2
S = S3/4. - L/4./K(9)
DNWRL = (L/K(9)/S3-1.)/4./K(9)
DMWRL = -(S2*(1./K(8)-(S1/K(8)+4.*FA*NH/K(5)))/S4)+(S4-S1)*4./
1K(5))/S2/S2
P(2) = L*L
P(3) = S*S
P(4) = NC*FA/(K(4)/L + 1.)
P(5) = L*M*M/K(5)
P(6) = P(4)*K(4)/L
P(7) = M*M
P(8) = L*M/K(8)
P(9) = S*L/K(9)
P(10) = L/K(2)
P(11) = M/K(7)
FL = 2.*(P(2) + P(4))+P(5) + P(6)+P(9)+P(8)+P(10) - FA*NO
FPRIML = FA*NC*K(4)/(L+K(4))/(L+K(4))+4.*L+M*M/K(5)+ M/K(8)+
1S/K(9)+1./K(2)+DMWRL*(L/K(8)+2.*L*M/K(5)) +DNWRL/K(9)*L
Q = -FL/L/FPRIML
WHENEVER .ABS.(Q) .L.16.
NEXTL = L*EXP.(Q)
OTHERWISE
NEXTL = L*EXP.(Q/.ABS.(Q))
END OF CONDITIONAL
WHENEVER .ABS.((NEXTL-L)/L).G.ERROR1,TRANSFER TO LOOP1
PHH = L
TRANSFER TO LOC2
H2OGAS WHENEVER (NC+NH/4.)/NO.LE..5,TRANSFER TO SIMPLE
S1 = NO/NC-1.
S3 = (1.+NH/NC/.7494)/S1-1.6688
L = (S3+SQRT.(S3*S3+6.6752/S1))/2.
P(2) = 0.
P(4) = FA*NC/L
P(5) = FA*(NO-NC)-P(4)
P(6) = .ABS.(FA*NC-P(4))
P(7) = .ABS.(FA*NH/2. -P(5))
TRANSFER TO HERR

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SIMPLE      P(2) = .ABS.(FA*(NO/2.-NC-NH/4.))
            P(4) = FA*NC
            P(5) = FA*NH/2.
            P(6) = 0.
            P(7) = 0.
HERR       P(8) = 0.
            P(9) = 0.
            P(10) = 0.
            P(11) = 0.
            P(3) = FA*NN/2.
            PHH = 1.
LOC2       MOLES = 0.
            THROUGH LOOP2, FOR I = 1,1,I.G.11
            N(I) = P(I)/FA
LOOP2      MOLES = MOLES + N(I)
            FUNCTION RETURN
            INTEGER I,J1
            END OF FUNCTION

$COMPILE MAD ,PUNCH OBJECT,PRINT OBJECT                                4FLAME01
            EXTERNAL FUNCTION (T0,TF,ST,R2)
            DIMENSION AL(87,ALDIM),AH(87,ALDIM),X(10),K(11),P(11),
            1RE(12),KK(5),QFACT(5),LOC(3)
            VECTOR VALUES ALDIM = 2,0,8
            PROGRAM COMMON AL,AH,P,K,R1,NC,NH,NO,NN,R,J2,RE,KK,X,TW,QFACT
            1,EA,RATIO,B0,RHO,SPEED,ERROR1,CPF
            ENTRY TO STURB.
            WHENEVER .ABS.(X-1.).GE..05
            J = 1
            OTHERWISE
            J = 2
            S2 = X
            X = .95
            END OF CONDITIONAL
LOC1       WHENEVER X.GE.1.05
            PHI = 1./X
            OTHERWISE
            PHI = X
            END OF CONDITIONAL
            S1 = R1*TF/EA*TF/(TF-T0)
            AE = S1*T0/TF
            BE = B0*T0/TF*(1.-PHI*(1.-S1))
            LAMDA = (TF.P..67)*(CPF+1.25*R2)
            WDT = AE*BE*SQRT.(TF)*S1* EXP.(-EA/R1/TF)
            BRAKT = 1.+1.25*R2/CPF
            ST = KK*BRAKT*RATIO*SQRT.(LAMDA*WDT/RHO/CPF)
            TRANSFER TO LOC(J)
LOC(1)     WHENEVER RE(7).E.$HELP $
            KK(1) = SPEED/ST
            KK = KK(1)
            ST = SPEED
            RE(7) = 0
            END OF CONDITIONAL
            FUNCTION RETURN
LOC(2)     J = 3
            X = 1.05
            Z = ST
            TRANSFER TO LOC1
LOC(3)     X = S2
            ST = Z+10.*(X-.95)*(ST-Z)
            TRANSFER TO LOC(1)
            INTEGER J,RE
            END OF FUNCTION

```

APPENDIX J

SPECIAL THERMODYNAMIC CONSIDERATIONS ASSOCIATED WITH INTER-RELATED STATE CHANGES OF TWO DISTINCT GASEOUS REGIONS

A. Isentropic Change of Volume of Each Region

Step 4 of the combustion process involves an isentropic change of volume of each of two gaseous regions. The two regions (burned and unburned) initially occupy a volume V and finally a volume V' . Pressure equilibrium is maintained between the regions. The fraction of the total volume occupied initially by each portion is known. It is desired to find what part of the total volume is occupied by the burned portion after the volume change.

The burned gas is presumed to be initially at P , T_b , V_b , and S_b whereas the unburned gas is at P , T_u , V_u , and S_u . After the volume change the burned gas is at P' , T'_b , V'_b , and S'_b and the unburned gas at P' , T'_u , V'_u , and S'_u . Let the ratio of the heat capacity at constant pressure to that at constant volume be k_u and k_b for the two regions at the initial volume and the difference between them $k_u - k_b = \delta$.

In the unburned gas, for this isentropic volume change,

$$\left(\frac{V'_u}{V_u}\right)^{k_u} = \frac{P}{P'}$$

and in the burned,

$$\left(\frac{V'_b}{V_b}\right)^{k_b} = \frac{P}{P'}$$

Equation these

$$\left(\frac{V'_u}{V_u}\right)^{k_u} = \left(\frac{V'_b}{V_b}\right)^{k_u+\delta} = \left(\frac{V'_b}{V_b}\right)^{k_u} \cdot \left(\frac{V'_b}{V_b}\right)^{\delta}$$

or

$$\frac{V'_u}{V_u} = \frac{V'_b}{V_b} \cdot \left(\frac{V'_b}{V_b}\right)^{\delta/k_u} = \frac{V'_b}{V_b} \cdot \kappa$$

where κ can be thought of as a correction factor to account for the differences in the heat capacity ratios.

Solving the above expression for V'_b and substitution $V' = V'_b + V'_u$

$$V'_b = \frac{V'}{1 + V_u/V_b \cdot \kappa}$$

The correction factor κ can be approximated by assuming that $\frac{V'_b}{V_b} = \frac{V'}{V}$.

Thus the final expression for the volume of the burned portion after the isentropic change of volume is

$$V'_b \approx \frac{V'}{1 + \left(\frac{V_u}{V_b}\right) \cdot \left(\frac{V'}{V}\right)^{\delta/k_u}} \quad (J-1)$$

B. Change of Volume with Known Values of Internal Energy and Entropy

Step 3 of the combustion process involves the recompression of the system to a specified final volume and internal energy while maintaining the entropy of the unburned portion constant. Pressure equilibrium is maintained between both portions. It is desired to estimate the pressure of the system after this compression.

The n_b moles of burned gas are presumed to be initially at P , T_b , V_b , and U_b and have heat capacity C_{P_b} . The n_u moles of unburned gas are presumed to be initially at P , T_u , V_u , and U_u and have heat capacity C_{P_u} . After the change in volume the burned gas has properties P , T'_b , V'_b ,

and U'_b and the unburned gas properties P , T'_u , V'_u , and U'_u . The initial total internal energy is $U = U_u + U_b$ and volume $V = V_u + V_b$. The final internal energy is $U' = U'_u + U'_b$ and volume is $V' = V'_u + V'_b$. These four quantities are known.

The change in internal energy of the unburned portion for this isentropic process is

$$U'_u - U_u = n_u C_{v_u} T_u \left[\left(\frac{P}{P'} \right)^{R/C_{p_u}} - 1 \right]$$

The change of internal energy in the burned portion is

$$\begin{aligned} U'_b - U_b &= n_b C_{v_b} \left[T'_b - T_b \right] = n_b C_{v_b} \left[\frac{P' V'_b}{n_b R} - T_b \right] \\ &= \frac{C_{v_b} P'}{R} (V' - V'_u) - n_b C_{v_b} T_b \end{aligned}$$

The total change of internal energy for the system is the sum of the changes in each portion and is

$$\begin{aligned} U' - U &= U'_b - U_b + U'_u - U_u \\ &= n_u C_{v_u} T_u \left(\frac{P}{P'} \right)^{R/C_{p_u}} - n_u T_u (C_{v_u} + C_{v_b}) - n_b T_b C_{v_b} + C_{v_b} \frac{P' V'}{R} \end{aligned}$$

Rearranging

$$P' = \frac{R}{V'} \left\{ \frac{U' - U}{C_{v_b}} + n_b T_b + n_u T_u + \frac{C_{v_u}}{C_{v_b}} n_u T_u \left[1 - \left(\frac{P}{P'} \right)^{R/C_{p_u}} \right] \right\}$$

Neglecting the bracketted term as being small compared to the other terms the expression for the final pressure becomes:

$$P' \cong \frac{R}{V'} \left\{ \frac{U' - U}{C_{v_b}} + n_b T_b + n_u T_u \right\} \quad (J-2)$$

Neglecting this term amounts to assuming that the entire internal energy decrease occurs in the burned portion.

APPENDIX K
CALCULATED DATA

The data sets presented are separated into three groups according to whether the flame propagation rate is normal, rapid, or reduced. These are referred to as Data Set A, B, and C respectively.

Each set of data presented consists of four pages. The first page describes the current operating conditions of the engine and presents the results of the calculations. The second and third pages present the total system thermodynamic properties and other information as a function of crank angle. The fourth page presents the thermodynamic properties of the burned gas region and other information during the course of the combustion process only as a function of crank angle. The data following are given in the following notation and units:

THETA	crank angle degrees, $0^\circ = \text{BDC}$
T	gas temperature, $^\circ\text{K}$
P	cylinder pressure, atm
V	volume, liters
H	enthalpy, cal
S	entropy, cal/ $^\circ\text{K}$
U	internal energy, cal
N	moles
Q	incremental heat transfer, cal
QTOT	total heat transfer, cal
W	incremental work, cal
WTOT	total work, cal

HB	heat transfer coefficient in burned gas region, $\text{cal/cm}^2\text{-}^\circ\text{K}$ (from Equation (5-18)),
FLAME SPEED	flame propagation rate, cm/sec (from Equation (4-24))
FLAME TEMPERATURE	effective flame temperature, $^\circ\text{K}$ (calculated in Step 7 of the combustion process)
EXTENT OF FLAME PROPAGATION	radius of spherical flame front, cm (corresponds to R'_0 in Figure 10-a)
MASS-FR. BURNED	mass fraction burned
VOL.-FR. BURNED	volume fraction occupied by burned gas

For all the data calculated using the analytical model in Data Sets A, B, and C, the exhaust temperature is 1000°K . During each cycle 0.00033 moles of fuel are inducted together with 0.0038 moles of O_2 and 0.0143 moles of N_2 . For the number of increments considered in the calculations about 25 minutes of computer time are required on the IBM 709 for each set of operating conditions.

DATA SET A

NORMAL FLAME PROPAGATION RATES

A MODEL SIMULATING THE INTERNAL-COMBUSTION ENGINE

ENGINE	FUEL	OPERATING CONDITIONS
FOUR STROKE AUTOMOTIVE TYPE	OCTANE C8H18	AIR-FUEL RATIO .92
BORE 3.94 IN.	ACTIVATION ENERGY 39000.00 CAL./MOLE	PISTON TEMP 500.00 K
STROKE 2.46 IN.	ENTHALPY OF COMBUSTION 1222770.00 CAL/MOLE	WALL TEMP 420.00 K
ROD 7.06 IN.		MAN. TEMP 320.00 K
CLEAR. VOL. 3.33 CU.IN.		MAN. PR. .99 ATM
SWEPT VOL. 30.00 CU.IN.		EXH. PR. 1.03 ATM
COMPRESSION RATIO 10.00 TO ONE		SPARK ADV. 16.00
		SPEED 2800.00 RPM

COMBUSTION OCCURS IN 39 INCREMENTS. ERRORS ARE LESS THAN .00100 PERCENT. FLAME SPEED FACTOR IS .13550000E 06
 HEAT TRANSFER FACTOR IS 1.00000 NORMAL RUN

SUMMARY OF RESULTS

PUMPING WORK	-.47611947E 00 CAL
NET CYCLE WORK	.14767102E 03 CAL
NET HEAT TRANSFERED	.70592567E 02 CAL
INDICATED THERMAL EFFICIENCY	.36569741E 02 PERCENT
MEAN EFFECTIVE PRESSURE	.12406213E 02 ATM
RESIDUAL WEIGHT FRACTION	.33289108E-01
VOLUMETRIC EFFICIENCY	.97627462E 00 PERCENT

THETA	T	P	V	H	S	U	N	Q	QTOT	W	WTOT	HB
.00	344.84	.99	.54611	-22.58769	.95518	-35.68175	.01911	.00000	.00000	.00000	.00000	.00000
10.00	345.51	1.00	.54302	-22.48808	.95518	-35.60766	.01911	.00000	.00000	.00000	.00000	.00000
20.00	347.55	1.02	.53380	-22.18514	.95518	-35.38210	.01911	.00000	.00000	.00000	.00000	.00000
30.00	351.01	1.06	.51855	-21.66938	.95518	-34.99796	.01911	.00000	.00000	.00000	.00000	.00000
40.00	356.02	1.12	.49750	-20.92281	.95518	-34.44157	.01911	.00000	.00000	.00000	.00000	.00000
50.00	362.73	1.21	.47096	-19.91986	.95518	-33.69353	.01911	.00000	.00000	.00000	.00000	.00000
60.00	371.37	1.33	.43941	-18.62534	.95518	-32.72705	.01911	.00000	.00000	.00000	.00000	.00000
70.00	382.23	1.49	.40348	-16.99272	.95518	-31.50656	.01911	.00000	.00000	.00000	.00000	.00000
80.00	395.67	1.70	.36400	-14.95164	.95518	-29.98581	.01911	.00000	.00000	.00000	.00000	.00000
90.00	412.16	2.01	.32199	-12.45483	.95518	-28.10524	.01911	.00000	.00000	.00000	.00000	.00000
100.00	432.28	2.43	.27867	-9.37447	.95518	-25.78904	.01911	.00000	.00000	.00000	.00000	.00000
110.00	456.74	3.04	.23542	-5.59927	.95518	-22.94252	.01911	.00000	.00000	.00000	.00000	.00000
120.00	486.33	3.94	.19371	-.98560	.95518	-19.45242	.01911	.00000	.00000	.00000	.00000	.00000
130.00	521.84	5.28	.15510	4.61714	.95518	-15.19795	.01911	.00000	.00000	.00000	.00000	.00000
140.00	563.67	7.30	.12105	11.31073	.95518	-10.09234	.01911	.00000	.00000	.00000	.00000	.00000
150.00	610.94	10.30	.09296	18.99131	.95518	-4.20704	.01911	.00000	.00000	.00000	.00000	.00000
160.00	659.41	14.36	.07199	26.99693	.95518	1.95803	.01911	.00000	.00000	.00000	.00000	.00000
164.00	677.09	16.13	.06580	29.94867	.95518	4.23851	.01911	.00000	.00000	.00000	.00000	.00000
166.95	688.74	17.40	.06209	31.90476	.95521	5.74418	.01911	.00000	.00000	.00000	.00000	.00000
169.07	696.63	18.30	.05987	33.23635	.95547	6.70610	.01911	.00000	.00000	.00000	.00000	.00000
171.43	705.94	19.41	.05785	34.82764	.95664	7.63212	.01912	.00000	.00000	.00000	.00000	.00000
172.60	711.73	20.13	.05704	35.83153	.95808	8.02685	.01912	.00000	.00000	.00000	.00000	.00000
174.20	720.72	21.29	.05611	37.42886	.96096	8.50117	.01914	.00000	.00000	.00000	.00000	.00000
176.53	734.82	23.22	.05515	39.91459	.96663	8.89990	.01917	.00000	.00000	.00000	.00000	.00000
177.99	746.13	24.87	.05480	42.02377	.97225	9.01625	.01920	.00000	.00000	.00000	.00000	.00000
179.06	756.12	26.41	.05465	43.98577	.97785	9.02613	.01923	.00000	.00000	.00000	.00000	.00000
179.91	765.28	27.89	.05461	45.87238	.98341	8.98449	.01927	.00000	.00000	.00000	.00000	.00000
180.96	777.84	30.03	.05465	48.59813	.99170	8.85165	.01931	.00000	.00000	.00000	.00000	.00000
181.84	789.33	32.11	.05476	51.24756	.99992	8.67122	.01936	.00000	.00000	.00000	.00000	.00000
182.51	800.05	34.15	.05488	53.87490	1.00810	8.48504	.01941	.00000	.00000	.00000	.00000	.00000
183.08	810.12	36.16	.05502	56.47580	1.01623	8.28802	.01946	.00000	.00000	.00000	.00000	.00000
183.57	819.61	38.15	.05516	59.05734	1.02431	8.08464	.01951	.00000	.00000	.00000	.00000	.00000
184.02	828.60	40.12	.05531	61.61343	1.03234	7.86884	.01956	.00000	.00000	.00000	.00000	.00000
184.47	837.06	42.05	.05548	64.11919	1.04033	7.61939	.01961	.00000	.00000	.00000	.00000	.00000
184.93	845.04	43.94	.05565	66.56620	1.04828	7.32792	.01966	.00000	.00000	.00000	.00000	.00000
185.40	852.56	45.79	.05587	68.95552	1.05619	6.99509	.01971	.00000	.00000	.00000	.00000	.00000
185.88	859.63	47.58	.05611	71.28861	1.06405	6.59611	.01976	.00000	.00000	.00000	.00000	.00000
186.21	866.66	49.42	.05629	73.63648	1.07188	6.26394	.01981	.00000	.00000	.00000	.00000	.00000
186.76	872.81	51.08	.05660	75.67922	1.07965	5.85715	.01986	.00000	.00000	.00000	.00000	.00000
187.43	878.28	52.60	.05701	77.46046	1.08736	5.43613	.01990	.00000	.00000	.00000	.00000	.00000
188.52	882.06	53.66	.05777	78.38318	1.09495	5.00118	.01995	.00000	.00000	.00000	.00000	.00000
190.17	883.26	54.01	.05911	77.96202	1.10242	.64654	.02000	.00000	.00000	.00000	.00000	.00000
191.85	883.35	54.03	.06071	76.98236	1.10983	-2.47204	.02004	.00000	.00000	.00000	.00000	.00000
193.51	882.61	53.82	.06254	75.57424	1.11722	-5.94615	.02009	.00000	.00000	.00000	.00000	.00000
195.13	881.26	53.44	.06454	73.82680	1.12458	-9.70149	.02013	.00000	.00000	.00000	.00000	.00000
196.44	880.80	53.31	.06632	72.64261	1.13201	-12.98484	.02018	.00000	.00000	.00000	.00000	.00000
197.88	879.18	52.85	.06835	70.78064	1.13940	-16.83239	.02022	.00000	.00000	.00000	.00000	.00000
199.42	876.56	52.12	.07090	68.32371	1.14675	-21.17425	.02026	.00000	.00000	.00000	.00000	.00000
201.49	870.44	50.44	.07451	63.36074	1.15393	-27.36808	.02030	.00000	.00000	.00000	.00000	.00000
203.19	866.02	49.25	.07771	60.02453	1.16124	-32.66965	.02035	.00000	.00000	.00000	.00000	.00000
204.63	862.83	48.41	.08060	57.17817	1.16864	-37.32830	.02039	.00000	.00000	.00000	.00000	.00000
206.30	857.98	47.16	.08417	53.23574	1.17598	-42.89579	.02043	.00000	.00000	.00000	.00000	.00000
208.12	852.04	45.66	.08828	48.52314	1.18329	-49.09771	.02047	.00000	.00000	.00000	.00000	.00000
210.07	845.15	43.97	.09296	43.14433	1.19058	-56.84725	.02051	.00000	.00000	.00000	.00000	.00000
212.11	837.57	42.17	.09817	37.23652	1.19788	-63.02759	.02055	.00000	.00000	.00000	.00000	.00000
214.77	826.30	39.61	.10335	28.71628	1.20504	-72.35293	.02059	.00000	.00000	.00000	.00000	.00000
219.21	804.27	35.12	.11840	13.87493	1.21206	-86.84225	.02063	.00000	.00000	.00000	.00000	.00000

225.00	2354.77	29.03	.13718	-9.21567	1.21028-105.66115	.02061	4.35938	28.90110	14.45951	41.07830	.03067
230.00	2270.55	24.79	.15485	-27.42208	1.20887-120.38681	.02060	3.29765	32.19875	11.42801	52.50631	.02729
235.00	2191.68	21.33	.17366	-44.04207	1.20756-133.76017	.02060	2.95586	35.15461	10.41750	62.92381	.02476
240.00	2118.57	18.51	.19344	-59.18444	1.20634-145.90009	.02060	2.66634	37.82095	9.47359	72.39739	.02257
245.00	2051.32	16.20	.21399	-72.94759	1.20520-156.90549	.02060	2.42177	40.24272	8.58363	80.98102	.02067
250.00	1989.72	14.30	.23513	-85.44770	1.20412-166.88152	.02060	2.21523	42.45795	7.76080	88.74182	.01903
255.00	1933.41	12.73	.25665	-96.80103	1.20309-175.92871	.02060	2.04035	44.49830	7.00684	95.74866	.01761
260.00	1881.95	11.42	.27837	-107.12217	1.20211-184.14319	.02059	1.89164	46.38993	6.32284	102.07150	.01638
265.00	1834.95	10.33	.30012	-116.51292	1.20117-191.60929	.02059	1.76448	48.15441	5.70162	107.77313	.01531
270.00	1791.97	9.41	.32170	-125.06549	1.20026-198.40270	.02059	1.65513	49.80954	5.13829	112.91142	.01437
275.00	1752.67	8.64	.34295	-132.86139	1.19939-204.58994	.02059	1.56056	51.37010	4.62668	117.53809	.01356
280.00	1716.71	7.98	.36372	-139.97358	1.19855-210.23030	.02059	1.47834	52.84844	4.16202	121.70011	.01284
285.00	1683.80	7.41	.38385	-146.46600	1.19772-215.37558	.02059	1.40647	54.25490	3.73882	125.43892	.01221
290.00	1653.67	6.93	.40322	-152.39512	1.19693-220.07146	.02059	1.34335	55.59825	3.35253	128.79145	.01166
295.00	1626.08	6.52	.42170	-157.81086	1.19614-224.35823	.02059	1.28768	56.88593	2.99909	131.79054	.01117
300.00	1600.71	6.16	.43918	-162.78120	1.19538-228.29032	.02059	1.23838	58.12431	2.69371	134.48425	.01074
305.00	1577.51	5.85	.45556	-167.31917	1.19462-231.87854	.02059	1.19436	59.31866	2.39386	136.87811	.01036
310.00	1556.30	5.59	.47076	-171.45878	1.19388-235.15024	.02059	1.15513	60.47379	2.11658	138.99469	.01003
315.00	1536.95	5.36	.48471	-175.22989	1.19314-238.12943	.02059	1.12009	61.59388	1.85910	140.85379	.00973
320.00	1519.33	5.16	.49734	-178.65873	1.19242-240.83716	.02059	1.08874	62.68261	1.61899	142.47278	.00947
325.00	1503.33	4.99	.50859	-181.76815	1.19172-243.29176	.02059	1.06067	63.74328	1.39394	143.86671	.00924
330.00	1488.61	4.85	.51843	-184.57576	1.19104-245.49704	.02059	1.03552	64.77880	1.16976	145.03647	.00905
335.00	1475.48	4.73	.52681	-187.09629	1.19036-247.48017	.02059	1.01262	65.79142	.97050	146.00697	.00887
340.00	1463.72	4.63	.53371	-189.35049	1.18969-249.25326	.02059	.99237	66.78379	.78073	146.78770	.00872
345.00	1453.28	4.56	.53911	-191.35099	1.18902-250.82639	.02059	.97432	67.75811	.59881	147.38651	.00860
350.00	1444.10	4.49	.54298	-193.10823	1.18836-252.20789	.02059	.95832	68.71642	.42318	147.80969	.00849
355.00	1436.14	4.45	.54531	-194.63063	1.18770-253.40450	.02059	.94422	69.66065	.25240	148.06208	.00841
360.00	1429.37	4.42	.54611	-195.92474	1.18705-254.42149	.02059	.93192	70.59257	.08506	148.14715	.00834
365.00	1036.55	1.03	1.70051	-268.95662	1.18705-311.37718	.02059	.00000	70.59257	.00000	148.14715	.00000
540.00	1000.00	1.03	.05461	-9.17099	.03930-10.53334	.00069	.00000	70.59257	.00000	148.14715	.00000

THETA	T	P	V	H	S	U	N	FLAME SPEED,	TEMP.,	AND EXTENT	MASS-FR.	BURN.-VOL.
166.95	2541.13	17.40	.00002	.00381	.00013	-.00663	.00000	677.4219	2541.13266	.18781	.00010	.00040
169.07	2541.51	18.30	.00024	.03787	.00125	-.06653	.00002	692.35579	2538.89896	.39187	.00100	.00394
171.43	2550.06	19.41	.00111	.19925	.00623	-.32458	.00010	715.48909	2548.37943	.65387	.00500	.01926
172.60	2554.17	20.13	.00215	.40749	.01246	-.64186	.00021	727.48067	2546.29190	.80122	.01000	.03774
174.20	2565.35	21.29	.00409	.86935	.02489	-1.23883	.00041	751.76242	2558.72079	1.04280	.02000	.07288
176.53	2572.67	23.22	.00752	1.79698	.04966	-2.43133	.00083	790.76913	2578.12265	1.35169	.04000	.13633
177.99	2585.69	24.87	.01058	2.88413	.07439	-3.49138	.00124	819.17786	2581.62408	1.56528	.06000	.19315
179.06	2598.75	26.41	.01336	4.10342	.09908	-4.44179	.00165	845.77429	2586.87708	1.70606	.08000	.24443
179.91	2611.26	27.89	.01589	5.44104	.12375	-5.29383	.00207	870.95643	2592.35233	1.83262	.10000	.29101
180.96	2628.34	30.03	.01932	7.62968	.16069	-6.42052	.00269	908.71444	2607.40726	1.97243	.13000	.35349
181.84	2643.99	32.11	.02238	10.02125	.19758	-7.37835	.00331	941.95177	2614.08182	2.08508	.16000	.40867
182.51	2658.76	34.15	.02513	12.61288	.23441	-8.16959	.00393	974.1654	2620.85892	2.17251	.19000	.45786
183.08	2672.52	36.16	.02762	15.37513	.27119	-8.81863	.00456	1005.25099	2627.35922	2.24560	.22000	.50207
183.57	2685.38	38.15	.02990	18.29329	.30793	-9.33773	.00518	1035.37379	2633.62735	2.28734	.25000	.54207
184.02	2697.35	40.12	.03200	21.34726	.34462	-9.74371	.00580	1064.55179	2639.61224	2.34508	.28000	.57849
184.47	2708.34	42.05	.03394	24.50578	.38126	-10.06258	.00642	1092.62779	2645.21103	2.40259	.31000	.61183
184.93	2718.37	43.94	.03576	27.74849	.41786	-10.31192	.00705	1119.66524	2650.52722	2.46055	.34000	.64250
185.40	2727.58	45.79	.03748	31.06672	.45443	-10.49855	.00767	1145.64869	2655.54974	2.51891	.37000	.67083
185.88	2735.85	47.58	.03912	34.42420	.49095	-10.65320	.00829	1170.52689	2660.26102	2.57856	.40000	.69712
186.21	2744.30	49.42	.04062	37.93510	.52743	-10.67959	.00891	1195.86458	2665.58032	2.62837	.43000	.72158
186.76	2750.36	51.08	.04213	41.26746	.56385	-10.85782	.00954	1218.35568	2669.13165	2.69646	.46000	.74441
187.43	2754.77	52.60	.04366	44.46488	.60022	-11.15100	.01016	1238.50146	2672.33578	2.78304	.49000	.76580
188.52	2754.27	53.66	.04540	47.00599	.63647	-11.99919	.01078	1251.89896	2672.99021	3.05689	.52000	.78587
190.17	2746.75	54.01	.04757	48.45000	.67259	-13.77300	.01140	1255.70917	2669.75861	3.28398	.55000	.80479
191.85	2737.07	54.03	.04995	49.43100	.70866	-15.93564	.01202	1255.64316	2668.16519	3.53648	.58000	.82269
193.51	2726.06	53.82	.05251	50.04192	.74470	-18.40787	.01264	1252.55498	2666.35791	3.78516	.61000	.83966
195.13	2714.21	53.44	.05523	50.34526	.78072	-21.13637	.01325	1247.21426	2664.38669	4.02183	.64000	.85577
196.44	2706.03	53.31	.05777	51.14469	.81680	-23.44572	.01387	1245.81732	2665.37772	4.20755	.67000	.87110
197.88	2695.19	52.85	.06062	51.32358	.85284	-26.27425	.01449	1239.26904	2662.36621	4.38379	.70000	.88568
199.42	2682.25	52.12	.06378	50.93790	.88885	-29.57320	.01510	1229.40932	2659.01791	4.56171	.73000	.89958
201.49	2659.69	50.44	.06801	48.44779	.92469	-34.62855	.01572	1205.23198	2648.91498	4.78342	.76000	.91283
203.19	2643.06	49.25	.07192	46.89510	.96065	-38.89337	.01633	1189.78374	2648.10764	4.97720	.79000	.92549
204.63	2630.69	48.41	.07557	46.02615	.99671	-42.58297	.01695	1178.56776	2647.42303	5.12571	.82000	.93759
206.30	2614.07	47.16	.07989	44.07098	1.03270	-47.17369	.01757	1160.50578	2641.16153	5.27697	.85000	.94916
208.12	2594.79	45.66	.08477	41.31735	1.06867	-52.42111	.01818	1138.92339	2634.84943	5.43334	.88000	.96022
210.07	2573.36	43.97	.09025	37.84918	1.10462	-58.25411	.01879	1114.17792	2627.88657	5.59139	.91000	.97082
212.11	2550.41	42.17	.09630	33.78652	1.14057	-64.56934	.01941	1087.72800	2620.84341	5.74637	.94000	.98097
214.77	2517.07	39.61	.10437	27.05063	1.17638	-73.07731	.02002	1048.18750	2606.30457	5.92646	.97000	.99069
219.21	2457.35	35.12	.11840	13.87493	1.21206	-86.84225	.02063	973.76212	2573.25613	6.19125	1.00000	1.00000

*** ALL INPUT DATA HAVE BEEN PROCESSED:
AT LOC 45270

A MODEL SIMULATING THE INTERNAL-COMBUSTION ENGINE

ENGINE	FUEL	OPERATING CONDITIONS
FOUR STROKE AUTOMOTIVE TYPE	OCTANE C8H18	AIR-FUEL RATIO .92
BORE 3.94 IN.	ACTIVATION ENERGY 39000.00 CAL./MOLE	PISTON TEMP 500.00 K
STROKE 2.46 IN.	ENTHALPY OF COMBUSTION 1222770.00 CAL/MOLE	WALL TEMP 420.00 K
ROD 7.06 IN.		MAN. TEMP 320.00 K
CLEAR. VOL. 3.33 CU.IN.		MAN. PR. .99 ATM
SWEPT VOL. 30.00 CU.IN.		EXH. PR. 1.03 ATM
COMPRESSION RATIO 10.00 TO ONE		SPARK ADV. 16.00
		SPEED 2800.00 RPM

COMBUSTION OCCURS IN 39 INCREMENTS. ERRORS ARE LESS THAN .00100 PERCENT FLAME SPEED FACTOR IS .135000000E 06
 HEAT TRANSFER FACTOR IS .00000 NORMAL RUN

SUMMARY OF RESULTS

PUMPING WORK	- .47611947E 00 CAL
NET CYCLE WORK	.16383728E 03 CAL
NET HEAT TRANSFERRED	.00000000E 00 CAL
INDICATED THERMAL EFFICIENCY	.40573207E 02 PERCENT
MEAN EFFECTIVE PRESSURE	.13764381E 02 ATM
RESIDUAL WEIGHT FRACTION	0.03329
VOLUMETRIC EFFICIENCY	.97627462E 00 PERCENT

THETA	T	P	V	H	S	U	N	Q	GTOT	W	WTOT	HE
.00	344.84	.99	.54611	-22.58769	.95518	-35.68175	.01911	.00000	.00000	.00000	.00000	.00000
10.00	345.51	1.00	.54302	-22.48808	.95518	-35.68210	.01911	.00000	.00000	.00000	.00000	.00000
20.00	347.59	1.02	.53360	-22.18514	.95518	-35.38210	.01911	.00000	.00000	.00000	.00000	.00000
30.00	351.01	1.06	.51855	-21.66938	.95518	-34.99726	.01911	.00000	.00000	.00000	.00000	.00000
40.00	356.02	1.12	.49750	-20.92281	.95518	-34.44157	.01911	.00000	.00000	.00000	.00000	.00000
50.00	362.73	1.21	.47026	-19.91986	.95518	-33.69353	.01911	.00000	.00000	.00000	.00000	.00000
60.00	371.37	1.33	.43941	-18.62534	.95518	-32.72705	.01911	.00000	.00000	.00000	.00000	.00000
70.00	382.23	1.49	.40348	-16.99272	.95518	-31.50656	.01911	.00000	.00000	.00000	.00000	.00000
80.00	395.67	1.70	.36300	-14.95164	.95518	-29.98581	.01911	.00000	.00000	.00000	.00000	.00000
90.00	412.16	2.01	.32199	-12.45483	.95518	-28.10524	.01911	.00000	.00000	.00000	.00000	.00000
100.00	432.28	2.43	.27867	-9.37447	.95518	-25.78904	.01911	.00000	.00000	.00000	.00000	.00000
110.00	456.74	3.04	.23542	-5.59927	.95518	-23.94252	.01911	.00000	.00000	.00000	.00000	.00000
120.00	486.33	3.94	.19371	-.98560	.95518	-19.45242	.01911	.00000	.00000	.00000	.00000	.00000
130.00	521.84	5.28	.15510	4.61714	.95518	-15.19795	.01911	.00000	.00000	.00000	.00000	.00000
140.00	563.67	7.30	.12108	11.31073	.95518	-10.05294	.01911	.00000	.00000	.00000	.00000	.00000
150.00	610.94	10.30	.09296	18.99131	.95518	-4.20704	.01911	.00000	.00000	.00000	.00000	.00000
160.00	659.41	14.36	.07199	26.99693	.95518	1.95803	.01911	.00000	.00000	.00000	.00000	.00000
170.00	677.09	16.13	.06580	29.94867	.95518	4.23851	.01911	.00000	.00000	.00000	.00000	.00000
180.00	688.78	17.40	.06208	31.91122	.95521	5.74918	.01911	.00000	.00000	.00000	.00000	.00000
190.00	696.69	18.30	.05985	33.24601	.95547	6.71358	.01911	.00000	.00000	.00000	.00000	.00000
200.00	706.01	19.42	.05784	34.83917	.95684	7.64106	.01912	.00000	.00000	.00000	.00000	.00000
210.00	711.80	20.14	.05702	35.84326	.95608	8.03595	.01912	.00000	.00000	.00000	.00000	.00000
220.00	720.78	21.30	.05609	37.43998	.95096	8.50981	.01914	.00000	.00000	.00000	.00000	.00000
230.00	735.01	23.25	.05514	40.08403	.96668	9.04059	.01917	.00000	.00000	.00000	.00000	.00000
240.00	746.39	24.97	.05439	42.30435	.97234	9.72996	.01920	.00000	.00000	.00000	.00000	.00000
250.00	756.43	26.46	.05465	44.36051	.97797	9.33911	.01923	.00000	.00000	.00000	.00000	.00000
260.00	765.62	27.95	.05461	46.33032	.98356	9.36778	.01927	.00000	.00000	.00000	.00000	.00000
270.00	778.23	30.10	.05465	49.18056	.99188	9.34016	.01932	.00000	.00000	.00000	.00000	.00000
280.00	793.76	32.19	.05479	51.94065	1.00015	9.25343	.01935	.00000	.00000	.00000	.00000	.00000
290.00	800.51	34.24	.05489	54.65872	1.00835	9.14455	.01941	.00000	.00000	.00000	.00000	.00000
300.00	810.59	36.26	.05503	57.34387	1.01650	9.01935	.01946	.00000	.00000	.00000	.00000	.00000
310.00	820.10	38.26	.05517	60.00130	1.02461	8.88097	.01951	.00000	.00000	.00000	.00000	.00000
320.00	829.09	40.23	.05532	62.63300	1.03267	8.72987	.01956	.00000	.00000	.00000	.00000	.00000
330.00	837.57	42.17	.05549	65.21992	1.04068	8.54962	.01961	.00000	.00000	.00000	.00000	.00000
340.00	845.57	44.07	.05568	67.77639	1.04856	8.33282	.01966	.00000	.00000	.00000	.00000	.00000
350.00	853.12	45.93	.05589	70.29279	1.05632	8.12717	.01971	.00000	.00000	.00000	.00000	.00000
360.00	860.26	47.74	.05612	72.76210	1.06452	7.86950	.01976	.00000	.00000	.00000	.00000	.00000
370.00	867.33	49.60	.05629	75.29512	1.07240	7.66963	.01981	.00000	.00000	.00000	.00000	.00000
380.00	873.54	51.28	.05661	77.57456	1.08024	7.26423	.01986	.00000	.00000	.00000	.00000	.00000
390.00	879.10	52.83	.05703	79.68719	1.08805	6.72494	.01991	.00000	.00000	.00000	.00000	.00000
400.00	882.94	53.92	.05762	81.77369	1.09584	5.66309	.01996	.00000	.00000	.00000	.00000	.00000
410.00	884.51	54.37	.05816	81.78600	1.10362	3.88925	.02001	.00000	.00000	.00000	.00000	.00000
420.00	885.01	54.51	.06076	81.96679	1.11139	1.75154	.02005	.00000	.00000	.00000	.00000	.00000
430.00	884.69	54.42	.06257	81.79891	1.11916	-.67587	.02010	.00000	.00000	.00000	.00000	.00000
440.00	884.14	54.26	.06446	81.52277	1.12692	-3.19319	.02015	.00000	.00000	.00000	.00000	.00000
450.00	884.09	54.25	.06621	81.46980	1.13470	-5.52423	.02019	.00000	.00000	.00000	.00000	.00000
460.00	882.92	53.91	.06830	80.88959	1.14247	-8.29499	.02024	.00000	.00000	.00000	.00000	.00000
470.00	880.76	53.30	.07073	79.81730	1.15025	-11.47630	.02029	.00000	.00000	.00000	.00000	.00000
480.00	875.33	51.77	.07427	77.08885	1.15803	-16.04000	.02033	.00000	.00000	.00000	.00000	.00000
490.00	871.64	50.76	.07735	75.18480	1.16584	-19.91560	.02037	.00000	.00000	.00000	.00000	.00000
500.00	869.01	50.05	.08015	73.79358	1.17367	-23.36426	.02042	.00000	.00000	.00000	.00000	.00000
510.00	864.82	48.93	.08359	71.53066	1.18151	-27.53979	.02046	.00000	.00000	.00000	.00000	.00000
520.00	859.64	47.98	.08734	68.67415	1.18937	-32.21182	.02050	.00000	.00000	.00000	.00000	.00000
530.00	853.60	46.05	.09201	65.24858	1.19725	-37.35772	.02055	.00000	.00000	.00000	.00000	.00000
540.00	846.87	44.39	.09698	61.35702	1.20515	-42.89187	.02059	.00000	.00000	.00000	.00000	.00000
550.00	836.81	41.99	.10378	55.42771	1.21308	-50.12120	.02063	.00000	.00000	.00000	.00000	.00000
560.00	817.02	37.75	.11584	44.04836	1.22108	-61.87004	.02066	.00000	.00000	.00000	.00000	.00000

220.00	2558.42	35.88	1.2084	38.70322	1.22108	-66.29910	.02065	.00000	.00000	4.42207	30.61735	.03258
225.00	2492.68	30.77	1.3718	22.77491	1.22109	-79.44677	.02064	.00000	.00000	13.14766	43.76502	.03162
230.00	2428.97	26.54	1.5485	7.89184	1.22108	-91.65689	.02062	.00000	.00000	12.21012	55.97514	.02891
235.00	2367.76	23.06	1.7366	-5.95078	1.22109	-102.94823	.02062	.00000	.00000	11.29135	67.26649	.02651
240.00	2309.85	20.19	1.9344	-18.69259	1.22108	-113.28922	.02061	.00000	.00000	10.34099	77.60747	.02439
245.00	2259.60	17.82	2.1399	-30.37516	1.22107	-122.73083	.02060	.00000	.00000	9.44161	87.04908	.02254
250.00	2205.32	15.85	2.3513	-41.02566	1.22107	-131.30954	.02060	.00000	.00000	8.57870	95.62779	.02093
255.00	2158.81	14.22	2.5635	-50.75101	1.22108	-139.12247	.02060	.00000	.00000	7.81294	103.44073	.01952
260.00	2116.03	12.85	2.7837	-59.61102	1.22109	-146.22553	.02060	.00000	.00000	7.10305	110.54378	.01829
265.00	2076.83	11.69	3.0012	-67.66924	1.22109	-152.67520	.02060	.00000	.00000	6.44967	116.99345	.01721
270.00	2041.02	10.72	3.2170	-74.98772	1.22109	-158.52501	.02060	.00000	.00000	5.84981	122.84326	.01627
275.00	2008.38	9.90	3.4295	-81.62607	1.22109	-163.82554	.02060	.00000	.00000	5.30053	128.14379	.01544
280.00	1978.70	9.19	3.6372	-87.63829	1.22109	-168.62182	.02060	.00000	.00000	4.79628	132.94007	.01472
285.00	1951.78	8.59	3.8385	-93.07412	1.22109	-172.95498	.02060	.00000	.00000	4.33317	137.27324	.01408
290.00	1927.43	8.08	4.0322	-97.97862	1.22110	-176.86208	.02060	.00000	.00000	3.90709	141.18033	.01352
295.00	1905.47	7.64	4.2170	-102.39223	1.22110	-180.37617	.02060	.00000	.00000	3.51409	144.69442	.01302
300.00	1885.73	7.26	4.3918	-106.35117	1.22110	-183.52672	.02060	.00000	.00000	3.15055	147.84497	.01259
305.00	1867.92	6.93	4.5556	-109.91497	1.22110	-186.36162	.02059	.00000	.00000	2.83490	150.67987	.01221
310.00	1852.09	6.65	4.7076	-113.08050	1.22108	-188.87877	.02059	.00000	.00000	2.51715	153.19702	.01188
315.00	1838.09	6.41	4.8471	-115.87372	1.22107	-191.09916	.02059	.00000	.00000	2.22038	155.41741	.01158
320.00	1825.84	6.20	4.9734	-118.31686	1.22107	-193.04065	.02059	.00000	.00000	1.94149	157.35890	.01133
325.00	1815.23	6.03	5.0859	-120.43025	1.22106	-194.71977	.02059	.00000	.00000	1.67912	159.03802	.01111
330.00	1806.19	5.89	5.1843	-122.23008	1.22105	-196.14946	.02059	.00000	.00000	1.42969	160.46771	.01092
335.00	1798.65	5.77	5.2681	-123.73052	1.22105	-197.34112	.02059	.00000	.00000	1.19166	161.65937	.01076
340.00	1792.55	5.68	5.3371	-124.94327	1.22104	-198.30416	.02059	.00000	.00000	.96304	162.62241	.01063
345.00	1787.89	5.60	5.3911	-125.87772	1.22104	-199.04612	.02059	.00000	.00000	.74196	163.36437	.01053
350.00	1784.51	5.55	5.4298	-126.54113	1.22104	-199.57282	.02059	.00000	.00000	.52670	163.89107	.01045
355.00	1782.50	5.52	5.4531	-126.93859	1.22103	-199.88836	.02059	.00000	.00000	.31554	164.20661	.01039
360.00	1781.83	5.51	5.4611	-127.07311	1.22103	-199.99516	.02059	.00000	.00000	.10680	164.31341	.01036
360.00	1247.85	1.03	2.04716	-230.23209	1.22103	-281.30019	.02059	.00000	.00000	.00000	164.31341	.00000
540.00	1000.00	1.03	.05461	-9.17099	.03930	-10.53334	.00069	.00000	.00000	.00000	164.31341	.00000

THETA	I	P	V	H	S	U	N	FLAME SPEED	TEMP.	AND EXTENT	MASS-FR.	BURN.-VOL.
166.96	2541.23	17.40	.00002	.00381	.00013	-.00663	.00000	675.03462	2541.23251	.18781	.00010	.00040
169.09	2541.59	18.30	.00024	.03789	.00125	-.06652	.00002	689.94530	2538.97421	.39184	.00100	.00394
171.46	2550.13	19.42	.00111	.19934	.00523	-.32450	.00010	713.02339	2548.45255	.65380	.00500	.01926
172.63	2554.24	20.14	.00215	.40765	.01246	-.64173	.00021	724.96620	2546.34534	.80120	.01000	.03774
174.23	2565.39	21.30	.00409	.86957	.02489	-1.23865	.00041	749.14967	2558.76233	1.04276	.02000	.07287
176.57	2585.17	23.25	.00755	1.93536	.04971	-2.31479	.00083	788.40847	2578.47577	1.35166	.04000	.13691
178.03	2599.95	24.91	.01063	3.12240	.07447	-3.29059	.00124	816.87914	2581.92993	1.56692	.06000	.19401
179.10	2613.29	26.46	.01341	4.42940	.09920	-4.16678	.00166	843.52698	2587.18536	1.70809	.08000	.24545
179.95	2625.62	27.95	.01595	5.84612	.12390	-4.95174	.00207	868.51958	2592.51596	1.83490	.10000	.29213
181.00	2642.48	30.10	.01938	8.15265	.16088	-5.97840	.00263	906.48457	2607.75507	1.97448	.13000	.35469
181.88	2657.73	32.19	.02245	10.65118	.19780	-6.84548	.00331	939.80029	2614.47516	2.08684	.16000	.40968
182.54	2671.89	34.24	.02520	13.33242	.23466	-7.56034	.00393	971.98961	2621.24246	2.17426	.19000	.45904
183.10	2685.10	36.26	.02769	16.17847	.27147	-8.13794	.00458	1003.05559	2627.75287	2.24752	.22000	.50319
183.60	2697.44	38.26	.02996	19.17372	.30822	-8.59114	.00518	1033.12535	2634.01050	2.28756	.25000	.54313
184.04	2709.00	40.23	.03206	22.30414	.34494	-8.93179	.00580	1062.27342	2640.01144	2.34663	.28000	.57948
184.50	2719.71	42.17	.03400	25.54450	.38161	-9.18097	.00643	1090.33157	2645.62628	2.40418	.31000	.61276
184.96	2729.80	44.07	.03582	28.89872	.41825	-9.33508	.00705	1117.35495	2650.95486	2.46217	.34000	.64340
185.43	2739.19	45.93	.03754	32.34172	.45485	-9.41535	.00767	1143.40410	2656.03308	2.52067	.37000	.67171
185.90	2747.92	47.74	.03917	35.86118	.49141	-9.43204	.00823	1168.47995	2660.84982	2.58041	.40000	.69797
186.23	2756.70	49.60	.04067	39.52596	.52795	-9.32716	.00892	1193.90981	2666.16632	2.63092	.43000	.72241
186.78	2763.63	51.28	.04219	43.09260	.56444	-9.30593	.00964	1216.26248	2669.60800	2.69876	.46000	.74524
187.46	2769.44	52.83	.04372	46.61716	.60091	-9.32073	.01016	1236.99683	2673.08319	2.78677	.49000	.76666
188.59	2771.72	53.92	.04550	49.72116	.63736	-9.69055	.01073	1250.69514	2673.40039	3.06524	.52000	.78666
190.23	2769.52	54.37	.04768	52.17376	.67379	-10.60914	.01141	1255.91722	2671.26492	3.29522	.55000	.80597
191.89	2765.46	54.51	.05007	54.29125	.71021	-11.80975	.01203	1257.45834	2670.06979	3.54926	.58000	.82404
193.54	2760.09	54.42	.05263	56.12216	.74664	-13.25121	.01265	1255.98044	2668.61090	3.79859	.61000	.84114
195.07	2754.65	54.26	.05527	57.85642	.78306	-14.77345	.01327	1253.81386	2667.83652	4.03481	.64000	.85733
196.36	2750.69	54.25	.05778	59.77872	.81949	-16.13706	.01389	1253.80103	2668.54456	4.21767	.67000	.87265
197.79	2744.62	53.91	.06060	61.23289	.85591	-17.89387	.01451	1248.97507	2665.96671	4.39401	.70000	.88722
199.32	2736.71	53.30	.06373	62.22390	.89235	-20.03378	.01513	1240.76724	2662.97079	4.57284	.73000	.90109
201.36	2721.91	51.77	.06791	61.68754	.92879	-23.46426	.01574	1219.42540	2653.93634	4.79572	.76000	.91434
203.01	2711.40	50.76	.07170	61.84615	.96526	-26.30374	.01636	1206.23943	2653.58295	4.98809	.79000	.92691
204.41	2703.59	50.05	.07525	62.44442	1.00174	-28.77381	.01698	1196.90425	2653.08716	5.13529	.82000	.93886
206.04	2692.49	48.93	.07944	62.18438	1.03824	-31.96027	.01760	1181.09462	2647.49860	5.28569	.85000	.95028
207.80	2679.43	47.58	.08414	61.30721	1.07476	-36.66173	.01821	1162.03128	2641.95386	5.44112	.88000	.96117
209.68	2664.56	46.05	.08939	59.81944	1.11128	-39.86973	.01883	1140.19296	2635.92868	5.59795	.91000	.97157
211.65	2648.38	44.39	.09518	57.80876	1.14784	-44.51070	.01944	1116.10751	2629.48132	5.75190	.94000	.98149
214.20	2624.65	41.99	.10284	53.70671	1.18442	-50.88895	.02005	1080.19279	2616.50229	5.93031	.97000	.99097
218.37	2579.93	37.75	.11584	44.04836	1.22108	-61.87003	.02068	1012.26093	2587.10577	6.18942	1.00000	1.00000

*** ALL INPUT DATA HAVE BEEN PROCESSED.
 AT LOC 45325

A MODEL SIMULATING THE INTERNAL-COMBUSTION ENGINE

ENGINE	FUEL	OPERATING CONDITIONS
FOUR STROKE AUTOMOTIVE TYPE	OCTANE C8H18	AIR-FUEL RATIO .92
BORE 3.94 IN.	ACTIVATION ENERGY 39000.00 CAL./MOLE	PISTON TEMP 500.00 K
STROKE 2.46 IN.	ENTHALPY OF COMBUSTION 1222770.00 CAL/MOLE	WALL TEMP 420.00 K
ROD 7.06 IN.		MAN. TEMP 320.00 K
CLEAR. VOL. 3.33 CU. IN.		MAN. PR. .99 ATM
SWEPT VOL. 30.00 CU. IN.		EXH. PR. 1.03 ATM
COMPRESSION RATIO 10.00 TO ONE		SPARK ADV. 16.00
		SPEED 2800.00 RPM

COMBUSTION OCCURS IN 39 INCREMENTS. ERRORS ARE LESS THAN .00100 PERCENT. FLAME SPEED FACTOR IS .13500000E 06
 HEAT TRANSFER FACTOR IS 2.000000 NORMAL RUN

SUMMARY OF RESULTS

PUMPING WORK	.47611947E 00 CAL
NET CYCLE WORK	.13428628E 03 CAL
NET HEAT TRANSFERED	.11785904E 03 CAL
INDICATED THERMAL EFFICIENCY	.33255097E 02 PERCENT
MEAN EFFECTIVE PRESSURE	.11281727E 02 ATM
RESIDUAL WEIGHT FRACTION	.33289107E-01
VOLUMETRIC EFFICIENCY	.97627462E 00 PERCENT

THEIR	T	P	V	H	S	U	N	Q	QTOT	M	WTOT	HE
.00	344.84	.99	54611	-22.58769	.95518	-35.68175	.01911	.00000	.00000	.00000	.00000	.00000
10.00	345.51	1.00	54302	-22.48808	.95518	-35.60766	.01911	.00000	.00000	.00000	.00000	.00000
20.00	347.55	1.02	53380	-22.18514	.95518	-35.38210	.01911	.00000	.00000	.00000	.00000	.00000
30.00	351.01	1.06	51855	-21.66238	.95518	-34.99796	.01911	.00000	.00000	.00000	.00000	.00000
40.00	356.02	1.12	49750	-20.92281	.95518	-34.44157	.01911	.00000	.00000	.00000	.00000	.00000
50.00	362.73	1.21	47036	-19.31986	.95518	-33.69353	.01911	.00000	.00000	.00000	.00000	.00000
60.00	371.37	1.33	43941	-18.62534	.95518	-32.72705	.01911	.00000	.00000	.00000	.00000	.00000
70.00	382.23	1.49	40348	-16.92272	.95518	-31.50656	.01911	.00000	.00000	.00000	.00000	.00000
80.00	395.67	1.70	36400	-14.96164	.95518	-29.98581	.01911	.00000	.00000	.00000	.00000	.00000
90.00	412.16	2.01	32199	-12.45483	.95518	-28.10524	.01911	.00000	.00000	.00000	.00000	.00000
100.00	432.28	2.43	27867	-9.37447	.95518	-25.78904	.01911	.00000	.00000	.00000	.00000	.00000
110.00	456.74	3.04	23542	-5.45927	.95518	-22.32452	.01911	.00000	.00000	.00000	.00000	.00000
120.00	486.33	3.94	19371	-1.98560	.95518	-19.45242	.01911	.00000	.00000	.00000	.00000	.00000
130.00	521.84	5.28	15510	4.61714	.95518	-15.19795	.01911	.00000	.00000	.00000	.00000	.00000
140.00	563.67	7.30	12106	11.31073	.95518	-10.09294	.01911	.00000	.00000	.00000	.00000	.00000
150.00	610.94	10.30	92296	18.99131	.95518	-4.20704	.01911	.00000	.00000	.00000	.00000	.00000
160.00	659.41	14.36	67199	26.99693	.95518	1.95803	.01911	.00000	.00000	.00000	.00000	.00000
164.00	677.09	16.13	66580	29.94867	.95518	4.23851	.01911	.00000	.00000	.00000	.00000	.00000
166.96	688.78	17.40	66208	31.91122	.95521	5.74918	.01911	.00000	.00000	.00000	.00000	.00000
169.09	696.69	18.30	65985	33.24601	.95547	6.71358	.01911	.00000	.00000	.00000	.00000	.00000
171.46	706.01	19.42	65784	34.83917	.95664	7.64106	.01912	.00000	.00000	.00000	.00000	.00000
172.63	711.80	20.14	65702	35.84326	.95808	8.03595	.01912	.00000	.00000	.00000	.00000	.00000
174.23	720.78	21.30	65609	37.43998	.96096	8.50981	.01914	.00000	.00000	.00000	.00000	.00000
176.58	734.73	23.21	65514	39.75951	.96657	8.77050	.01917	.00000	.00000	.00000	.00000	.00000
178.04	745.92	24.84	65479	41.74612	.97216	8.78479	.01920	.00000	.00000	.00000	.00000	.00000
179.13	755.84	26.37	65465	43.60594	.97772	8.79872	.01923	.00000	.00000	.00000	.00000	.00000
179.98	764.92	27.83	65461	45.40099	.98326	8.79080	.01927	.00000	.00000	.00000	.00000	.00000
181.05	777.40	29.95	65466	48.00251	.99151	8.35354	.01931	.00000	.00000	.00000	.00000	.00000
181.94	788.80	32.01	65477	50.52974	.99970	8.07009	.01936	.00000	.00000	.00000	.00000	.00000
182.62	799.47	34.03	65491	53.05557	1.00785	7.79792	.01941	.00000	.00000	.00000	.00000	.00000
183.18	809.49	36.03	65505	55.56479	1.01595	7.52293	.01946	.00000	.00000	.00000	.00000	.00000
183.68	818.95	38.01	65520	58.06094	1.02401	7.24686	.01951	.00000	.00000	.00000	.00000	.00000
184.11	827.93	39.97	65534	60.59308	1.03202	6.97609	.01956	.00000	.00000	.00000	.00000	.00000
184.56	836.36	41.89	65551	62.97158	1.03999	6.65207	.01961	.00000	.00000	.00000	.00000	.00000
185.02	844.30	43.76	65571	65.31770	1.04791	6.27479	.01965	.00000	.00000	.00000	.00000	.00000
185.50	851.77	45.59	65592	67.57427	1.05578	5.82913	.01970	.00000	.00000	.00000	.00000	.00000
185.99	858.76	47.36	65617	69.70860	1.06360	5.28704	.01975	.00000	.00000	.00000	.00000	.00000
186.32	865.74	49.18	65635	71.95492	1.07139	4.83284	.01980	.00000	.00000	.00000	.00000	.00000
186.88	871.79	50.80	65667	73.74064	1.07908	4.01810	.01985	.00000	.00000	.00000	.00000	.00000
187.53	877.10	52.27	65709	75.15102	1.08669	2.89304	.01990	.00000	.00000	.00000	.00000	.00000
188.65	880.59	53.25	65786	75.43065	1.09409	.80481	.01995	.00000	.00000	.00000	.00000	.00000
190.32	881.28	53.44	65924	73.89112	1.10122	-2.79194	.01999	.00000	.00000	.00000	.00000	.00000
192.02	880.83	53.32	66089	71.71039	1.10828	-6.91954	.02003	.00000	.00000	.00000	.00000	.00000
193.71	879.51	52.94	66277	69.01791	1.11529	-11.46934	.02008	.00000	.00000	.00000	.00000	.00000
195.35	877.57	52.40	66483	65.94116	1.12226	-16.33493	.02012	.00000	.00000	.00000	.00000	.00000
196.69	876.60	52.13	66668	63.60964	1.12935	-20.55233	.02016	.00000	.00000	.00000	.00000	.00000
198.17	874.37	51.51	66890	60.42368	1.13635	-25.52248	.02020	.00000	.00000	.00000	.00000	.00000
199.68	871.51	50.73	67133	56.83023	1.14332	-30.80437	.02025	.00000	.00000	.00000	.00000	.00000
201.79	864.54	48.86	67505	50.26841	1.14993	-38.54107	.02028	.00000	.00000	.00000	.00000	.00000
203.55	859.27	47.49	67842	44.96883	1.15675	-45.22041	.02033	.00000	.00000	.00000	.00000	.00000
205.04	855.40	46.50	68146	40.70928	1.16374	-51.03528	.02037	.00000	.00000	.00000	.00000	.00000
206.77	849.75	45.09	68522	35.13890	1.17060	-57.92459	.02041	.00000	.00000	.00000	.00000	.00000
208.67	842.86	43.42	68959	28.64188	1.17738	-65.56756	.02045	.00000	.00000	.00000	.00000	.00000
210.71	834.97	41.57	69457	21.38747	1.18413	-73.82078	.02049	.00000	.00000	.00000	.00000	.00000
212.85	826.37	39.63	70090	13.57090	1.19085	-82.52167	.02053	.00000	.00000	.00000	.00000	.00000
215.66	813.63	36.89	70787	2.57031	1.19726	-93.80233	.02057	.00000	.00000	.00000	.00000	.00000
220.48	788.83	32.14	72235	-15.70533	1.20333	-110.94678	.02061	.00000	.00000	.00000	.00000	.00000

225.00	2229.48	27.47	.13718	-36.21605	1.20074-127.48591	.02060	5.96779	53.63952	10.57133	38.16463	.02854
230.00	2130.20	23.25	.15485	-56.84116	1.19811-144.03248	.02060	5.83074	59.47026	10.71583	48.88047	.02583
235.00	2032.47	19.85	.17366	-75.38710	1.19570-158.85847	.02060	5.11651	64.58678	9.70948	58.58994	.02323
240.00	1957.11	17.10	.19344	-92.04609	1.19346-172.14369	.02060	4.52706	69.11383	8.75816	67.34811	.02100
245.00	1882.51	14.87	.21399	-107.02155	1.19139-184.06460	.02059	4.04029	73.15412	7.88062	75.22872	.01909
250.00	1814.93	13.04	.23513	-120.50565	1.18944-194.78232	.02059	3.63647	76.79059	7.08126	82.30998	.01746
255.00	1753.64	11.55	.25665	-132.67167	1.18761-204.43920	.02059	3.29924	80.08983	6.35833	88.66832	.01606
260.00	1697.99	10.31	.27837	-143.66987	1.18589-213.16033	.02059	3.01554	83.10536	5.70489	94.37321	.01485
265.00	1647.36	9.28	.30012	-153.63601	1.18424-221.05406	.02059	2.77509	85.88045	5.11864	99.49185	.01381
270.00	1601.20	8.41	.32170	-162.68539	1.18268-228.21463	.02059	2.56975	88.45020	4.59083	104.08267	.01290
275.00	1559.07	7.68	.34295	-170.91788	1.18118-234.72288	.02059	2.39312	90.84332	4.11513	108.19780	.01211
280.00	1520.56	7.06	.36372	-178.42008	1.17973-240.64876	.02059	2.24017	93.08349	3.68571	111.88351	.01142
285.00	1484.70	6.54	.38385	-185.32762	1.17834-246.08865	.02059	2.10620	95.12040	3.33292	115.21650	.01082
290.00	1452.01	6.08	.40322	-191.59506	1.17700-251.01829	.02059	1.98815	97.17854	2.94149	118.15798	.01028
295.00	1422.03	5.70	.42170	-197.32683	1.17570-255.52312	.02059	1.88409	99.06263	2.62074	120.77872	.00994
300.00	1394.50	5.37	.43918	-202.57340	1.17443-259.64334	.02059	1.79200	100.85463	2.32823	123.10695	.00940
305.00	1369.22	5.08	.45556	-207.37864	1.17320-263.41397	.02059	1.71011	102.56475	2.06051	125.16746	.00903
310.00	1346.00	4.83	.47076	-211.78074	1.17201-266.86552	.02059	1.63702	104.20176	1.81454	126.98200	.00870
315.00	1324.66	4.62	.48471	-215.81306	1.17083-270.02462	.02059	1.57153	105.77330	1.58763	128.56963	.00842
320.00	1305.08	4.43	.49734	-219.50468	1.16969-272.91477	.02059	1.51271	107.28601	1.37736	129.94699	.00816
325.00	1287.12	4.28	.50859	-222.88101	1.16857-275.55610	.02059	1.45975	108.74576	1.18158	131.12857	.00794
330.00	1270.67	4.14	.51843	-225.96416	1.16747-277.96640	.02059	1.41200	110.15776	.99831	132.12687	.00774
335.00	1255.66	4.03	.52681	-228.77333	1.16639-280.16108	.02059	1.36890	111.52666	.82578	132.95265	.00757
340.00	1241.99	3.93	.53371	-231.32511	1.16532-282.15345	.02059	1.32998	112.85664	.66238	133.61503	.00742
345.00	1229.60	3.85	.53911	-233.63372	1.16428-283.95491	.02059	1.29488	114.15151	.50659	134.12163	.00729
350.00	1218.43	3.79	.54298	-235.71124	1.16325-285.57519	.02059	1.26325	115.41477	.35703	134.47865	.00719
355.00	1208.42	3.74	.54531	-237.56777	1.16223-287.02241	.02059	1.23485	116.64961	.21237	134.69102	.00710
360.00	1199.55	3.71	.54611	-239.21157	1.16122-288.30322	.02059	1.20944	117.85905	.07138	134.76240	.00702
360.00	895.70	1.03	1.46944	-293.87312	1.16122-330.52951	.02059	.00000	117.85905	.00000	134.76240	.00000
540.00	1000.00	1.03	.05461	-9.17099	.03930 -10.53334	.00069	.00000	117.85905	.00000	134.76240	.00000

THETA	T	R	V	H	S	U	N	FLAME SPEED, TEMP., AND EXTENT	MASS-FR.	BURN.-VOL.
166.96	2541.23	17.40	.00002	.00381	.00013	-.00663	.00000	675.034622541.23251	.00010	.00040
169.09	2541.59	18.30	.00024	.03789	.00125	-.06652	.00002	682.945302538.97421	.00100	.00394
171.46	2550.13	19.42	.00111	.19934	.00623	-.32450	.00010	713.023392548.45255	.00500	.01926
172.63	2554.24	20.14	.00215	.40765	.01246	-.64173	.00021	724.966202546.34534	.01000	.03774
174.23	2565.39	21.30	.00409	.86957	.02489	-1.23865	.00041	749.149672558.76233	.02000	.07287
176.58	2559.88	23.21	.00748	1.65727	.04960	-2.54880	.00083	787.523182577.77811	.04000	.13573
178.04	2570.83	24.84	.01053	2.63968	.07429	-3.69700	.00124	815.544772581.27966	.06000	.19235
179.13	2583.53	26.37	.01330	3.76738	.09896	-4.72468	.00165	841.622882586.37708	.08000	.24355
179.98	2596.26	27.83	.01583	5.02470	.12360	-5.64462	.00207	866.466642591.78738	.10000	.28985
181.05	2613.81	29.25	.01925	7.10050	.16051	-6.86679	.00269	904.052882606.94626	.13000	.35227
181.94	2629.80	32.01	.02232	9.38069	.19735	-7.91914	.00331	936.869772613.55429	.16000	.40744
182.62	2645.11	34.03	.02578	11.87568	.23416	-8.79253	.00393	968.722262620.31250	.19000	.45668
183.18	2659.40	36.03	.02758	14.54972	.27091	-9.51676	.00455	999.529852626.80408	.22000	.50095
183.68	2672.74	38.01	.02986	17.38460	.30762	-10.10685	.00518	1029.372482633.05392	.25000	.54102
184.11	2685.30	39.97	.03196	20.37083	.34430	-10.57071	.00580	1058.431352639.11267	.28000	.57751
184.56	2696.54	41.82	.03391	23.44285	.38092	-10.96352	.00642	1086.225572644.62439	.31000	.61091
185.02	2706.65	43.76	.03574	26.58592	.41749	-11.29765	.00704	1112.970472649.89545	.34000	.64163
185.50	2715.64	45.59	.03747	29.77385	.45402	-11.59515	.00767	1138.610402654.85202	.37000	.66999
185.99	2723.43	47.36	.03911	32.96706	.49049	-11.88934	.00829	1163.093432659.48666	.40000	.69629
186.32	2731.74	49.18	.04061	36.34620	.52694	-12.02789	.00891	1188.081882664.79422	.43000	.72079
186.88	2736.77	50.80	.04214	39.42682	.56329	-12.41989	.00953	1210.070592668.19183	.46000	.74361
187.55	2739.49	52.27	.04367	42.26245	.59955	-13.01923	.01015	1229.504712671.20404	.49000	.76496
188.65	2735.71	53.25	.04542	44.17893	.63561	-14.39687	.01077	1241.715762671.05820	.52000	.78492
190.32	2722.20	53.44	.04761	44.53704	.67139	-17.08724	.01139	1243.393162667.44315	.55000	.80362
192.02	2706.47	53.32	.05001	44.34684	.70711	-20.23540	.01201	1241.317762665.39664	.58000	.82134
193.71	2689.30	52.94	.05261	43.70057	.74277	-23.76197	.01262	1236.326932663.22443	.61000	.83818
195.35	2671.38	52.40	.05538	42.69584	.77839	-27.58404	.01324	1228.840532660.79025	.64000	.85419
196.69	2658.57	52.13	.05798	42.35821	.81415	-30.84226	.01386	1225.275922661.33151	.67000	.86951
198.17	2642.35	51.51	.06091	41.22326	.84980	-34.76254	.01447	1216.812062657.86435	.70000	.88410
199.68	2625.11	50.73	.06406	39.68678	.88542	-39.01279	.01509	1206.035962654.71237	.73000	.89804
201.79	2594.29	48.86	.06839	35.32605	.92068	-45.60464	.01570	1178.906342643.02560	.76000	.91128
203.55	2570.91	47.49	.07246	32.09012	.95616	-51.24720	.01631	1160.718802641.55038	.79000	.92402
205.04	2553.55	46.50	.07627	29.79376	.99180	-56.10423	.01693	1147.370862640.59128	.82000	.93627
206.77	2530.84	45.09	.08079	26.19225	1.02732	-62.03124	.01754	1126.786352633.49783	.85000	.94799
208.67	2504.72	43.42	.08594	21.63027	1.06276	-68.73860	.01816	1102.106692626.06119	.88000	.95923
210.71	2476.07	41.57	.09174	16.25354	1.09816	-76.10124	.01877	1074.736502618.31833	.91000	.97003
212.85	2445.81	39.63	.09816	10.23883	1.13354	-83.97101	.01938	1045.758302610.51782	.94000	.98041
215.66	2401.98	36.89	.10683	.97113	1.16861	-94.47467	.02000	1002.277992593.97488	.97000	.99038
220.48	2325.81	32.14	.12235	-15.70532	1.20333	-110.94678	.02061	920.747312556.23593	1.00000	1.00000

*** ALL INPUT DATA HAVE BEEN PROCESSED.
 AT LOC 45267

DATA SET B

RAPID FLAME PROPAGATION RATES

A MODEL SIMULATING THE INTERNAL-COMBUSTION ENGINE

ENGINE	FUEL		OPERATING CONDITIONS	
FOUR STROKE AUTOMOTIVE TYPE	OCTANE C8H18	AIR-FUEL RATIO	.92	
BORE 3.94 IN.	ACTIVATION ENERGY 39000.00 CAL./MOLE	PISTON TEMP	500.00 K	
STROKE 2.46 IN.	ENTHALPY OF COMBUSTION 1222770.00 CAL/MOLE	WALL TEMP	420.00 K	
ROD 7.06 IN.		MAN. TEMP	320.00 K	
CLEAR VOL. 3.33 CU.IN.		MAN. PR.	.99 ATM	
SWEPT VOL. 30.00 CU.IN.		EXH. PR.	1.03 ATM	
COMPRESSION RATIO 10.00 TO ONE		SPARK ADV.	16.00	
		SPEED	2800.00 RPM	

COMBUSTION OCCURS IN 39 INCREMENTS. ERRORS ARE LESS THAN .00100 PERCENT. FLAME SPEED FACTOR IS .15525000E 06
 HEAT TRANSFER FACTOR IS 1.00000 NORMAL RUN

SUMMARY OF RESULTS

PUMPING WORK	-.47611947E 00 CAL
NET CYCLE WORK	.14891748E 03 CAL
NET HEAT TRANSFERRED	.72684486E 02 CAL
INDICATED THERMAL EFFICIENCY	.36878417E 02 PERCENT
MEAN EFFECTIVE PRESSURE	.12510931E 02 ATM
RESIDUAL WEIGHT FRACTION	.33289108E-01
VOLUMETRIC EFFICIENCY	.97627462E 00 PERCENT

THETA	T	P	V	H	S	U	N	Q	QIOT	W	MIOT	H8
.00	344.84	.99	.54611	-22.58769	.95518	-35.68175	.01911	.00000	.00000	.00000	.00000	.00000
10.00	345.51	1.00	.54302	-22.48808	.95518	-35.60766	.01911	.00000	.00000	.00000	.00000	.00000
20.00	347.55	1.02	.53380	-22.18514	.95518	-35.38210	.01911	.00000	.00000	.00000	.00000	.00000
30.00	351.01	1.06	.51855	-21.68938	.95518	-34.99796	.01911	.00000	.00000	.00000	.00000	.00000
40.00	356.02	1.12	.49750	-20.92281	.95518	-34.44157	.01911	.00000	.00000	.00000	.00000	.00000
50.00	362.73	1.21	.47096	-19.91986	.95518	-33.69353	.01911	.00000	.00000	.00000	.00000	.00000
60.00	371.37	1.33	.43941	-18.62534	.95518	-32.72705	.01911	.00000	.00000	.00000	.00000	.00000
70.00	382.23	1.49	.40348	-16.99272	.95518	-31.50656	.01911	.00000	.00000	.00000	.00000	.00000
80.00	395.67	1.70	.36400	-14.96164	.95518	-29.98581	.01911	.00000	.00000	.00000	.00000	.00000
90.00	412.16	2.01	.32199	-12.45483	.95518	-28.10524	.01911	.00000	.00000	.00000	.00000	.00000
100.00	432.28	2.43	.27867	-9.37447	.95518	-25.78904	.01911	.00000	.00000	.00000	.00000	.00000
110.00	456.74	3.04	.23542	-5.59927	.95518	-22.94252	.01911	.00000	.00000	.00000	.00000	.00000
120.00	486.33	3.94	.19371	-9.85660	.95518	-19.45242	.01911	.00000	.00000	.00000	.00000	.00000
130.00	521.84	5.28	.15510	4.61714	.95518	-15.19795	.01911	.00000	.00000	.00000	.00000	.00000
140.00	563.67	7.30	.12106	11.31073	.95518	-10.09294	.01911	.00000	.00000	.00000	.00000	.00000
150.00	610.94	10.30	.09296	18.99131	.95518	-4.20704	.01911	.00000	.00000	.00000	.00000	.00000
160.00	659.41	14.36	.07199	26.99693	.95518	1.95803	.01911	.00000	.00000	.00000	.00000	.00000
164.00	677.09	16.13	.06580	29.94867	.95518	4.23851	.01911	.00000	.00000	.00000	.00000	.00000
166.59	687.41	17.25	.06250	31.68026	.95521	5.57040	.01911	.00000	.00000	.00000	.00000	.00000
168.46	694.60	18.06	.06047	32.89473	.95547	6.44144	.01911	.00000	.00000	.00000	.00000	.00000
170.54	703.47	19.11	.05856	34.40938	.95664	7.30770	.01912	.00000	.00000	.00000	.00000	.00000
172.97	718.22	20.96	.05680	37.00419	.96097	8.17112	.01914	.00000	.00000	.00000	.00000	.00000
175.02	732.77	22.93	.05571	39.59162	.96665	8.65004	.01917	.00000	.00000	.00000	.00000	.00000
176.30	744.60	24.64	.05522	41.81181	.97229	8.85403	.01920	.00000	.00000	.00000	.00000	.00000
177.23	755.07	26.25	.05496	43.88790	.97789	8.95470	.01923	.00000	.00000	.00000	.00000	.00000
177.98	764.68	27.79	.05480	45.88285	.98346	8.99964	.01927	.00000	.00000	.00000	.00000	.00000
178.89	777.90	30.04	.05467	48.78124	.99176	9.00586	.01931	.00000	.00000	.00000	.00000	.00000
179.70	790.02	32.23	.05462	51.59148	.99999	8.95507	.01936	.00000	.00000	.00000	.00000	.00000
180.22	801.25	34.38	.05461	54.37510	1.00817	8.89659	.01941	.00000	.00000	.00000	.00000	.00000
180.69	811.78	36.51	.05463	57.11790	1.01630	8.81556	.01946	.00000	.00000	.00000	.00000	.00000
181.11	821.71	38.61	.05466	59.83092	1.02437	8.72009	.01951	.00000	.00000	.00000	.00000	.00000
181.44	831.14	40.69	.05470	62.53645	1.03240	8.62762	.01956	.00000	.00000	.00000	.00000	.00000
181.82	840.06	42.75	.05475	65.21341	1.04039	8.51972	.01961	.00000	.00000	.00000	.00000	.00000
182.21	848.52	44.79	.05482	67.85229	1.04834	8.38773	.01966	.00000	.00000	.00000	.00000	.00000
182.60	856.55	46.79	.05490	70.42085	1.05623	8.20356	.01971	.00000	.00000	.00000	.00000	.00000
183.00	864.16	48.76	.05500	72.92402	1.06408	7.97081	.01976	.00000	.00000	.00000	.00000	.00000
183.38	871.45	50.71	.05510	75.39425	1.07189	7.71607	.01981	.00000	.00000	.00000	.00000	.00000
183.81	878.32	52.61	.05524	77.75526	1.07965	7.37494	.01986	.00000	.00000	.00000	.00000	.00000
184.31	884.74	54.43	.05541	79.99041	1.08737	6.93314	.01991	.00000	.00000	.00000	.00000	.00000
184.96	890.53	56.13	.05568	81.95088	1.09502	6.26530	.01996	.00000	.00000	.00000	.00000	.00000
187.72	890.51	56.12	.05720	80.45500	1.10224	2.70643	.02000	.00000	.00000	.00000	.00000	.00000
189.18	892.48	56.71	.05828	80.37848	1.10967	.34013	.02005	.00000	.00000	.00000	.00000	.00000
190.64	893.64	57.05	.05953	79.89698	1.11705	-2.36230	.02009	.00000	.00000	.00000	.00000	.00000
192.19	893.59	57.04	.06107	78.74187	1.12438	-5.62603	.02014	.00000	.00000	.00000	.00000	.00000
193.50	893.93	57.14	.06252	77.90079	1.13175	-8.62485	.02018	.00000	.00000	.00000	.00000	.00000
194.74	894.00	57.16	.06404	76.94101	1.13912	-11.72129	.02023	.00000	.00000	.00000	.00000	.00000
195.96	893.74	57.08	.06565	75.81867	1.14649	-14.95234	.02027	.00000	.00000	.00000	.00000	.00000
197.41	891.87	56.52	.06772	73.68669	1.15377	-19.02499	.02032	.00000	.00000	.00000	.00000	.00000
199.28	887.11	55.12	.07067	69.66226	1.16091	-24.61636	.02036	.00000	.00000	.00000	.00000	.00000
200.58	885.15	54.55	.07287	67.52435	1.16825	-28.75001	.02040	.00000	.00000	.00000	.00000	.00000
202.01	882.09	53.67	.07545	64.62853	1.17553	-33.45416	.02045	.00000	.00000	.00000	.00000	.00000
203.52	878.22	52.58	.07837	61.17201	1.18280	-38.62345	.02049	.00000	.00000	.00000	.00000	.00000
205.19	873.17	51.18	.08178	56.90371	1.19002	-44.46590	.02053	.00000	.00000	.00000	.00000	.00000
207.00	866.99	49.51	.08573	51.81745	1.19722	-50.98375	.02057	.00000	.00000	.00000	.00000	.00000
209.24	858.01	47.17	.09094	44.73105	1.20430	-59.13387	.02061	.00000	.00000	.00000	.00000	.00000
212.67	841.03	43.16	.09964	33.02007	1.21128	-71.12848	.02064	.00000	.00000	.00000	.00000	.00000
215.00	2496.65	39.87	.10600	22.89440	1.21050	-79.46541	.02063	.00000	.00000	.00000	.00000	.00000

220.00	2405.30	33.67	.12084	1.91183	1.20888	-96.62883	.02062	4.02338	29.01330	13.14004	31.93379	.03293
225.00	2316.82	28.56	.13718	-17.54491	1.20738	-112.41795	.02061	3.57032	32.58362	12.21880	44.15259	.02971
230.00	2233.11	24.38	.15485	-35.38738	1.20600	-126.80840	.02060	3.17933	35.76299	11.21111	55.36370	.02685
235.00	2154.93	20.97	.17366	-51.69755	1.20473	-139.90520	.02060	2.84695	38.60990	10.24985	65.61355	.02435
240.00	2082.75	18.20	.19344	-66.54113	1.20354	-151.78737	.02060	2.56607	41.17598	9.31610	74.92965	.02219
245.00	2016.53	15.92	.21399	-80.02390	1.20242	-162.55591	.02060	2.32956	43.50554	8.43897	83.36862	.02032
250.00	1955.96	14.06	.23513	-92.26486	1.20136	-172.31599	.02060	2.13025	45.63579	7.62983	90.99845	.01871
255.00	1900.65	12.51	.25665	-103.38026	1.20036	-181.16647	.02059	1.96171	47.59750	6.88877	97.88721	.01731
260.00	1850.15	11.23	.27837	-113.48287	1.19940	-189.20127	.02059	1.81850	49.41601	6.21630	104.10351	.01610
265.00	1804.01	10.16	.30012	-122.67315	1.19848	-196.50318	.02059	1.69612	51.11212	5.60579	109.70930	.01505
270.00	1761.86	9.25	.32170	-131.04151	1.19759	-203.14612	.02059	1.59092	52.70304	5.05203	114.76133	.01413
275.00	1723.32	8.49	.34295	-138.66824	1.19674	-209.19545	.02059	1.49997	54.20301	4.54937	119.31069	.01333
280.00	1688.07	7.84	.36372	-145.62468	1.19592	-214.70907	.02059	1.42091	55.62392	4.09271	123.40340	.01263
285.00	1655.81	7.29	.38385	-151.97346	1.19511	-219.73763	.02059	1.35183	56.97574	3.67673	127.08013	.01201
290.00	1626.29	6.82	.40322	-157.77007	1.19433	-224.32598	.02059	1.29117	58.26691	3.29718	130.37731	.01147
295.00	1599.27	6.41	.42170	-163.06317	1.19357	-228.51333	.02059	1.23768	59.50460	2.94967	133.32698	.01099
300.00	1574.55	6.06	.43918	-167.89594	1.19282	-232.33447	.02059	1.19033	60.69493	2.63081	135.95779	.01057
305.00	1551.95	5.76	.45556	-172.30633	1.19209	-235.81992	.02059	1.14828	61.84320	2.33717	138.29495	.01019
310.00	1531.19	5.50	.47076	-176.35016	1.19137	-239.01422	.02059	1.11081	62.95402	2.08349	140.37844	.00987
315.00	1512.26	5.27	.48471	-180.03237	1.19066	-241.92163	.02059	1.07717	64.03119	1.83024	142.20868	.00958
320.00	1494.65	5.08	.49734	-183.41496	1.18995	-244.58355	.02059	1.04709	65.07828	1.61482	143.82350	.00932
325.00	1478.92	4.91	.50859	-186.43637	1.18927	-246.96100	.02059	1.01954	66.09782	1.35792	145.18142	.00909
330.00	1464.69	4.77	.51843	-189.16477	1.18860	-249.10720	.02059	.99528	67.09310	1.15092	146.33233	.00890
335.00	1451.89	4.66	.52681	-191.61727	1.18793	-251.03575	.02059	.97359	68.06669	.95496	147.28729	.00873
340.00	1440.44	4.56	.53371	-193.80841	1.18727	-252.75827	.02059	.95422	69.02091	.76830	148.05559	.00858
345.00	1430.28	4.48	.53911	-195.75050	1.18662	-254.28457	.02059	.93699	69.95790	.58932	148.64490	.00846
350.00	1421.36	4.42	.54298	-197.45373	1.18597	-255.62281	.02059	.92172	70.87961	.41653	149.06143	.00836
355.00	1413.64	4.38	.54531	-198.92629	1.18533	-256.77954	.02059	.90829	71.78790	.24845	149.30987	.00828
360.00	1407.10	4.35	.54611	-200.17452	1.18470	-257.75986	.02059	.89659	72.68449	.08373	149.39360	.00821
360.00	1023.03	1.03	1.67833	-271.38074	1.18470	-313.24813	.02059	.00000	72.68449	.00000	149.39360	.00000
540.00	1000.00	1.03	.05461	-9.17099	.03930	-10.53334	.00069	.00000	72.68449	.00000	149.39360	.00000

THEIA	T	P	V	H	S	U	N	FLAME SPEED, TEMP., AND EXTENT MASS-FR. BURN.-VOL.
166.59	2537.66	17.25	.00002	.00372	.00013	-.00671	.00000	771.67351 2537.66077 .18781 .00010 .00040
168.46	2538.47	18.06	.00024	.03712	.00125	-.06715	.00002	787.31929 2536.13147 .39292 .00100 .00394
170.54	2547.11	19.11	.00113	.19570	.00624	-.32750	.00010	812.65478 2545.75754 .65649 .00500 .01931
171.57	2551.59	19.81	.00218	.40142	.01246	-.64684	.00021	826.30920 2544.25040 .80191 .01000 .03783
172.97	2563.24	20.96	.00415	.85976	.02490	-1.24666	.00041	854.19674 2556.95749 .1.04391 .02000 .07306
175.02	2573.47	22.93	.00762	1.80903	.04968	-2.42080	.00083	900.19459 2577.39182 1.35295 .04000 .13670
176.30	2587.55	24.64	.01069	2.91892	.07442	-3.46161	.00124	934.16209 2581.35593 1.56672 .06000 .19360
177.23	2601.62	26.25	.01346	4.17107	.09913	-4.38433	.00165	966.13378 2586.98367 1.70732 .08000 .24491
177.98	2614.82	27.79	.01597	5.54397	.12380	-5.20662	.00207	996.17546 2592.62112 1.83381 .10000 .29148
178.89	2633.05	30.04	.01935	7.80324	.16076	-6.27375	.00269	1041.68782 2608.50833 1.97336 .13000 .35392
179.70	2649.42	32.23	.02234	10.26546	.19764	-7.17237	.00331	1082.21582 2615.77118 2.09153 .16000 .40898
180.22	2665.00	34.38	.02502	12.94443	.23448	-7.88988	.00393	1120.86934 2622.70496 2.17344 .19000 .45811
180.69	2679.28	36.51	.02744	15.79047	.27126	-8.46850	.00456	1158.00291 2629.33017 2.24359 .22000 .50223
181.11	2692.54	38.61	.02964	18.79210	.30799	-8.91741	.00518	1194.47279 2635.98874 2.28757 .25000 .54215
181.44	2705.09	40.69	.03164	21.94964	.34468	-9.23616	.00580	1229.78461 2642.33673 2.33990 .28000 .57849
181.82	2716.81	42.75	.03350	25.23620	.38132	-9.44749	.00642	1264.00150 2648.33151 2.39527 .31000 .61177
182.21	2727.68	44.79	.03522	28.62974	.41792	-9.56973	.00705	1297.18150 2654.07062 2.45077 .34000 .64239
182.60	2737.52	46.79	.03682	32.08846	.45447	-9.63831	.00767	1329.32805 2659.54349 2.50575 .37000 .67066
183.00	2746.52	48.76	.03833	35.60845	.49097	-9.65641	.00829	1360.11256 2664.59454 2.56121 .40000 .69688
183.38	2755.00	50.71	.03975	39.20807	.52744	-9.60845	.00892	1390.41466 2669.67252 2.60628 .43000 .72130
183.81	2762.51	52.61	.04110	42.81483	.56386	-9.55570	.00954	1419.41899 2674.30014 2.66690 .46000 .74411
184.31	2769.07	54.43	.04242	46.40862	.60023	-9.51507	.01016	1446.94548 2678.54718 2.73771 .49000 .76548
184.96	2773.92	56.13	.04374	49.84914	.63654	-9.60519	.01079	1472.09947 2682.07141 2.83260 .52000 .78554
187.72	2759.65	56.12	.04601	50.36148	.67241	-12.17069	.01140	1470.01303 2673.53775 3.14277 .55000 .80429
189.18	2754.42	56.71	.04791	52.14348	.70849	-13.66146	.01202	1479.92172 2678.21100 3.42090 .58000 .82216
190.64	2747.74	57.05	.04995	53.59828	.74453	-15.42713	.01264	1484.45322 2677.75681 3.65930 .61000 .83912
192.19	2738.35	57.04	.05223	54.46889	.78051	-17.68371	.01326	1483.82405 2675.95715 3.90481 .64000 .85521
193.50	2731.28	57.14	.05443	55.63075	.81654	-19.69332	.01388	1485.47699 2676.78320 4.11216 .67000 .87054
194.74	2724.24	57.16	.05669	56.69188	.85257	-21.78640	.01450	1485.69456 2676.44659 4.28580 .70000 .88513
195.96	2716.99	57.08	.05903	57.60670	.88859	-24.00131	.01511	1484.44780 2675.77728 4.45759 .73000 .89905
197.41	2705.66	56.52	.06179	57.57859	.92453	-27.00529	.01573	1475.46739 2671.91589 4.64830 .76000 .91233
199.28	2686.53	55.12	.06537	55.74592	.96032	-31.51882	.01635	1453.77093 2664.15781 4.87979 .79000 .92501
200.58	2676.61	54.55	.06829	55.65901	.99632	-34.56545	.01696	1445.74739 2666.53363 5.05306 .82000 .93716
202.01	2663.94	53.67	.07159	54.82228	1.03226	-38.23622	.01758	1432.34364 2662.69824 5.21242 .85000 .94877
203.52	2649.49	52.58	.07523	53.40975	1.06818	-42.38402	.01819	1415.25578 2658.45670 5.37413 .88000 .95990
205.19	2632.27	51.18	.07937	51.16260	1.10406	-47.22247	.01881	1393.21266 2652.79474 5.54231 .91000 .97056
207.00	2612.34	49.51	.08408	48.05586	1.13991	-52.77007	.01942	1366.58902 2646.24710 5.71462 .94000 .98078
209.24	2584.88	47.17	.09008	42.89794	1.17564	-60.00957	.02003	1327.92632 2634.93097 5.90755 .97000 .99059
212.67	2539.10	43.16	.09964	33.02008	1.21128	-71.12848	.02064	1258.50751 2610.84439 6.17306 1.00000 1.00000

8*** ALL INPUT DATA HAVE BEEN PROCESSED.

AT LOC 45267

A MODEL SIMULATING THE INTERNAL-COMBUSTION ENGINE

ENGINE	FUEL	OPERATING CONDITIONS
FOUR STROKE AUTOMOTIVE TYPE	OCTANE C8H18	AIR-FUEL RATIO .92
BORE 3.54 IN.	ACTIVATION ENERGY 39000.00 CAL./MOLE	PISTON TEMP 500.00 K
STROKE 2.46 IN.	ENTHALPY OF COMBUSTION 1222770.00 CAL./MOLE	WALL TEMP 420.00 K
ROD 7.06 IN.		MAN. TEMP 320.00 K
CLEAR. VOL. 3.33 CU.IN.		MAN. PR. .99 ATM
SWEPT VOL. 30.00 CU.IN.		EXH. PR. 1.03 ATM
COMPRESSION RATIO 10.00 TO ONE		SPARK ADV. 16.00
		SPEED 2800.00 RPM

COMBUSTION OCCURS IN 39 INCREMENTS. ERRORS ARE LESS THAN .00100 PERCENT. FLAME SPEED FACTOR IS .15525000E 06
 HEAT TRANSFER FACTOR IS .00000 NORMAL RUN

SUMMARY OF RESULTS

PUMPING WORK	-.47611947E 00 CAL
NET CYCLE WORK	.16641193E 03 CAL
NET HEAT TRANSFERRED	.00000000E 00 CAL
INDICATED THERMAL EFFICIENCY	.41210800E 02 PERCENT
MEAN EFFECTIVE PRESSURE	.13980684E 02 ATM
RESIDUAL WEIGHT FRACTION	.33269108E-01
VOLUMETRIC EFFICIENCY	.97627462E 00 PERCENT

215.00	2608.10	41.72	.10600	50.95418	1.21963	-56.15980	.02067	.00000	.00000	8.76006	20.47806	.03644
220.00	2540.76	35.62	.12084	34.21611	1.21963	-70.03143	.02065	.00000	.00000	13.87163	34.34968	.03443
225.00	2474.68	30.54	.13718	18.40763	1.21962	-83.05276	.02063	.00000	.00000	13.02133	47.37101	.03140
230.00	2410.54	26.34	.15485	3.60244	1.21963	-95.17372	.02062	.00000	.00000	12.12097	59.49198	.02870
235.00	2349.07	22.88	.17366	-10.14951	1.21963	-106.36929	.02061	.00000	.00000	11.19557	70.68755	.02630
240.00	2291.04	20.03	.19344	-22.80630	1.21963	-116.62364	.02061	.00000	.00000	10.25434	80.94189	.02420
245.00	2236.77	17.67	.21399	-34.40623	1.21963	-125.98561	.02060	.00000	.00000	9.36197	90.30386	.02236
250.00	2186.52	15.72	.23513	-44.98645	1.21962	-134.49653	.02060	.00000	.00000	8.51093	98.81478	.02075
255.00	2140.13	14.09	.25665	-54.64090	1.21963	-142.24466	.02060	.00000	.00000	7.74813	106.56291	.01935
260.00	2097.51	12.73	.27837	-63.43280	1.21964	-149.28709	.02060	.00000	.00000	7.04243	113.60534	.01813
265.00	2058.50	11.59	.30012	-71.42675	1.21964	-155.68066	.02060	.00000	.00000	6.39358	119.99892	.01706
270.00	2022.88	10.63	.32170	-78.68540	1.21964	-161.47898	.02060	.00000	.00000	5.79831	125.79723	.01612
275.00	1990.43	9.81	.34295	-85.26850	1.21964	-166.73242	.02060	.00000	.00000	5.25344	131.05067	.01530
280.00	1960.94	9.11	.36372	-91.22992	1.21964	-171.48573	.02060	.00000	.00000	4.75331	135.80398	.01458
285.00	1934.20	8.51	.38385	-96.61953	1.21964	-175.78003	.02060	.00000	.00000	4.29430	140.09828	.01395
290.00	1910.01	8.00	.40322	-101.48194	1.21964	-179.65188	.02060	.00000	.00000	3.87185	143.97014	.01340
295.00	1888.20	7.57	.42170	-105.85745	1.21964	-183.13416	.02060	.00000	.00000	3.48228	147.45241	.01291
300.00	1868.59	7.19	.43918	-109.78214	1.21964	-186.25621	.02059	.00000	.00000	3.12205	150.57446	.01248
305.00	1850.93	6.87	.45556	-113.31330	1.21964	-189.06406	.02059	.00000	.00000	2.80785	153.38231	.01210
310.00	1835.21	6.59	.47076	-116.45000	1.21963	-191.55734	.02059	.00000	.00000	2.49329	155.87560	.01177
315.00	1821.32	6.35	.48471	-119.21792	1.21962	-193.75680	.02059	.00000	.00000	2.19946	158.07506	.01148
320.00	1809.16	6.15	.49734	-121.63934	1.21961	-195.68040	.02059	.00000	.00000	1.92360	159.99866	.01123
325.00	1798.64	5.98	.50859	-123.73337	1.21961	-197.34348	.02059	.00000	.00000	1.66308	161.66174	.01101
330.00	1789.66	5.83	.51843	-125.51704	1.21960	-198.75984	.02059	.00000	.00000	1.41636	163.07809	.01082
335.00	1782.18	5.72	.52681	-127.00403	1.21959	-199.94041	.02059	.00000	.00000	1.18057	164.25866	.01067
340.00	1776.12	5.62	.53371	-128.20592	1.21959	-200.89450	.02059	.00000	.00000	.95409	165.21275	.01054
345.00	1771.46	5.55	.53911	-129.13202	1.21959	-201.62957	.02059	.00000	.00000	.73507	165.94782	.01043
350.00	1768.14	5.50	.54298	-129.78951	1.21958	-202.15139	.02059	.00000	.00000	.52183	166.46965	.01035
355.00	1766.16	5.47	.54531	-130.18340	1.21958	-202.46399	.02059	.00000	.00000	.31260	166.78224	.01030
360.00	1765.48	5.46	.54611	-130.31673	1.21958	-202.56980	.02059	.00000	.00000	.10581	166.88805	.01026
360.00	1238.17	1.03	2.03129	-232.03647	1.21958	-282.70864	.02059	.00000	.00000	.00000	166.88805	.00000
340.00	1000.00	1.03	.05461	-9.17099	.03930	-10.53334	.00069	.00000	.00000	.00000	166.88805	.00000

THETA	T	P	V	H	S	U	N	FLAME SPEED, TEMP., AND EXTENT MASS-FR. BURN.	-VOL.		
166.59	2537.66	17.25	.00002	.00372	.00013	-.00671	.00000	771.673512537.66077	.18781	.00010	.00040
168.46	2538.47	18.06	.00024	.03712	.00125	-.06715	.00002	787.319232536.13147	.39292	.00100	.00394
170.54	2547.11	19.11	.00113	.19570	.00624	-.32750	.00010	812.654782545.75754	.65649	.00500	.01931
171.57	2551.59	19.81	.00218	.40142	.01246	-.64684	.00021	826.309202544.25040	.80191	.01000	.03783
172.97	2563.24	20.96	.00415	.85976	.02490	-1.24666	.00041	854.196742556.95749	1.04391	.02000	.07306
175.02	2584.30	22.95	.00764	1.92912	.04973	-2.31963	.00083	900.627652577.69019	1.35295	.04000	.13721
176.30	2600.08	24.67	.01073	3.12872	.07450	-3.28475	.00124	934.809582581.62927	1.56817	.06000	.19436
177.22	2614.36	26.28	.01351	4.45758	.09924	-4.14251	.00166	966.964332587.26184	1.70912	.08000	.24581
177.96	2627.40	27.84	.01603	5.90005	.12393	-4.90579	.00207	997.158742592.91687	1.83582	.10000	.29246
178.88	2645.34	30.10	.01941	8.25965	.16092	-5.88789	.00269	1042.898862608.86655	1.97516	.13000	.35495
179.68	2661.50	32.31	.02240	10.82178	.19784	-6.70147	.00331	1083.307502615.96823	2.09316	.16000	.41005
180.19	2676.42	34.47	.02507	13.57289	.23470	-7.35754	.00394	1122.168842622.94498	2.17499	.19000	.45913
180.66	2690.22	36.60	.02749	16.49191	.27150	-7.87382	.00456	1159.696672629.74106	2.24493	.22000	.50319
181.08	2703.06	38.71	.02968	19.56288	.30825	-8.26353	.00518	1195.942552636.22156	2.28755	.25000	.54305
181.42	2715.31	40.80	.03159	22.79269	.34496	-8.52061	.00580	1231.318052642.56088	2.34112	.28000	.57934
181.80	2726.77	42.87	.03354	26.15042	.38162	-8.67103	.00643	1265.905242648.71710	2.39650	.31000	.61257
182.19	2737.54	44.91	.03526	29.62587	.41825	-8.72330	.00705	1299.204712654.47815	2.45200	.34000	.64315
182.58	2747.47	46.93	.03686	33.18624	.45483	-8.70512	.00767	1331.496412659.97931	2.50698	.37000	.67139
182.99	2756.76	48.91	.03836	36.83384	.49137	-8.61440	.00830	1362.806872665.23969	2.56246	.40000	.69760
183.34	2765.61	50.88	.03978	40.57693	.52788	-8.44411	.00892	1393.430362670.40411	2.60762	.43000	.72200
183.77	2773.74	52.80	.04113	44.36818	.56435	-8.23413	.00954	1422.437332674.92868	2.66831	.46000	.74480
184.27	2781.33	54.66	.04245	48.21875	.60080	-7.97468	.01017	1450.399632679.28439	2.73937	.49000	.76618
184.92	2787.68	56.40	.04376	52.00648	.63721	-7.76917	.01079	1476.118482682.93909	2.83541	.52000	.78628
187.69	2782.77	56.57	.04606	54.16165	.67360	-8.94010	.01141	1477.032092675.64716	3.14814	.55000	.80544
189.14	2782.37	57.27	.04796	56.96879	.71000	-9.56140	.01203	1488.437852680.20718	3.42962	.58000	.82346
190.58	2780.64	57.75	.04999	59.54084	.74638	-10.38094	.01265	1494.920262680.15588	3.66925	.61000	.84051
192.11	2776.91	57.89	.05225	61.72704	.78275	-11.52528	.01327	1496.620332678.90005	3.91476	.64000	.85669
193.39	2774.25	58.13	.05441	64.05293	.81913	-12.55191	.01390	1500.284652679.99045	4.11751	.67000	.87202
194.62	2771.21	58.27	.05664	66.26236	.85550	-13.67640	.01452	1502.263962679.88040	4.29478	.70000	.88659
195.82	2767.82	58.32	.05894	68.35210	.89186	-14.90139	.01514	1502.891132679.56683	4.46671	.73000	.90044
197.25	2761.33	57.89	.06166	69.74002	.92823	-16.71625	.01576	1496.010992676.17554	4.65845	.76000	.91367
199.07	2749.59	56.70	.06515	69.89343	.96461	-19.56728	.01637	1477.760972669.51492	4.88957	.79000	.92631
200.33	2743.81	56.28	.06797	71.20387	1.00100	-21.44498	.01699	1472.070282671.99750	5.06122	.82000	.93832
201.72	2735.96	55.56	.07116	71.94901	1.03739	-23.79633	.01761	1460.765372668.50565	5.21959	.85000	.94979
203.19	2726.71	54.62	.07466	72.24518	1.07380	-26.52307	.01823	1446.700292665.04987	5.38031	.88000	.96075
204.80	2715.32	53.39	.07864	71.88298	1.11022	-29.80028	.01884	1427.577992660.10248	5.54752	.91000	.97124
206.56	2701.78	51.88	.08315	70.81988	1.14667	-33.66172	.01946	1403.979022654.30023	5.71934	.94000	.98126
208.70	2682.89	49.74	.08884	68.19744	1.18313	-38.82385	.02007	1369.248322644.27747	5.91089	.97000	.99085
211.95	2649.17	46.00	.09773	61.47587	1.21963	-47.39974	.02068	1306.696232623.09903	6.17132	1.00000	1.00000

**** ALL INPUT DATA HAVE BEEN PROCESSED.
AT LOC 45270

A MODEL SIMULATING THE INTERNAL-COMBUSTION ENGINE

OPERATING CONDITIONS

FUEL

ENGINE

FOUR STROKE AUTOMOTIVE TYPE OCTANE C8H18 AIR-FUEL RATIO .92
BORE 3.94 IN. ACTIVATION ENERGY 39000.00 CAL./MOLE PISTON TEMP 500.00 K
STROKE 2.46 IN. ENTHALPY OF WALL TEMP 420.00 K
ROD 7.06 IN. COMBUSTION 1222770.00 CAL/MOLE MAN. TEMP 320.00 K
CLEAR. VOL. 3.33 CU.IN. MAN. PR. .99 ATM
SWEEP VOL. 30.00 CU.IN. EXH. PR. 1.03 ATM
COMPRESSION RATIO 10.00 TO ONE SPARK ADV. 16.00
SPEED 2800.00 RPM

COMBUSTION OCCURS IN 39 INCREMENTS. ERRORS ARE LESS THAN .00100 PERCENT. FLAME SPEED FACTOR IS .15525000E 06
HEAT TRANSFER FACTOR IS 2.00000 NORMAL RUN

SUMMARY OF RESULTS

PUMPING WORK -.47611947E 00 CAL
NET CYCLE WORK .13452925E 03 CAL
NET HEAT TRANSFERED .12129069E 03 CAL
INDICATED THERMAL EFFICIENCY .33315267E 02 PERCENT
MEAN EFFECTIVE PRESSURE .11302140E 02 ATM
RESIDUAL WEIGHT FRACTION .33289107E-01
VOLUMETRIC EFFICIENCY .97627462E 00 PERCENT

THETA	T	P	V	H	S	U	N	Q	QTOT	W	WTOT	HB
.00	344.84	.99	.54611	-22.58769	.95518	-35.68175	.01911	.00000	.00000	.00000	.00000	.00000
10.00	345.51	1.00	.54302	-22.48808	.95518	-35.60766	.01911	.00000	.00000	.00000	-.07409	.00000
20.00	347.55	1.02	.53380	-22.18514	.95518	-35.38210	.01911	.00000	.00000	.00000	-.22555	.00000
30.00	351.01	1.06	.51855	-21.66938	.95518	-34.99796	.01911	.00000	.00000	.00000	-.68380	.00000
40.00	356.02	1.12	.49750	-20.92281	.95518	-34.44157	.01911	.00000	.00000	.00000	-1.24019	.00000
50.00	362.73	1.21	.47096	-19.91986	.95518	-33.69353	.01911	.00000	.00000	.00000	-.74803	.00000
60.00	371.37	1.33	.43941	-18.62534	.95518	-32.72705	.01911	.00000	.00000	.00000	-1.98822	.00000
70.00	382.23	1.49	.40348	-16.99272	.95518	-31.50656	.01911	.00000	.00000	.00000	-4.17519	.00000
80.00	395.67	1.70	.36400	-14.96164	.95518	-29.98581	.01911	.00000	.00000	.00000	-1.52076	.00000
90.00	412.16	2.01	.32199	-12.45483	.95518	-28.10524	.01911	.00000	.00000	.00000	-1.88057	.00000
100.00	432.28	2.43	.27867	-9.37447	.95518	-25.78904	.01911	.00000	.00000	.00000	-2.31620	.00000
110.00	456.74	3.04	.23542	-5.59927	.95518	-22.94252	.01911	.00000	.00000	.00000	-2.84652	.00000
120.00	486.33	3.94	.19371	-.98550	.95518	-19.45242	.01911	.00000	.00000	.00000	-3.49010	.00000
130.00	521.84	5.28	.15510	4.61714	.95518	-15.19795	.01911	.00000	.00000	.00000	-4.25447	.00000
140.00	563.67	7.30	.12106	11.31073	.95518	-10.09294	.01911	.00000	.00000	.00000	-5.10500	.00000
150.00	610.94	10.30	.09296	18.99131	.95518	-4.20704	.01911	.00000	.00000	.00000	-5.88590	.00000
160.00	659.41	14.36	.07199	26.99693	.95518	1.95803	.01911	.00000	.00000	.00000	-6.16508	.00000
164.00	677.09	16.13	.06580	29.94867	.95518	4.23851	.01911	.00000	.00000	.00000	-2.28048	.00000
166.59	687.41	17.25	.06250	31.68026	.95521	5.57040	.01911	.00000	.00000	.00000	-1.33189	.00000
168.46	694.60	18.06	.06047	32.89473	.95547	6.44144	.01911	.00000	.00000	.00000	-.87104	.00000
170.54	703.47	19.11	.05856	34.40938	.95664	7.30770	.01912	.00000	.00000	.00000	-.86625	.00000
171.57	709.18	19.81	.05775	35.39930	.95809	7.69132	.01912	.00000	.00000	.00000	-.38362	.00000
172.97	718.22	20.96	.05680	37.00419	.96097	8.17112	.01914	.00000	.00000	.00000	-.47981	.00000
176.30	744.39	24.61	.05522	41.56945	.97221	8.65213	.01920	.00000	.00000	.00000	-2.9033	.00000
177.24	754.82	26.21	.05495	43.56069	.97779	8.68166	.01923	.00000	.00000	.00000	-1.7213	.00000
177.99	764.39	27.74	.05480	45.48048	.98333	8.66325	.01927	.00000	.00000	.00000	-.10763	.00000
178.91	777.55	29.98	.05467	48.27117	.99160	8.57852	.01931	.00000	.00000	.00000	-.09482	.00000
179.72	789.61	32.16	.05462	50.97629	.99980	8.43896	.01936	.00000	.00000	.00000	-0.3687	.00000
180.74	800.82	34.30	.05461	53.68590	1.00796	8.31738	.01941	.00000	.00000	.00000	-0.0217	.00000
180.72	811.33	36.41	.05463	56.35429	1.01606	8.17298	.01946	.00000	.00000	.00000	-.02000	.00000
181.14	821.24	38.51	.05467	58.99815	1.02411	8.01854	.01951	.00000	.00000	.00000	-.03764	.00000
181.44	830.68	40.59	.05470	61.65984	1.03213	7.88811	.01956	.00000	.00000	.00000	-.04066	.00000
181.82	839.58	42.64	.05475	64.26532	1.04010	7.71909	.01961	.00000	.00000	.00000	-.04858	.00000
182.21	848.03	44.67	.05482	66.82491	1.04802	7.51935	.01966	.00000	.00000	.00000	-.06402	.00000
182.61	856.02	46.66	.05490	69.28779	1.05588	7.24517	.01971	.00000	.00000	.00000	-.09987	.00000
183.01	863.61	48.62	.05500	71.66936	1.06369	6.90882	.01976	.00000	.00000	.00000	-.12108	.00000
183.40	870.84	50.54	.05511	73.98711	1.07146	6.52444	.01981	.00000	.00000	.00000	-.14074	.00000
183.83	877.65	52.42	.05524	76.16504	1.07917	6.02762	.01986	.00000	.00000	.00000	-.18199	.00000
184.33	883.99	54.22	.05542	78.14676	1.08681	5.37058	.01990	.00000	.00000	.00000	-.22383	.00000
184.97	889.70	55.88	.05568	79.80318	1.09437	4.44446	.01995	.00000	.00000	.00000	-.33470	.00000
187.72	889.09	55.70	.05720	76.67012	1.10108	-.49692	.01999	.00000	.00000	.00000	2.09000	.00000
189.19	890.71	56.18	.05829	75.62439	1.10820	-3.67987	.02004	.00000	.00000	.00000	1.49720	.00000
190.66	891.50	56.41	.05955	74.11623	1.11527	-7.24564	.02008	.00000	.00000	.00000	1.74670	.00000
192.23	890.99	56.26	.06111	71.74547	1.12223	-11.52874	.02013	.00000	.00000	.00000	2.16231	.00000
193.56	890.86	56.22	.06260	69.75398	1.12926	-15.49055	.02017	.00000	.00000	.00000	2.05133	.00000
194.83	890.51	56.12	.06415	67.67433	1.13629	-19.52270	.02021	.00000	.00000	.00000	2.13499	.00000
196.07	889.83	55.92	.06580	65.41665	1.14331	-23.70063	.02026	.00000	.00000	.00000	2.25921	.00000
197.53	887.47	55.23	.06791	61.94257	1.15018	-28.88820	.02030	.00000	.00000	.00000	2.42422	.00000
199.45	881.87	53.61	.07095	56.01897	1.15674	-36.10619	.02034	.00000	.00000	.00000	4.09627	.00000
200.87	878.83	52.75	.07338	52.11953	1.16362	-41.63524	.02038	.00000	.00000	.00000	3.20164	.00000
202.34	875.14	51.72	.07608	47.74868	1.17047	-47.54948	.02042	.00000	.00000	.00000	3.47787	.00000
203.91	870.54	50.46	.07914	42.70534	1.17727	-54.01541	.02046	.00000	.00000	.00000	3.86274	.00000
205.63	864.69	48.90	.08272	36.70350	1.18398	-61.26536	.02050	.00000	.00000	.00000	4.39646	.00000
207.51	857.62	47.06	.08689	29.75418	1.19061	-69.28667	.02054	.00000	.00000	.00000	3.08621	.00000
209.85	847.44	44.53	.09242	20.38366	1.19701	-79.27808	.02058	.00000	.00000	.00000	6.24980	.00000
213.50	828.38	40.24	.10187	5.62875	1.20311	-93.64177	.02062	.00000	.00000	.00000	4.48885	.00000
215.00	2388.98	38.12	.10600	-1.97214	1.20217	-99.82658	.02061	.00000	.00000	.00000	3.90831	.00000
					1.20217	-99.82658	.02061	2.27649	47.18162		16.96321	.03259

220.00	2276.53	31.85	1.2084	-26.32447	1.19909-119.52989	.02060	7.28943	54.47105	12.41388	29.37709	.03150
225.00	2171.15	26.75	1.3718	-48.40530	1.19629-137.27710	.02060	6.30939	60.78044	11.43782	40.81491	.02811
230.00	2074.28	22.64	1.5485	-68.30939	1.19375-153.20713	.02060	5.49101	66.27145	10.43902	51.25393	.02516
235.00	1986.15	19.33	1.7366	-86.19291	1.19141-167.47982	.02060	4.81568	71.08712	9.45701	60.71094	.02262
240.00	1906.32	16.65	1.9344	-102.25384	1.18925-180.27188	.02059	4.26032	75.34745	8.53173	69.24268	.02045
245.00	1834.09	14.48	2.1399	-116.69198	1.18725-191.75277	.02059	3.80250	79.14995	7.67839	76.92107	.01860
250.00	1768.68	12.71	2.3513	-129.69328	1.18537-202.67681	.02059	3.42296	82.57291	6.90108	83.82215	.01701
255.00	1709.39	11.25	2.5665	-141.42221	1.18361-211.37902	.02059	3.10611	85.67901	6.19610	90.01825	.01565
260.00	1655.54	10.05	2.7837	-152.02805	1.18194-219.78105	.02059	2.83962	88.51863	5.56241	95.58067	.01448
265.00	1606.55	9.05	3.0012	-161.63865	1.18035-227.38673	.02059	2.61372	91.13235	4.99196	100.57263	.01346
270.00	1561.91	8.20	3.2170	-170.36466	1.17884-234.28572	.02059	2.42081	93.55315	4.47818	105.05081	.01258
275.00	1521.16	7.49	3.4295	-178.30217	1.17739-240.55567	.02059	2.25488	95.80803	4.01507	109.06588	.01182
280.00	1483.30	6.89	3.6372	-185.59553	1.17600-246.29945	.02059	2.11121	97.91924	3.63258	112.69845	.01115
285.00	1448.82	6.38	3.8385	-192.20501	1.17465-251.49784	.02059	1.98402	99.90326	3.21436	115.91282	.01055
290.00	1417.23	5.94	4.0322	-198.24159	1.17336-256.24172	.02059	1.87314	101.77640	2.87075	118.78356	.01003
295.00	1388.26	5.56	4.2170	-203.76101	1.17210-260.57552	.02059	1.77549	103.55188	2.55831	121.34187	.00958
300.00	1361.67	5.24	4.3918	-208.81164	1.17088-264.53783	.02059	1.68905	105.24093	2.27327	123.61514	.00917
305.00	1337.25	4.96	4.5556	-213.43558	1.16969-268.16232	.02059	1.61219	106.85311	2.01230	125.62744	.00882
310.00	1314.82	4.72	4.7076	-217.66955	1.16854-271.47836	.02059	1.54358	108.39669	1.77246	127.39990	.00850
315.00	1294.22	4.51	4.8471	-221.54559	1.16741-274.51161	.02059	1.48214	109.87883	1.55111	128.95101	.00822
320.00	1275.33	4.33	4.9734	-225.09172	1.16630-277.28452	.02059	1.42697	111.30580	1.34595	130.29696	.00798
325.00	1258.02	4.18	5.0859	-228.33237	1.16522-279.81667	.02059	1.37732	112.68311	1.15483	131.45179	.00776
330.00	1242.18	4.05	5.1843	-231.28885	1.16416-282.12514	.02059	1.33257	114.01568	.97590	132.42769	.00757
335.00	1227.74	3.94	5.2681	-233.97968	1.16311-284.22478	.02059	1.29222	115.30790	.80742	133.23512	.00740
340.00	1214.60	3.85	5.3371	-236.42078	1.16209-286.12837	.02059	1.25582	116.56372	.64777	133.88288	.00726
345.00	1202.72	3.77	5.3911	-238.62585	1.16108-287.84691	.02059	1.22302	117.78673	.49552	134.37840	.00713
350.00	1192.02	3.71	5.4298	-240.60650	1.16008-289.38971	.02059	1.19351	118.98024	.34929	134.72769	.00703
355.00	1182.46	3.66	5.4531	-242.37241	1.15910-290.76456	.02059	1.16704	120.14728	.20782	134.93551	.00694
360.00	1174.01	3.63	5.4611	-243.93148	1.15813-291.97784	.02059	1.14342	121.29070	.06986	135.00537	.00687
360.00	879.88	1.03	1.44349	-296.62187	1.15813-332.63102	.02059	.00000	121.29070	.00000	135.00537	.00000
540.00	1000.00	1.03	.05461	-9.17099	.03930 -10.53334	.00069	.00000	121.29070	.00000	135.00537	.00000

THETA	T	P	V	H	S	U	N	FLAME SPEED,	TEMP.,	AND EXTENT	MASS-FR.	BURN.-VOL.
								771.67351	2537.66077	.18781	.00010	.00040
166.59	2537.66	17.25	.00002	.00372	.00013	-.00671	.00000	787.31929	2536.13147	.39292	.00100	.00394
168.46	2538.47	18.06	.00024	.03712	.00125	-.06715	.00002	812.65478	2545.75754	.65649	.00500	.01931
170.54	2547.11	19.11	.00113	.19570	.00624	-.32750	.00010	826.30920	2544.25040	.80191	.01000	.03783
171.57	2551.59	19.81	.00218	.40142	.01246	-.64684	.00021	854.19674	2556.95749	1.04391	.02000	.07306
172.97	2563.24	20.96	.00415	.85976	.02490	-1.24666	.00041	899.75685	2577.09015	1.35295	.04000	.13619
175.02	2562.50	22.91	.00759	1.68882	.04963	-2.52191	.00083	933.50958	2581.08075	1.56525	.06000	.19283
176.30	2574.86	24.61	.01065	2.70918	.07434	-3.63814	.00124	965.27422	2586.68924	1.70550	.08000	.24399
177.24	2588.68	26.21	.01341	3.88398	.09902	-4.62612	.00165	995.17088	2592.31448	1.83175	.10000	.29048
177.99	2602.04	27.74	.01592	5.18697	.12367	-5.50765	.00207	1040.24724	2608.00507	1.97152	.13000	.35286
178.91	2620.56	29.98	.01929	7.34562	.16059	-6.66017	.00269	1080.70000	2615.32202	2.08987	.16000	.40791
179.72	2637.17	32.16	.02228	9.70875	.19745	-7.64281	.00331	1119.12622	2622.21634	2.17188	.19000	.45709
180.24	2653.45	34.30	.02496	12.31622	.23426	-8.42142	.00393	1156.57430	2629.08603	2.24223	.22000	.50126
180.72	2668.20	36.41	.02739	15.08787	.27102	-9.06334	.00455	1192.74619	2635.61893	2.28760	.25000	.54123
181.14	2681.90	38.51	.02959	18.02057	.30773	-9.57101	.00518	1228.04591	2642.00534	2.33843	.28000	.57765
181.44	2695.07	40.59	.03160	21.13108	.34440	-9.93018	.00580	1262.15971	2647.95618	2.39403	.31000	.61098
181.82	2707.01	42.64	.03345	24.34585	.38102	-10.20284	.00642	1295.23021	2653.67886	2.44918	.34000	.64164
182.21	2718.00	44.67	.03518	27.65979	.41760	-10.39305	.00705	1327.20909	2659.10870	2.50452	.37000	.66994
182.61	2727.70	46.66	.03678	31.01391	.45412	-10.55075	.00767	1357.79106	2664.11917	2.55995	.40000	.69618
183.00	2736.45	48.62	.03829	34.41290	.49059	-10.67197	.00829	1387.77905	2669.10239	2.60491	.43000	.72061
183.40	2744.48	50.54	.03971	37.86331	.52701	-10.75104	.00891	1416.49739	2673.69110	2.66540	.46000	.74342
183.83	2751.37	52.42	.04107	41.28928	.56337	-10.85215	.00954	1443.61513	2677.83835	2.73549	.49000	.76477
184.33	2756.92	54.22	.04239	44.63353	.59967	-11.02371	.01016	1468.68739	2681.45169	2.82965	.52000	.78481
184.97	2760.53	55.88	.04370	47.77283	.63588	-11.36984	.01078	1463.89445	2671.74725	3.13650	.55000	.80313
187.72	2736.88	55.70	.04594	46.69024	.67125	-15.28460	.01140	1471.58069	2676.14819	3.41047	.58000	.82087
189.19	2727.00	56.18	.04785	47.52176	.70703	-17.57725	.01201	1474.83195	2675.56134	3.64818	.61000	.83773
190.66	2715.67	56.41	.04989	47.96632	.74275	-20.19335	.01263	1472.04071	2673.23132	3.89288	.64000	.85373
192.23	2700.86	56.26	.05217	47.63982	.77836	-23.45470	.01325	1471.63460	2673.70422	4.10159	.67000	.86904
193.56	2689.12	56.22	.05440	47.66465	.81405	-26.41676	.01386	1470.00739	2673.10687	4.27556	.70000	.88366
194.83	2677.83	56.12	.05669	47.61189	.84973	-29.44085	.01448	1466.86111	2672.08551	4.44765	.73000	.89763
196.07	2666.49	55.92	.05907	47.39325	.88541	-32.60115	.01510	1430.59120	2667.92209	4.63694	.76000	.91095
197.53	2650.12	55.23	.06186	46.02265	.92094	-36.72038	.01571	1418.21089	2660.16757	4.86879	.79000	.92367
199.45	2623.16	53.61	.06554	42.29852	.95616	-42.79453	.01632	1402.09901	2656.31854	5.04402	.82000	.93593
200.87	2606.77	52.75	.06868	40.45646	.99169	-47.29157	.01694	1382.18069	2651.38156	5.20544	.85000	.94769
202.34	2589.03	51.72	.07210	38.12787	1.02720	-52.18570	.01755	1357.08076	2644.93701	5.36800	.88000	.95898
203.91	2569.06	50.46	.07589	35.10663	1.06265	-57.64741	.01817	1326.97392	2637.40720	5.53710	.91000	.96983
205.63	2545.65	48.90	.08022	31.09778	1.09801	-63.91603	.01878	1284.39039	2624.76984	5.90439	.97000	.99031
207.51	2518.91	47.06	.08517	26.09212	1.13330	-70.99281	.01940	1207.79457	2597.43137	6.17494	1.00000	1.00000
209.85	2482.42	44.53	.09152	18.60653	1.16835	-80.08986	.02001					
213.50	2423.04	40.24	.09187	5.62875	1.20311	-93.64177	.02062					

**** ALL INPUT DATA HAVE BEEN PROCESSED.
AT LOC 45267

DATA SET C

REDUCED FLAME PROPAGATION RATES

A MODEL SIMULATING THE INTERNAL-COMBUSTION ENGINE

ENGINE	FUEL	OPERATING CONDITIONS
FOUR STROKE AUTOMOTIVE TYPE	OCTANE C8H18	AIR-FUEL RATIO .92
BORE 3.94 IN.	ACTIVATION ENERGY 39000.00 CAL./MOLE	PISTON TEMP 500.00 K
STROKE 2.46 IN.	ENTHALPY OF COMBUSTION 1222770.00 CAL./MOLE	WALL TEMP 420.00 K
ROD 7.06 IN.		MAN. TEMP 320.00 K
CLEAR. VOL. 3.33 CU.IN.		MAN. PR. .99 ATM
SWEPT VOL. 30.00 CU.IN.		EXH. PR. 1.03 ATM
COMPRESSION RATIO 10.00 TO ONE		SPARK ADV. 16.00
		SPEED 2800.00 RPM

COMBUSTION OCCURS IN 39 INCREMENTS. ERRORS ARE LESS THAN .00100 PERCENT. FLAME SPEED FACIDR IS .11475000E 06
 HEAT TRANSFER FACTOR IS 1.00000 NORMAL RUN

SUMMARY OF RESULTS

PUMPING WORK	-.47611947E 00 CAL
NET CYCLE WORK	.14192007E 03 CAL
NET HEAT TRANSFERRED	.66320603E 02 CAL
INDICATED THERMAL EFFICIENCY	.35145557E 02 PERCENT
MEAN EFFECTIVE PRESSURE	.11923061E 02 ATM
RESIDUAL WEIGHT FRACTION	.33289107E-01
VOLUMETRIC EFFICIENCY	.97627462E 00 PERCENT

THETA	T	P	V	H	S	U	N	Q	QTOT	W	WTOT	HB
0.00	344.84	.99	.54611	-22.58769	.95518	-35.68175	.01911	.00000	.00000	.00000	.00000	.00000
10.00	345.51	1.00	.54302	-22.48808	.95518	-35.60766	.01911	.00000	.00000	.00000	.00000	.00000
20.00	347.55	1.02	.53380	-22.18514	.95518	-35.38210	.01911	.00000	.00000	.00000	.00000	.00000
30.00	351.01	1.06	.51855	-21.66938	.95518	-34.99796	.01911	.00000	.00000	.00000	.00000	.00000
40.00	356.02	1.12	.49750	-20.92281	.95518	-34.44157	.01911	.00000	.00000	.00000	.00000	.00000
50.00	362.73	1.21	.47096	-19.91986	.95518	-33.69353	.01911	.00000	.00000	.00000	.00000	.00000
60.00	371.37	1.33	.43941	-18.62534	.95518	-32.72705	.01911	.00000	.00000	.00000	.00000	.00000
70.00	382.23	1.49	.40348	-16.99272	.95518	-31.50656	.01911	.00000	.00000	.00000	.00000	.00000
80.00	395.67	1.70	.36400	-14.96164	.95518	-29.98581	.01911	.00000	.00000	.00000	.00000	.00000
90.00	412.16	2.01	.32199	-12.45483	.95518	-28.10524	.01911	.00000	.00000	.00000	.00000	.00000
100.00	432.28	2.43	.27867	-9.37447	.95518	-25.78904	.01911	.00000	.00000	.00000	.00000	.00000
110.00	456.74	3.04	.23542	-5.59927	.95518	-22.94252	.01911	.00000	.00000	.00000	.00000	.00000
120.00	486.33	3.94	.19371	-.98560	.95518	-19.45242	.01911	.00000	.00000	.00000	.00000	.00000
130.00	521.84	5.28	.15510	4.61714	.95518	-15.19795	.01911	.00000	.00000	.00000	.00000	.00000
140.00	563.67	7.30	.12106	11.31073	.95518	-10.09294	.01911	.00000	.00000	.00000	.00000	.00000
150.00	610.94	10.30	.09296	18.99131	.95518	-4.20704	.01911	.00000	.00000	.00000	.00000	.00000
160.00	659.41	14.36	.07199	26.99693	.95518	1.95803	.01911	.00000	.00000	.00000	.00000	.00000
164.00	677.09	16.13	.06580	29.94867	.95518	4.23851	.01911	.00000	.00000	.00000	.00000	.00000
167.45	690.58	17.60	.06153	32.21366	.95521	5.98332	.01911	.00000	.00000	.00000	.00000	.00000
169.93	699.32	18.61	.05908	33.69022	.95547	7.05779	.01911	.00000	.00000	.00000	.00000	.00000
172.68	709.03	19.79	.05698	35.35205	.95663	8.03895	.01912	.00000	.00000	.00000	.00000	.00000
174.04	714.79	20.52	.05619	36.35057	.95808	8.42980	.01912	.00000	.00000	.00000	.00000	.00000
175.91	723.46	21.65	.05536	37.89117	.96095	8.86026	.01914	.00000	.00000	.00000	.00000	.00000
178.66	736.46	23.45	.05470	40.14294	.96659	9.07455	.01917	.00000	.00000	.00000	.00000	.00000
180.38	746.70	24.96	.05462	42.01612	.97221	9.400379	.01920	.00000	.00000	.00000	.00000	.00000
181.61	755.73	26.35	.05472	43.76666	.97779	8.84557	.01923	.00000	.00000	.00000	.00000	.00000
182.60	764.02	27.68	.05490	45.44841	.98335	8.63999	.01927	.00000	.00000	.00000	.00000	.00000
183.82	775.43	29.61	.05524	47.89305	.99164	8.28125	.01931	.00000	.00000	.00000	.00000	.00000
184.89	785.77	31.45	.05565	50.24318	.99988	7.85851	.01936	.00000	.00000	.00000	.00000	.00000
185.69	795.57	33.28	.05602	52.62136	1.00807	7.46968	.01941	.00000	.00000	.00000	.00000	.00000
186.38	804.80	35.09	.05638	54.98821	1.01621	7.08130	.01946	.00000	.00000	.00000	.00000	.00000
186.94	813.68	36.88	.05671	57.37174	1.02432	6.71540	.01951	.00000	.00000	.00000	.00000	.00000
187.51	821.83	38.63	.05706	59.69606	1.03238	6.30862	.01956	.00000	.00000	.00000	.00000	.00000
188.08	829.49	40.32	.05745	61.95167	1.04041	5.85252	.01961	.00000	.00000	.00000	.00000	.00000
188.66	836.59	41.94	.05787	64.08827	1.04838	5.30383	.01965	.00000	.00000	.00000	.00000	.00000
189.25	843.17	43.49	.05833	66.10613	1.05631	4.66207	.01970	.00000	.00000	.00000	.00000	.00000
189.79	849.50	45.03	.05878	68.10725	1.06420	4.01102	.01975	.00000	.00000	.00000	.00000	.00000
190.48	854.85	46.36	.05939	69.75437	1.07203	3.07043	.01980	.00000	.00000	.00000	.00000	.00000
191.36	859.08	47.44	.06022	70.94065	1.07980	1.75024	.01985	.00000	.00000	.00000	.00000	.00000
194.87	851.33	45.48	.06420	66.36100	1.08711	-4.35642	.01989	.00000	.00000	.00000	.00000	.00000
196.90	848.52	44.79	.06698	64.19504	1.09465	-8.45549	.01993	.00000	.00000	.00000	.00000	.00000
198.85	845.23	43.99	.06997	61.77002	1.10219	-12.76670	.01998	.00000	.00000	.00000	.00000	.00000
200.92	840.55	42.87	.07347	58.55350	1.10968	-17.73275	.02002	.00000	.00000	.00000	.00000	.00000
202.52	837.94	42.26	.07643	56.40046	1.11727	-21.81617	.02006	.00000	.00000	.00000	.00000	.00000
204.23	834.36	41.43	.07978	53.69459	1.12484	-26.35529	.02011	.00000	.00000	.00000	.00000	.00000
205.96	830.29	40.50	.08342	50.67718	1.13241	-31.15113	.02015	.00000	.00000	.00000	.00000	.00000
207.92	824.57	39.23	.08781	46.68102	1.13992	-36.75425	.02019	.00000	.00000	.00000	.00000	.00000
210.29	816.16	37.42	.09351	41.09173	1.14735	-43.70840	.02023	.00000	.00000	.00000	.00000	.00000
212.25	810.14	36.17	.09851	36.75588	1.15493	-49.54172	.02028	.00000	.00000	.00000	.00000	.00000
214.10	804.71	35.07	.10348	32.76098	1.16256	-55.13235	.02032	.00000	.00000	.00000	.00000	.00000
216.14	798.10	33.77	.10925	27.97823	1.17017	-61.36817	.02036	.00000	.00000	.00000	.00000	.00000
218.43	790.14	32.26	.11601	22.26584	1.17775	-68.36222	.02040	.00000	.00000	.00000	.00000	.00000
220.75	782.14	30.79	.12320	16.43606	1.18536	-75.44677	.02044	.00000	.00000	.00000	.00000	.00000
222.95	774.99	29.53	.13033	11.06298	1.19305	-82.15270	.02049	.00000	.00000	.00000	.00000	.00000
225.72	765.00	27.84	.13965	3.69811	1.20064	-90.47902	.02053	.00000	.00000	.00000	.00000	.00000
229.34	751.08	25.62	.15243	-6.48157	1.20808	-101.08393	.02057	.00000	.00000	.00000	.00000	.00000
236.88	719.46	21.21	.18101	-27.29611	1.21531	-120.26429	.02060	.00000	.00000	.00000	.00000	.00000
240.00	224.61	19.44	.19344	-37.05705	1.21452	-128.13207	.02060	.00000	.00000	.00000	.00000	.00000

245.00	2154.89	17.02	.21399	-51.63729	1.21331-139.84531	.02060	2.70522	32.33511	9.00803	71.82845	.02171
250.00	2090.54	15.03	.23513	-64.90686	1.21216-150.47359	.02060	2.47796	34.81307	8.15032	79.97877	.01999
255.00	2031.42	13.38	.25665	-76.97657	1.21106-160.11945	.02060	2.28430	37.09737	7.36156	87.34033	.01850
260.00	1977.24	12.00	.27837	-87.95834	1.21002-168.88120	.02060	2.11891	39.21629	6.64283	93.98316	.01721
265.00	1927.63	10.85	.30012	-97.95552	1.20902-176.84669	.02060	1.97711	41.19340	5.98837	99.97153	.01608
270.00	1882.22	9.89	.32170	-107.06621	1.20805-184.09779	.02059	1.85497	43.04837	5.39613	105.36767	.01510
275.00	1840.63	9.07	.34295	-115.37619	1.20712-190.70532	.02059	1.74919	44.79756	4.85834	110.22601	.01424
280.00	1802.55	8.37	.36372	-122.96221	1.20622-196.73218	.02059	1.65711	46.45467	4.36974	114.59575	.01349
285.00	1767.65	7.78	.38385	-129.89193	1.20534-202.23350	.02059	1.57655	48.03122	3.92478	118.52053	.01282
290.00	1735.54	7.27	.40322	-136.25044	1.20449-207.27794	.02059	1.50573	49.53695	3.53871	122.05924	.01224
295.00	1706.13	6.84	.42170	-142.06296	1.20364-211.88637	.02059	1.44297	50.97991	3.16546	125.22470	.01173
300.00	1679.17	6.46	.43918	-147.37658	1.20282-216.09684	.02059	1.38734	52.36725	2.82314	128.04783	.01127
305.00	1654.49	6.14	.45556	-152.23259	1.20201-219.94270	.02059	1.33786	53.70511	2.50800	130.55583	.01087
310.00	1631.91	5.86	.47076	-156.66730	1.20121-223.45321	.02059	1.29372	54.99884	2.21678	132.77261	.01052
315.00	1611.28	5.62	.48471	-160.71251	1.20043-226.65396	.02059	1.25426	56.25309	1.94649	134.71911	.01021
320.00	1592.46	5.41	.49734	-164.39613	1.19967-229.56742	.02059	1.21890	57.47200	1.69455	136.41366	.00993
325.00	1575.34	5.23	.50859	-167.74256	1.19891-232.21319	.02059	1.18719	58.65919	1.45859	137.87224	.00969
330.00	1559.81	5.08	.51843	-170.77274	1.19817-234.60806	.02059	1.15874	59.81792	1.23613	139.10838	.00948
335.00	1545.80	4.96	.52681	-173.50529	1.19743-236.76710	.02059	1.13322	60.95114	1.02582	140.13419	.00930
340.00	1533.22	4.85	.53371	-175.95594	1.19671-238.70285	.02059	1.11036	62.06150	.82538	140.95958	.00914
345.00	1522.01	4.77	.53911	-178.13795	1.19599-240.42595	.02059	1.08995	63.15145	.63316	141.59273	.00901
350.00	1512.11	4.71	.54298	-180.06231	1.19528-241.94525	.02059	1.07178	64.22323	.44751	142.04025	.00889
355.00	1503.48	4.66	.54531	-181.73795	1.19458-243.26791	.02059	1.05572	65.27896	.26694	142.30718	.00881
360.00	1495.88	4.63	.54611	-183.17993	1.19391-244.39857	.02059	1.04165	66.32060	.08902	142.39620	.00874
360.00	1076.70	1.03	1.76639	-261.71504	1.19391-305.77901	.02059	.00000	66.32060	.00000	142.39620	.00000
540.00	1000.00	1.03	.05461	-9.17099	.03930 -10.53334	.00069	.00000	66.32060	.00000	142.39620	.00000

THETA	T	P	V	H	S	U	N	FLAME SPEED,	TEMP.,	AND EXTENT	MASS-FR.	BURN.-VOL.
								578.08720	2545.72406	.18781	.00010	.00040
167.45	2545.72	17.60	.00002	.00393	.00013	-.00653	.00000	592.18315	2542.46326	.39044	.00100	.00393
169.93	2545.38	18.61	.00023	.03883	.00125	-.06574	.00002	612.50503	2551.40607	.65041	.00500	.01920
172.68	2553.45	19.79	.00109	.20330	.00623	-.32124	.00010	622.45848	2548.51422	.80034	.01000	.03762
174.04	2556.93	20.52	.00211	.41386	.01245	-.63665	.00021	642.34013	2560.18518	1.04130	.02000	.07266
175.91	2566.85	21.65	.00402	.87543	.02488	-1.23398	.00043	672.76707	2577.38882	1.35023	.04000	.13590
178.66	2569.26	23.45	.00743	1.75694	.04962	-2.24633	.00083	694.50113	2580.17596	1.56373	.06000	.19266
180.38	2579.81	24.96	.01052	2.78550	.07434	-3.57463	.00124	714.81129	2584.87314	1.71380	.08000	.24399
181.61	2591.61	26.35	.01335	3.94602	.09903	-4.57429	.00165	733.99926	2589.78619	1.83215	.10000	.29065
182.60	2602.86	27.68	.01596	5.21152	.12369	-5.48679	.00207	763.10695	2603.66779	1.97342	.13000	.35331
183.82	2618.63	29.61	.01952	7.28701	.16064	-6.70820	.00269	788.37389	2609.65405	2.08953	.16000	.40867
184.89	2632.75	31.45	.02274	9.53519	.19753	-7.78633	.00331	813.13393	2616.11981	2.17998	.19000	.45806
185.69	2646.63	33.28	.02566	11.99049	.23437	-8.69171	.00393	837.00906	2622.14270	2.25598	.22000	.50244
186.38	2659.62	35.09	.02833	14.61007	.27118	-9.46025	.00455	860.36851	2628.10770	2.29947	.25000	.54259
186.94	2672.03	36.88	.03077	17.39487	.30794	-10.09086	.00518	882.58956	2633.40228	2.36213	.28000	.57914
187.51	2683.23	38.63	.03305	20.28375	.34466	-10.63516	.00580	903.72313	2638.34152	2.42543	.31000	.61260
188.08	2693.33	40.32	.03519	23.25568	.38134	-11.11046	.00642	923.69524	2642.89377	2.48909	.34000	.64334
188.66	2702.00	41.94	.03723	26.25148	.41796	-11.56680	.00704	942.51939	2647.11191	2.55440	.37000	.67172
189.25	2709.48	43.49	.03918	29.25993	.45455	-12.01355	.00767	960.97351	2651.54623	2.61194	.40000	.69804
189.79	2716.61	45.03	.04103	32.34709	.49109	-12.39500	.00829	976.40470	2654.03885	2.68796	.43000	.72254
190.48	2721.22	46.36	.04291	35.24367	.52758	-12.93797	.00891	988.79188	2655.58490	2.78704	.46000	.74541
191.36	2722.97	47.44	.04489	37.84277	.56400	-13.73244	.00953	961.73959	2631.52084	3.08247	.49000	.76686
194.87	2690.09	45.48	.04923	35.79969	.59997	-18.43115	.01014	955.05754	2636.84543	3.45078	.52000	.78712
196.90	2674.00	44.79	.05272	35.66959	.63617	-21.51543	.01076	945.18372	2633.55383	3.71376	.55000	.80623
198.85	2657.74	43.99	.05641	35.28828	.67236	-24.80573	.01138	930.85149	2627.65616	3.95535	.58000	.82427
200.92	2638.22	42.87	.06056	34.18302	.70850	-28.69796	.01199	924.13770	2628.88889	4.14893	.61000	.84134
202.52	2625.86	42.26	.06430	33.95009	.74475	-31.85751	.01261	913.52208	2624.62906	4.31976	.64000	.85751
204.23	2611.38	41.43	.06841	33.19733	.78098	-35.44694	.01323	901.72520	2620.67700	4.48777	.67000	.87285
205.96	2596.16	40.50	.07282	32.12410	.81720	-39.30014	.01384	884.83929	2613.53311	4.66277	.70000	.88742
207.92	2576.76	39.23	.07793	30.11509	.85337	-43.92703	.01446	860.56046	2602.76517	4.85467	.73000	.90127
210.29	2549.96	37.42	.08428	26.52967	.88946	-49.85287	.01507	845.04574	2601.76526	5.01345	.76000	.91445
212.25	2531.13	36.17	.09009	24.10846	.92569	-54.80613	.01569	830.69186	2598.43643	5.14118	.79000	.92699
214.10	2514.59	35.07	.09593	21.89341	.96197	-59.58313	.01631	812.72359	2591.37323	5.27472	.82000	.93894
216.14	2494.95	33.77	.10258	18.87000	.99824	-65.02144	.01692	791.58764	2583.03009	5.41356	.85000	.95034
218.43	2471.64	32.26	.11025	14.88286	1.03447	-71.24472	.01754	771.21184	2576.68668	5.54733	.88000	.96121
220.75	2448.63	30.79	.11842	10.69592	1.07074	-77.62298	.01815	753.62059	2572.37665	5.66695	.91000	.97159
222.95	2428.59	29.53	.12663	6.86913	1.10708	-83.69805	.01877	728.05232	2559.94394	5.80086	.94000	.98149
225.72	2400.05	27.84	.13707	1.00551	1.14333	-91.42872	.01938	692.95245	2542.15146	5.96103	.97000	.99096
229.34	2359.44	25.62	.15105	-7.75615	1.17942	-101.50292	.01999	614.26028	2487.33002	6.24214	1.00000	1.00000
236.88	2270.56	21.21	.18101	-27.29611	1.21531	-120.26429	.02060					

A MODEL SIMULATING THE INTERNAL-COMBUSTION ENGINE

OPERATING CONDITIONS

FUEL

ENGINE									
FOUR STROKE AUTOMOTIVE TYPE	OCTANE C8H18			AIR-FUEL RATIO	.92				
BORE	3.94 IN.	ACTIVATION ENERGY	39000.00 CAL./MOLE	PISTON TEMP	500.00 K				
STROKE	2.46 IN.	ENTHALPY OF		WALL TEMP	420.00 K				
ROD	7.06 IN.	COMBUSTION	1222770.00 CAL/MOLE	MAN. TEMP	320.00 K				
CLEAR. VOL.	3.33 CU.IN.			MAN. PR.	.99 ATM				
SWEPT VOL.	30.00 CU.IN.			EXH. PR.	1.03 ATM				
COMPRESSION RATIO	10.00 TO ONE			SPARK ADV.	16.00				
				SPEED	2800.00 RPM				

COMBUSTION OCCURS IN 39 INCREMENTS. ERRORS ARE LESS THAN .00100 PERCENT. FLAME SPEED FACTOR IS .11475000E 06
 HEAT TRANSFER FACTOR IS .00000 NORMAL RUN

SUMMARY OF RESULTS

PUMPING WORK	-.47611947E 00 CAL
NET CYCLE WORK	.15566373E 03 CAL
NET HEAT TRANSFERED	.00000000E 00 CAL
INDICATED THERMAL EFFICIENCY	.38549080E 02 PERCENT
MEAN EFFECTIVE PRESSURE	.13077700E 02 ATM
RESIDUAL WEIGHT FRACTION	.33289108E-01
VOLUMETRIC EFFICIENCY	.97627462E 00 PERCENT

THETA	T	P	V	H	S	U	N	Q	QTDT	W	WTDT	HB
10.00	344.84	.99	.54611	-22.58769	.95518	-35.68175	.01911	.00000	.00000	.00000	.00000	.00000
10.00	345.51	1.00	.54302	-22.48808	.95518	-35.60766	.01911	.00000	.00000	.00000	-.07409	.00000
20.00	347.55	1.02	.53380	-22.18514	.95518	-35.38210	.01911	.00000	.00000	.00000	-.22555	.00000
30.00	351.01	1.06	.51855	-21.66938	.95518	-34.99796	.01911	.00000	.00000	.00000	-.38415	.00000
40.00	356.02	1.12	.49750	-20.92281	.95518	-34.44157	.01911	.00000	.00000	.00000	-.55639	.00000
50.00	362.73	1.21	.47096	-19.91986	.95518	-33.69353	.01911	.00000	.00000	.00000	-.74803	.00000
60.00	371.37	1.33	.43941	-18.62534	.95518	-32.72705	.01911	.00000	.00000	.00000	-.96649	.00000
70.00	382.23	1.49	.40348	-16.99272	.95518	-31.50656	.01911	.00000	.00000	.00000	-1.22048	.00000
80.00	395.67	1.70	.36400	-14.96164	.95518	-29.98581	.01911	.00000	.00000	.00000	-1.52076	.00000
90.00	412.16	2.01	.32199	-12.45483	.95518	-28.10524	.01911	.00000	.00000	.00000	-1.88057	.00000
100.00	432.28	2.43	.27867	-9.37447	.95518	-25.78904	.01911	.00000	.00000	.00000	-2.31620	.00000
110.00	456.74	3.04	.23542	-5.59927	.95518	-22.94252	.01911	.00000	.00000	.00000	-2.84652	.00000
120.00	486.33	3.94	.19371	-.98560	.95518	-19.45242	.01911	.00000	.00000	.00000	-3.49010	.00000
130.00	521.84	5.28	.15510	4.61714	.95518	-15.19795	.01911	.00000	.00000	.00000	-4.25447	.00000
140.00	563.67	7.30	.12106	11.31073	.95518	-10.09294	.01911	.00000	.00000	.00000	-5.10500	.00000
150.00	610.94	10.30	.09296	18.99131	.95518	-4.20704	.01911	.00000	.00000	.00000	-5.88590	.00000
160.00	659.41	14.36	.07199	26.99693	.95518	1.95803	.01911	.00000	.00000	.00000	-6.16508	.00000
164.00	677.09	16.13	.06580	29.94867	.95518	4.23851	.01911	.00000	.00000	.00000	-2.28048	.00000
167.45	690.58	17.60	.06153	32.21366	.95521	5.98332	.01911	.00000	.00000	.00000	-1.74481	.00000
169.93	699.32	18.61	.05908	33.69022	.95547	7.05779	.01911	.00000	.00000	.00000	-1.07447	.00000
172.68	709.03	19.79	.05698	35.35205	.95663	8.03895	.01912	.00000	.00000	.00000	-.98117	.00000
174.04	714.79	20.52	.05619	36.35057	.95808	8.42980	.01912	.00000	.00000	.00000	-.39085	.00000
175.91	723.46	21.65	.05536	37.89117	.96095	8.86026	.01914	.00000	.00000	.00000	-.43045	.00000
178.66	736.63	23.48	.05470	40.33485	.96666	9.23429	.01917	.00000	.00000	.00000	-.37404	.00000
180.37	746.97	25.00	.05462	42.34777	.97231	9.28034	.01920	.00000	.00000	.00000	-.04605	.00000
181.61	756.09	26.40	.05472	44.21102	.97794	9.21660	.01923	.00000	.00000	.00000	.06374	.00000
182.59	764.44	27.75	.05490	45.99293	.98353	9.09521	.01927	.00000	.00000	.00000	.12140	.00000
183.79	775.94	29.70	.05523	48.58010	.99186	8.85646	.01931	.00000	.00000	.00000	.23875	.00000
184.86	786.34	31.55	.05564	51.06843	1.00014	8.55076	.01936	.00000	.00000	.00000	.30576	.00000
185.66	796.20	33.40	.05600	53.56069	1.00836	8.25848	.01941	.00000	.00000	.00000	.29222	.00000
186.34	805.47	35.22	.05636	56.03376	1.01654	7.96030	.01946	.00000	.00000	.00000	.29819	.00000
186.85	814.43	37.06	.05665	58.54546	1.02467	7.70218	.01951	.00000	.00000	.00000	.25811	.00000
187.41	822.70	38.82	.05700	60.97175	1.03276	7.38241	.01956	.00000	.00000	.00000	.31978	.00000
187.98	830.39	40.52	.05738	63.31671	1.04081	7.00232	.01961	.00000	.00000	.00000	.37988	.00000
188.56	837.55	42.17	.05779	65.59001	1.04882	6.57000	.01966	.00000	.00000	.00000	.42684	.00000
189.15	844.22	43.74	.05825	67.79362	1.05680	6.08598	.01971	.00000	.00000	.00000	.43252	.00000
189.67	850.66	45.31	.05867	70.00824	1.06475	5.61579	.01976	.00000	.00000	.00000	.48411	.00000
190.36	856.13	46.69	.05928	71.95383	1.07267	4.92825	.01981	.00000	.00000	.00000	.47009	.00000
191.22	860.59	47.83	.06008	73.58503	1.08056	3.98436	.01986	.00000	.00000	.00000	.68754	.00000
194.71	853.72	46.08	.06400	70.87076	1.08844	-.55354	.01990	.00000	.00000	.00000	.94389	.00000
196.71	851.51	45.53	.06671	69.94768	1.09634	-3.60945	.01994	.00000	.00000	.00000	4.53790	.00000
198.65	848.78	44.85	.06964	68.78059	1.10426	-6.86784	.01999	.00000	.00000	.00000	3.05591	.00000
200.69	844.73	43.87	.07308	67.02077	1.11219	-10.61803	.02003	.00000	.00000	.00000	3.25839	.00000
202.23	842.84	43.41	.07588	66.15816	1.12014	-13.62461	.02008	.00000	.00000	.00000	3.75019	.00000
203.90	839.85	42.71	.07913	64.77960	1.12810	-17.05995	.02013	.00000	.00000	.00000	3.00658	.00000
205.59	836.38	41.89	.08264	63.12082	1.13607	-20.72871	.02017	.00000	.00000	.00000	3.43534	.00000
207.51	831.34	40.74	.08688	60.67106	1.14405	-25.05440	.02021	.00000	.00000	.00000	3.66876	.00000
209.81	823.89	39.08	.09233	56.97514	1.15206	-30.41421	.02026	.00000	.00000	.00000	4.32569	.00000
211.69	818.71	37.96	.09706	54.32340	1.16010	-34.90705	.02030	.00000	.00000	.00000	5.35982	.00000
213.53	813.61	36.88	.10195	51.64920	1.16817	-39.41942	.02034	.00000	.00000	.00000	4.49284	.00000
215.50	807.79	35.69	.10740	48.53179	1.17627	-44.29915	.02039	.00000	.00000	.00000	4.51237	.00000
217.68	800.78	34.29	.11375	44.69444	1.18439	-49.77492	.02043	.00000	.00000	.00000	4.87973	.00000
219.88	793.74	32.93	.12047	40.75315	1.19255	-55.33587	.02047	.00000	.00000	.00000	5.47577	.00000
221.97	787.44	31.75	.12712	37.13013	1.20074	-60.63279	.02051	.00000	.00000	.00000	5.56095	.00000
224.59	778.53	30.15	.13579	31.91554	1.20897	-67.24328	.02055	.00000	.00000	.00000	5.29691	.00000
227.98	766.12	28.03	.14756	24.50374	1.21724	-75.66112	.02059	.00000	.00000	.00000	6.41049	.00000
234.78	738.03	23.77	.17281	7.86010	1.22559	-91.64307	.02063	.00000	.00000	.00000	8.41784	.00000
240.00	2367.83	20.71	.19344	-5.74319	1.22559	-102.75016	.02062	.00000	.00000	.00000	11.10710	.00000
											67.06842	.02508

245.00	2313.92	18.29	.21399	-17.67298	1.22558-112.44208	.02061	.00000	9.69191	76.76033	.02311
250.00	2263.51	16.28	.23513	-28.58864	1.22558-121.27368	.02061	.00000	8.83160	85.59193	.02147
255.00	2216.88	14.60	.25665	-38.51925	1.22557-129.28151	.02060	.00000	8.00783	93.59976	.02004
260.00	2173.78	13.20	.27837	-47.58189	1.22559-136.57005	.02060	.00000	7.28854	100.88830	.01878
265.00	2134.13	12.02	.30012	-55.83420	1.22559-143.19288	.02060	.00000	6.62284	107.51114	.01768
270.00	2097.81	11.02	.32170	-63.33560	1.22560-149.20294	.02060	.00000	6.01006	113.52119	.01672
275.00	2064.64	10.17	.34295	-70.14353	1.22560-154.64998	.02060	.00000	5.44704	118.96823	.01587
280.00	2034.43	9.45	.36372	-76.31217	1.22560-159.58000	.02060	.00000	4.93002	123.89825	.01513
285.00	2007.00	8.84	.38385	-81.89165	1.22560-164.03511	.02060	.00000	4.45510	128.35336	.01448
290.00	1982.16	8.31	.40322	-86.92705	1.22560-168.05267	.02060	.00000	4.01756	132.37092	.01390
295.00	1959.74	7.85	.42170	-91.45946	1.22560-171.66655	.02060	.00000	3.61388	135.98480	.01339
300.00	1939.58	7.46	.43918	-95.52568	1.22560-174.90691	.02060	.00000	3.24036	139.22516	.01295
305.00	1921.53	7.13	.45556	-99.15836	1.22560-177.80038	.02060	.00000	2.89347	142.11863	.01256
310.00	1905.47	6.84	.47076	-102.38671	1.22560-180.37074	.02060	.00000	2.57036	144.68899	.01222
315.00	1891.14	6.59	.48471	-105.26145	1.22560-182.65874	.02060	.00000	2.28800	146.97700	.01192
320.00	1878.59	6.38	.49734	-107.77556	1.22559-184.65905	.02059	.00000	2.00031	148.97730	.01166
325.00	1867.73	6.21	.50859	-109.94998	1.22558-186.38869	.02059	.00000	1.72964	150.70695	.01143
330.00	1858.47	6.06	.51843	-111.80164	1.22558-187.86126	.02059	.00000	1.47256	152.17951	.01124
335.00	1850.74	5.94	.52681	-113.34516	1.22557-189.08854	.02059	.00000	1.22728	153.40679	.01108
340.00	1844.50	5.84	.53371	-114.59267	1.22556-190.08030	.02059	.00000	.99176	154.39855	.01094
345.00	1839.68	5.77	.53911	-115.55388	1.22556-190.84436	.02059	.00000	.76406	155.16261	.01083
350.00	1836.26	5.71	.54298	-116.23626	1.22556-191.38674	.02059	.00000	.54238	155.70499	.01075
355.00	1834.21	5.68	.54531	-116.64505	1.22555-191.71163	.02059	.00000	.32489	156.02988	.01069
360.00	1833.52	5.67	.54611	-116.78341	1.22555-191.82160	.02059	.00000	.10997	156.13985	.01066
360.00	1278.38	1.03	2.09724	-224.52061	1.22555-276.83809	.02059	.00000	.00000	156.13985	.00000
540.00	1000.00	1.03	.05461	-9.17099	.03930 -10.53334	.00069	.00000	.00000	156.13985	.00000

THETA	T	P	V	H	S	U	N	FLAME SPEED, TEMP., AND EXTENT	MASS-FR. BURN.-VOL.			
167.45	2545.72	17.60	.00002	.00393	.00013	-.00653	.00000	578.08720	2545.72406	.18781	.00010	.00040
169.93	2545.38	18.61	.00023	.03883	.00125	-.06574	.00002	592.18315	2542.46326	.39044	.00100	.00393
172.68	2553.45	19.79	.00109	.20330	.00623	-.32124	.00010	612.50503	2551.40607	.65041	.00500	.01920
174.04	2556.93	20.52	.00211	.41386	.01245	-.63665	.00021	622.45848	2548.51422	.80034	.01000	.03762
175.91	2566.85	21.65	.00402	.87543	.02488	-1.23398	.00041	642.34013	2560.18518	1.04140	.02000	.07266
178.66	2584.12	23.48	.00747	1.92084	.04969	-2.32737	.00083	673.22456	2577.81104	1.35023	.04000	.13660
180.37	2597.08	25.00	.01058	3.07264	.07445	-3.33271	.00124	695.13030	2580.51791	1.56572	.06000	.19371
181.61	2609.02	26.40	.01342	4.33404	.09917	-4.24713	.00166	715.82738	2585.40479	1.71625	.08000	.24522
182.59	2619.97	27.75	.01603	5.69070	.12387	-5.08249	.00207	735.03692	2590.21384	1.83472	.10000	.29197
183.79	2635.28	29.70	.01959	7.89766	.16085	-6.19258	.00269	764.37862	2604.18008	1.97567	.13000	.35470
184.86	2649.09	31.55	.02282	10.27751	.19779	-7.15889	.00331	789.85182	2610.19501	2.09127	.16000	.41010
185.66	2662.29	33.40	.02573	12.84132	.23467	-7.97206	.00393	814.75306	2616.66458	2.18191	.19000	.45943
186.34	2674.70	35.22	.02839	15.56443	.27150	-8.65256	.00456	838.79264	2622.71014	2.25769	.22000	.50375
186.85	2686.79	37.06	.03081	18.46067	.30829	-9.18853	.00518	862.45916	2628.82419	2.30110	.25000	.54381
187.41	2697.59	38.82	.03308	21.45009	.34503	-9.64700	.00580	884.99043	2634.12054	2.36330	.28000	.58028
187.98	2707.25	40.52	.03521	24.51151	.38174	-10.04602	.00642	906.26539	2639.07407	2.42648	.31000	.61365
188.56	2716.00	42.17	.03724	27.64155	.41840	-10.38817	.00705	926.45988	2643.70993	2.48996	.34000	.64435
189.15	2723.98	43.74	.03918	30.83146	.45504	-10.68068	.00767	945.57439	2648.02441	2.55512	.37000	.67272
189.67	2731.76	45.31	.04101	34.12457	.49164	-10.88715	.00829	964.44866	2652.61188	2.61282	.40000	.69902
190.36	2737.62	46.69	.04289	37.31379	.52822	-11.18168	.00891	980.54672	2655.37570	2.68823	.43000	.72354
191.22	2741.49	47.83	.04485	40.34232	.56476	-11.61206	.00954	993.45390	2657.09464	2.78745	.46000	.74646
194.71	2720.57	46.08	.04919	40.09467	.60130	-14.79685	.01015	969.18217	2634.70956	3.08272	.49000	.76853
196.71	2711.18	45.53	.05264	41.16885	.63786	-16.86822	.01077	964.14987	2640.09195	3.45513	.52000	.78900
198.65	2701.23	44.85	.05629	42.01737	.67443	-19.12770	.01139	955.84516	2637.16284	3.71901	.55000	.80826
200.69	2688.86	43.87	.06040	42.34135	.71102	-21.82553	.01201	943.50600	2632.03912	3.96184	.58000	.82648
202.23	2681.94	43.41	.06401	43.37145	.74763	-23.92970	.01263	938.62103	2633.80780	4.15615	.61000	.84355
203.90	2672.89	42.71	.06803	43.93496	.78424	-26.42392	.01325	929.55212	2629.80783	4.32679	.64000	.85971
205.59	2662.99	41.89	.07231	44.21465	.82086	-29.15444	.01386	919.34918	2626.35846	4.49550	.67000	.88953
207.51	2649.82	40.74	.07728	43.74904	.85750	-32.50614	.01448	904.42387	2620.04617	4.67164	.70000	.90334
209.81	2631.37	39.08	.08340	42.09789	.89416	-36.84459	.01510	882.36642	2610.35510	4.86408	.73000	.91638
211.69	2618.62	37.96	.08894	41.31693	.93086	-40.45244	.01571	868.95744	2609.93253	5.02241	.76000	.92875
213.53	2606.34	36.88	.09468	40.45574	.96759	-44.12512	.01633	854.95669	2606.04041	5.14973	.79000	.94052
215.50	2592.59	35.69	.10101	39.12011	1.00434	-48.18968	.01695	838.77357	2600.23218	5.28362	.82000	.95172
217.68	2576.18	34.29	.10826	37.03450	1.04111	-52.87383	.01756	819.59791	2592.82236	5.42165	.85000	.96236
219.88	2559.93	32.93	.11594	34.77195	1.07793	-57.70032	.01818	801.17214	2587.26434	5.55447	.88000	.97247
221.97	2545.63	31.75	.12362	32.74257	1.11478	-62.32931	.01879	785.29438	2583.53806	5.67318	.91000	.98211
224.59	2525.03	30.15	.13336	29.08295	1.15166	-68.30213	.01941	761.74409	2572.36276	5.80623	.94000	.99129
227.98	2495.74	28.03	.14627	23.15170	1.18859	-76.14045	.02002	729.62302	2556.78836	5.96489	.97000	.99129
234.78	2427.65	23.77	.17281	7.86011	1.22559	-91.64306	.02063	657.46908	2508.96545	6.23693	1.00000	1.00000

**** ALL INPUT DATA HAVE BEEN PROCESSED.
AT LOC 45267

A MODEL SIMULATING THE INTERNAL-COMBUSTION ENGINE

ENGINE	FUEL		OPERATING CONDITIONS	
FOUR STROKE AUTOMOTIVE TYPE	OCTANE C8H18	AIR-FUEL RATIO	.92	
BORE 3.94 IN.	ACTIVATION ENERGY 39000.00 CAL./MOLE	PISTON TEMP	500.00 K	
STROKE 2.46 IN.	ENTHALPY OF	WALL TEMP	420.00 K	
ROD 7.06 IN.	COMBUSTION 1222770.00 CAL/MOLE	MAN. TEMP	320.00 K	
CLEAR VOL. 3.33 CU.IN.		MAN. PR.	.99 ATM	
SWEPT VOL. 30.00 CU.IN.		EXH. PR.	1.03 ATM	
COMPRESSION RATIO 10.00 TO ONE		SPARK ADV.	16.00	
		SPEED	2800.00 RPM	

COMBUSTION OCCURS IN 39 INCREMENTS. ERRORS ARE LESS THAN .00100 PERCENT. FLAME SPEED FACTOR IS .11475000E 06
 HEAT TRANSFER FACTOR IS 2.00000 NORMAL RUN

SUMMARY OF RESULTS

PUMPING WORK	- .47611947E 00 CAL
NET CYCLE WORK	.13045873E 03 CAL
NET HEAT TRANSFERED	.11131061E 03 CAL
INDICATED THERMAL EFFICIENCY	.32307231E 02 PERCENT
MEAN EFFECTIVE PRESSURE	.10960165E 02 ATM
RESIDUAL WEIGHT FRACTION	.33289108E-01
VOLUMETRIC EFFICIENCY	.97627462E 00 PERCENT

THETA	T	P	V	H	S	U	N	Q	QTOT	W	WTOT	HB
-0.00	344.84	.99	.54611	-22.58769	.95518	-35.68175	.01911	.00000	.00000	.00000	.00000	.00000
10.00	345.51	1.00	.54302	-22.48808	.95518	-35.60766	.01911	.00000	.00000	.00000	.00000	.00000
20.00	347.55	1.02	.53380	-22.18514	.95518	-35.38210	.01911	.00000	.00000	.00000	.00000	.00000
30.00	351.01	1.06	.51855	-21.66938	.95518	-34.99796	.01911	.00000	.00000	.00000	.00000	.00000
40.00	356.02	1.12	.49750	-20.92281	.95518	-34.44157	.01911	.00000	.00000	.00000	.00000	.00000
50.00	362.73	1.21	.47096	-19.91986	.95518	-33.69353	.01911	.00000	.00000	.00000	.00000	.00000
60.00	371.37	1.33	.43941	-18.62534	.95518	-32.72705	.01911	.00000	.00000	.00000	.00000	.00000
70.00	382.23	1.49	.40348	-16.99272	.95518	-31.50656	.01911	.00000	.00000	.00000	.00000	.00000
80.00	395.67	1.70	.36400	-14.96164	.95518	-29.98581	.01911	.00000	.00000	.00000	.00000	.00000
90.00	412.16	2.01	.32199	-12.45483	.95518	-28.10524	.01911	.00000	.00000	.00000	.00000	.00000
100.00	432.28	2.43	.27867	-9.37447	.95518	-25.78904	.01911	.00000	.00000	.00000	.00000	.00000
110.00	456.74	3.04	.23542	-5.59927	.95518	-22.94252	.01911	.00000	.00000	.00000	.00000	.00000
120.00	486.33	3.94	.19371	-0.98560	.95518	-19.45242	.01911	.00000	.00000	.00000	.00000	.00000
130.00	521.84	5.28	.15510	4.61714	.95518	-15.19795	.01911	.00000	.00000	.00000	.00000	.00000
140.00	563.67	7.30	.12106	11.31073	.95518	-10.09294	.01911	.00000	.00000	.00000	.00000	.00000
150.00	610.94	10.30	.09296	18.99131	.95518	-4.20704	.01911	.00000	.00000	.00000	.00000	.00000
160.00	659.41	14.36	.07199	26.99693	.95518	1.95803	.01911	.00000	.00000	.00000	.00000	.00000
170.00	714.78	19.52	.05619	36.34925	.95518	8.42871	.01912	.00000	.00000	.00000	.00000	.00000
180.00	776.92	29.52	.05525	47.20319	.99142	7.70474	.01931	.00000	.00000	.00000	.00000	.00000
184.91	785.19	31.34	.05566	49.41230	.99961	7.16335	.01936	.00000	.00000	.00000	.00000	.00000
185.72	794.93	33.16	.05603	51.67616	1.00777	6.67553	.01941	.00000	.00000	.00000	.00000	.00000
186.41	804.12	34.95	.05640	53.93731	1.01589	6.19936	.01946	.00000	.00000	.00000	.00000	.00000
187.01	812.82	36.72	.05675	56.19663	1.02396	5.72863	.01951	.00000	.00000	.00000	.00000	.00000
187.58	821.01	38.45	.05711	58.42386	1.03200	5.23914	.01955	.00000	.00000	.00000	.00000	.00000
188.15	828.63	40.13	.05750	60.57227	1.04000	4.69160	.01960	.00000	.00000	.00000	.00000	.00000
188.74	835.64	41.72	.05794	62.55630	1.04793	4.01387	.01965	.00000	.00000	.00000	.00000	.00000
189.34	842.14	43.25	.05841	64.40021	1.05580	3.22467	.01970	.00000	.00000	.00000	.00000	.00000
189.90	848.33	44.74	.05887	66.17990	1.06364	2.38658	.01975	.00000	.00000	.00000	.00000	.00000
190.60	853.54	46.03	.05950	67.53168	1.07139	1.96111	.01980	.00000	.00000	.00000	.00000	.00000
191.48	857.61	47.06	.06034	68.30288	1.07903	-0.47441	.01984	.00000	.00000	.00000	.00000	.00000
195.00	849.01	44.91	.06437	61.90255	1.08578	-8.10384	.01988	.00000	.00000	.00000	.00000	.00000
197.05	845.61	44.08	.06720	58.52039	1.09295	-13.21535	.01992	.00000	.00000	.00000	.00000	.00000
199.04	841.61	43.12	.07028	54.77580	1.10010	-18.62106	.01997	.00000	.00000	.00000	.00000	.00000
201.15	836.20	41.85	.07389	50.13541	1.10715	-24.76101	.02001	.00000	.00000	.00000	.00000	.00000
202.89	832.42	40.98	.07713	46.46241	1.11436	-30.09580	.02005	.00000	.00000	.00000	.00000	.00000
204.63	828.21	40.04	.08061	42.50438	1.12155	-35.66126	.02009	.00000	.00000	.00000	.00000	.00000
206.42	823.42	38.98	.08443	38.15291	1.12872	-41.54577	.02013	.00000	.00000	.00000	.00000	.00000
208.42	816.97	37.59	.08900	32.69817	1.13580	-48.33051	.02018	.00000	.00000	.00000	.00000	.00000
210.38	807.55	35.64	.09499	25.25604	1.14268	-56.73137	.02022	.00000	.00000	.00000	.00000	.00000
212.92	800.63	34.26	.10030	19.43157	1.14981	-63.80002	.02026	.00000	.00000	.00000	.00000	.00000
214.73	795.17	33.20	.10525	14.52778	1.15708	-70.11257	.02030	.00000	.00000	.00000	.00000	.00000
216.87	787.71	31.81	.11136	8.20674	1.16423	-77.57412	.02034	.00000	.00000	.00000	.00000	.00000
219.27	778.74	30.19	.11859	.76372	1.17128	-85.93923	.02039	.00000	.00000	.00000	.00000	.00000
221.73	769.71	28.63	.12633	-6.80674	1.17838	-94.39674	.02043	.00000	.00000	.00000	.00000	.00000
224.07	761.63	27.29	.13402	-13.78698	1.18559	-102.27410	.02047	.00000	.00000	.00000	.00000	.00000
227.00	750.51	25.54	.14411	-23.08885	1.19256	-112.21681	.02051	.00000	.00000	.00000	.00000	.00000
230.88	735.04	23.25	.15807	-35.71984	1.19921	-124.72738	.02056	.00000	.00000	.00000	.00000	.00000
239.34	699.58	18.72	.19077	-60.33552	1.20534	-146.82207	.02060	.00000	.00000	.00000	.00000	.00000
245.00	2020.42	15.96	.21399	-79.23451	1.20273	-161.92594	.02060	.00000	.00000	.00000	.00000	.00000

250.00	1946.72	13.99	.23513	-94.12656	1.20060	-173.79926	.02060	4.27935	63.23549	7.59397	74.88201	.01874
255.00	1879.77	12.38	.25665	-107.56360	1.19860	-184.49478	.02059	3.88090	67.11640	6.81462	81.69663	.01723
260.00	1818.90	11.04	.27837	-119.71199	1.19670	-194.15137	.02059	3.54525	70.66164	6.11134	87.80797	.01592
265.00	1763.48	9.93	.30012	-130.72164	1.19490	-202.89242	.02059	3.26056	73.92220	5.48050	93.28847	.01479
270.00	1712.94	9.00	.32170	-140.72046	1.19318	-210.82252	.02059	3.01733	76.93953	4.91277	98.20124	.01381
275.00	1666.77	8.21	.34295	-149.81958	1.19153	-218.03208	.02059	2.80806	79.74758	4.40150	102.60275	.01296
280.00	1624.53	7.55	.36372	-158.11469	1.18994	-224.59872	.02059	2.62679	82.37437	3.93985	106.54259	.01221
285.00	1585.85	6.98	.38385	-165.68957	1.18842	-230.59023	.02059	2.46879	84.84316	3.52272	110.06531	.01156
290.00	1550.36	6.50	.40322	-172.61667	1.18694	-236.06513	.02059	2.33026	87.17342	3.14464	113.20995	.01098
295.00	1517.78	6.08	.42170	-178.95901	1.18551	-241.07426	.02059	2.20815	89.38157	2.80098	116.01093	.01048
300.00	1487.28	5.72	.43918	-184.83060	1.18412	-245.69757	.02059	2.09998	91.48155	2.52334	118.53426	.01003
305.00	1459.46	5.41	.45556	-190.16793	1.18276	-249.89612	.02059	2.00186	93.48341	2.19668	120.73094	.00963
310.00	1433.88	5.15	.47076	-195.06300	1.18145	-253.74431	.02059	1.91433	95.39834	1.93326	122.66420	.00928
315.00	1410.36	4.92	.48471	-199.55308	1.18016	-257.27183	.02059	1.83704	97.23538	1.69048	124.35469	.00897
320.00	1388.74	4.72	.49734	-203.67062	1.17891	-260.50458	.02059	1.76702	99.00240	1.46573	125.82042	.00869
325.00	1368.88	4.55	.50859	-207.44390	1.17767	-263.46515	.02059	1.70393	100.70633	1.25664	127.07706	.00845
330.00	1350.66	4.40	.51843	-210.89750	1.17647	-266.17321	.02059	1.64696	102.35328	1.06110	128.13816	.00824
335.00	1333.98	4.28	.52681	-214.05278	1.17528	-268.64588	.02059	1.59545	103.94873	.87721	129.01538	.00805
340.00	1318.75	4.18	.53371	-216.92820	1.17412	-270.89793	.02059	1.54886	105.49759	.70320	129.71857	.00789
345.00	1304.89	4.09	.53911	-219.53961	1.17297	-272.94210	.02059	1.50671	107.00430	.53746	130.25603	.00775
350.00	1292.34	4.02	.54298	-221.90051	1.17184	-274.78923	.02059	1.46862	108.47292	.37851	130.63454	.00763
355.00	1281.03	3.97	.54531	-224.02221	1.17073	-276.44844	.02059	1.43428	109.90720	.22493	130.85947	.00753
360.00	1270.94	3.93	.54611	-225.91406	1.16963	-277.92725	.02059	1.40342	111.31061	.07539	130.93485	.00744
360.00	939.75	1.03	1.54171	-286.16306	1.16963	-324.62229	.02059	.00000	111.31061	.00000	130.93485	.00000
540.00	1000.00	1.03	.05461	-9.17099	.03930	-10.53334	.00069	.00000	111.31061	.00000	130.93485	.00000

THETA	T	P	V	H	S	U	N	FLAME SPEED,	TEMP.,	AND EXTENT	MASS-FR.	BURN.-VOL.
167.45	2545.72	17.60	.00002	.00393	.00013	-.06653	.00000	578.08189	2545.71854	.18781	.00010	.00040
169.93	2544.12	18.61	.00023	.03848	.00125	-.06603	.00002	592.18032	2542.46326	.39037	.00100	.00392
172.68	2552.94	19.79	.00109	.20259	.00623	-.32184	.00010	612.50201	2551.40607	.65036	.00500	.01920
174.03	2556.54	20.52	.00211	.41280	.01245	-.63754	.00021	622.45556	2548.51422	.80029	.01000	.03762
175.91	2566.87	21.65	.00402	.87556	.02488	-1.23387	.00041	642.33797	2560.18518	1.04140	.02000	.07266
178.66	2554.11	23.43	.00739	1.59251	.04956	-2.60344	.00083	672.42058	2577.06775	1.35023	.04000	.13520
180.39	2562.29	24.91	.01046	2.49957	.07423	-3.81481	.00124	693.91210	2579.86728	1.56170	.06000	.19160
181.62	2573.85	26.29	.01328	3.55750	.09889	-4.90090	.00165	714.02116	2584.55426	1.71128	.08000	.24273
182.62	2585.38	27.61	.01588	4.73065	.12352	-5.89135	.00207	733.08243	2589.46973	1.82952	.10000	.28929
183.83	2601.61	29.52	.01944	6.67308	.16042	-7.22547	.00269	761.82874	2603.15701	1.97115	.13000	.35187
184.91	2616.06	31.34	.02266	8.78908	.19727	-8.41531	.00331	786.90739	2609.11688	2.08740	.16000	.40721
185.72	2630.62	33.16	.02559	11.13482	.23408	-9.41390	.00393	811.51015	2615.58597	2.17800	.19000	.45665
186.41	2644.23	34.95	.02826	13.65132	.27085	-10.27013	.00455	835.24036	2621.59430	2.25422	.22000	.50110
187.01	2656.97	36.72	.03072	16.32314	.30758	-10.99673	.00517	858.00027	2627.18500	2.29765	.25000	.54133
187.58	2668.65	38.45	.03300	19.11482	.34427	-11.62378	.00580	880.35580	2632.76038	2.36054	.28000	.57796
188.15	2679.03	40.13	.03516	21.98035	.38092	-12.18984	.00642	901.35139	2637.66772	2.42412	.31000	.61148
188.74	2687.51	41.72	.03721	24.83024	.41751	-12.76996	.00704	920.73200	2641.87405	2.48789	.34000	.64227
189.34	2694.59	43.25	.03917	27.66814	.45404	-13.36143	.00766	939.31845	2646.07965	2.55345	.37000	.67068
189.90	2701.03	44.74	.04104	30.54363	.49053	-13.92223	.00828	957.29089	2650.30685	2.61108	.40000	.69703
190.60	2704.42	46.03	.04293	33.15259	.52694	-14.70898	.00891	972.52499	2652.87997	2.68748	.43000	.72150
191.48	2704.17	47.06	.04491	35.34476	.56323	-15.84735	.00953	984.30093	2654.21085	2.78669	.46000	.74431
195.00	2659.09	44.91	.04925	31.55030	.59864	-22.01452	.01014	954.47726	2628.36087	3.08049	.49000	.76514
197.05	2636.03	44.08	.05276	30.24136	.63446	-26.08195	.01075	946.07736	2633.55182	3.44493	.52000	.78514
199.04	2612.60	43.12	.05651	28.58063	.67027	-30.43532	.01137	934.29723	2629.61795	3.70677	.55000	.80407
201.15	2585.51	41.85	.06073	26.08630	.70598	-35.47425	.01198	918.11971	2623.21246	3.94782	.58000	.82194
202.89	2565.82	40.98	.06471	24.39021	.74184	-39.84072	.01260	907.80484	2622.72958	4.14098	.61000	.83898
204.63	2545.78	40.04	.06894	22.39555	.77768	-44.44852	.01321	895.78753	2618.84201	4.31388	.64000	.85515
206.42	2524.77	38.98	.07349	19.99686	.81351	-49.38377	.01383	882.14403	2614.20193	4.48116	.67000	.87053
208.42	2498.87	37.59	.07878	16.53131	.84924	-55.19084	.01444	863.65404	2606.56165	4.65512	.70000	.88514
210.88	2463.43	35.64	.08539	11.14979	.88478	-62.55834	.01506	836.62564	2594.28659	4.84603	.73000	.89901
212.92	2438.21	34.26	.09151	7.18161	.92056	-68.75363	.01567	819.37247	2592.95264	5.00496	.76000	.91233
214.73	2419.00	33.20	.09737	4.00852	.95649	-74.29110	.01629	805.55334	2590.92242	5.13293	.79000	.92509
216.87	2393.34	31.81	.10437	-.57695	.99229	-80.97386	.01690	785.56991	2581.93442	5.26567	.82000	.93723
219.27	2363.00	30.19	.11252	-6.32311	1.02801	-88.59055	.01752	762.24449	2572.42535	5.40526	.85000	.94884
221.73	2333.21	28.63	.12127	-12.28664	1.06376	-96.37194	.01814	739.91488	2565.23666	5.54000	.88000	.95996
224.07	2307.49	27.29	.13009	-17.77362	1.09962	-103.75791	.01875	720.82283	2560.47116	5.66047	.91000	.97062
227.00	2271.38	25.54	.14134	-25.63213	1.13525	-113.05019	.01937	693.17662	2546.62494	5.79524	.94000	.98081
230.88	2220.36	23.25	.15658	-36.91211	1.17055	-125.08233	.01998	655.46177	2526.66293	5.95695	.97000	.99059
239.34	2112.99	18.72	.19077	-60.33551	1.20534	-146.82207	.02060	570.11234	2463.79385	6.24834	1.00000	1.00000

**** ALL INPUT DATA HAVE BEEN PROCESSED.
AT LOC 45267

DATA SET D

FUEL-AIR CYCLE ENGINE CALCULATIONS

IDEALIZED THERMODYNAMIC CYCLE FOR THE INTERNAL COMBUSTION ENGINE

THIS IS A NORMAL RUN EMPLOYING OCTANE, C8H18 AS A FUEL. ADDITIONAL COMMENTS ARE.

CONVERGENCE ERRORS ARE LESS THAN .001 PERCENT

MANIFOLD PRESSURE = 14.54900 PSI EXHAUST PRESSURE = 15.13700 PSI AIR-FUEL RATIO = .92000

MANIFOLD TEMPERATURE = 574.99999 °R COMPRESSION RATIO = 10.00000 H OF COMBUSTION = 1222770.00 CAL/MOLE

CLEARANCE VOLUME = 1.00000 CU.IN. SWEEP VOLUME = 9.00000 CU.IN.

SUMMARY OF RESULTS IN ENGLISH SYSTEM

STATION	T - °R	P - PSI.	H - BTU.	S - BTU./°R	U - BTU.
0	575.000	14.549	-.01598258	.00000000	.00000000
1	625.262	14.549	-.02365288	.00062745	-.03923317
2	1294.103	301.120	.04632732	.00062745	.01408081
3	5234.630	1326.517	.15613554	.00079917	.01408089
4	3125.905	78.430	-.16076147	.00079917	-.24475101
4S	2201.272	15.137	-.27756449	.00079917	-.33670970
5	2201.272	15.137	-.00760723	.00002190	-.00922822

RESULTS

- UNDER THE FOLLOWING ASSUMPTIONS
- 1) ISENTROPIC COMPRESSION AND EXPANSION
- 2) CONSTANT VOLUME COMBUSTION
- 3) NO HEAT TRANSFER

COMPRESSION WORK	=	.53313985E-01 BTU
EXPANSION WORK	=	.25883190E 00 BTU
PUMPING WORK	=	-.56667733E-03 BTU
NET CYCLE WORK	=	.20495123E 00 BTU
INDICATED THERMAL EFFICIENCY	=	.42688828E 02 PERCENT
MEAN EFFECTIVE PRESSURE	=	.212666302E 03 PSI
RESIDUAL WEIGHT FRACTION	=	.27406932E-01 = F
VOLUMETRIC EFFICIENCY	=	.89984529E 00 BASED ON 1 ATM. AND 25°K

**** ALL INPUT DATA HAVE BEEN PROCESSED.
AT LOC 22062

DATA SET E

NORMAL RATES OF FLAME PROPAGATION AND HEAT
TRANSFER - 1200°K EXHAUST TEMPERATURE

A MODEL SIMULATING THE INTERNAL-COMBUSTION ENGINE

ENGINE	FUEL	OPERATING CONDITIONS
FOUR STROKE AUTOMOTIVE TYPE	OCTANE C8H18	AIR-FUEL RATIO .92
BORE 3.94 IN.	ACTIVATION ENERGY 39000.00 CAL./MOLE	PISTON TEMP 500.00 K
STROKE 2.46 IN.	ENTHALPY OF COMBUSTION	WALL TEMP 420.00 K
ROD 7.06 IN.	1222770.00 CAL/MOLE	MAN. TEMP 320.00 K
CLEAR. VOL. 3.33 CU.IN.		MAN. PR. .99 ATM
SWEPT VOL. 30.00 CU.IN.		EXH. PR. 1.03 ATM
COMPRESSION RATIO 10.00 TO ONE		SPARK ADV. 16.00
		SPEED 2800.00 RPM

COMBUSTION OCCURS IN 39 INCREMENTS. ERRORS ARE LESS THAN .00100 PERCENT. FLAME SPEED FACTOR IS .13500000E 06
 HEAT TRANSFER FACTOR IS 1.00000 NORMAL RUN

SUMMARY OF RESULTS

PUMPING WORK	- .47611947E 00 CAL
NET CYCLE WORK	.14762896E 03 CAL
NET HEAT TRANSFERED	.71429917E 02 CAL
INDICATED THERMAL EFFICIENCY	.36641216E 02 PERCENT
MEAN EFFECTIVE PRESSURE	.12402679E 02 ATM
RESIDUAL WEIGHT FRACTION	.27956433E-01
VOLUMETRIC EFFICIENCY	.97409263E 00 PERCENT

IHETA	T	P	V	H	S	U	N	Q	QIOT	M	MIOT	HB
.00	347.66	.99	.54611	-20.02049	.94852	-33.11455	.01895	.00000	.00000	.00000	.00000	.00000
10.00	348.34	1.00	.54302	-19.92079	.94852	-33.04032	.01895	.00000	.00000	.00000	.00000	.00000
20.00	350.39	1.02	.53380	-19.61799	.94852	-32.81476	.01895	.00000	.00000	.00000	.00000	.00000
30.00	353.88	1.06	.51855	-19.10252	.94852	-32.43063	.01895	.00000	.00000	.00000	.00000	.00000
40.00	358.92	1.12	.49750	-18.35633	.94852	-31.87427	.01895	.00000	.00000	.00000	.00000	.00000
50.00	365.67	1.21	.47096	-17.35397	.94852	-31.12629	.01895	.00000	.00000	.00000	.00000	.00000
60.00	374.36	1.32	.43941	-16.06025	.94852	-30.15992	.01895	.00000	.00000	.00000	.00000	.00000
70.00	385.28	1.48	.40348	-14.42873	.94852	-28.93965	.01895	.00000	.00000	.00000	.00000	.00000
80.00	398.80	1.70	.36400	-12.39918	.94852	-27.41926	.01895	.00000	.00000	.00000	.00000	.00000
90.00	415.39	2.01	.32199	-9.89447	.94852	-25.53928	.01895	.00000	.00000	.00000	.00000	.00000
100.00	435.63	2.43	.27867	-6.81701	.94852	-23.22403	.01895	.00000	.00000	.00000	.00000	.00000
110.00	460.22	3.04	.23542	-3.04581	.94852	-20.37899	.01895	.00000	.00000	.00000	.00000	.00000
120.00	489.96	3.93	.19371	1.56231	.94852	-16.89117	.01895	.00000	.00000	.00000	.00000	.00000
130.00	525.65	5.27	.15510	7.15741	.94852	-12.64012	.01895	.00000	.00000	.00000	.00000	.00000
140.00	567.69	7.29	.12106	13.84065	.94852	-7.54011	.01895	.00000	.00000	.00000	.00000	.00000
150.00	615.17	10.29	.09296	21.50783	.94852	-1.66111	.01895	.00000	.00000	.00000	.00000	.00000
160.00	663.85	14.34	.07199	29.49792	.94852	4.49536	.01895	.00000	.00000	.00000	.00000	.00000
164.00	681.60	16.11	.06580	32.44356	.94852	6.77263	.01895	.00000	.00000	.00000	.00000	.00000
166.88	693.05	17.34	.06217	34.35307	.94855	8.24312	.01895	.00000	.00000	.00000	.00000	.00000
168.96	700.84	18.22	.05998	35.66051	.94881	9.18718	.01895	.00000	.00000	.00000	.00000	.00000
171.26	710.11	19.32	.05798	37.23493	.94997	10.10203	.01896	.00000	.00000	.00000	.00000	.00000
172.40	715.91	20.04	.05717	38.23393	.95141	10.49496	.01897	.00000	.00000	.00000	.00000	.00000
173.97	724.93	21.19	.05623	39.82725	.95427	10.96998	.01898	.00000	.00000	.00000	.00000	.00000
176.25	739.18	23.13	.05524	42.32347	.95990	11.38438	.01902	.00000	.00000	.00000	.00000	.00000
177.67	750.62	24.78	.05486	44.44241	.96550	11.51563	.01905	.00000	.00000	.00000	.00000	.00000
178.72	760.74	26.33	.05469	46.41781	.97106	11.54354	.01908	.00000	.00000	.00000	.00000	.00000
179.55	770.00	27.81	.05462	48.31103	.97659	11.51428	.01911	.00000	.00000	.00000	.00000	.00000
180.58	782.72	29.97	.05463	51.05936	.98483	11.41071	.01916	.00000	.00000	.00000	.00000	.00000
181.44	794.37	32.06	.05470	53.73497	.99301	11.26275	.01921	.00000	.00000	.00000	.00000	.00000
182.09	805.21	34.12	.05480	56.37642	1.00113	11.09917	.01926	.00000	.00000	.00000	.00000	.00000
182.64	815.39	36.14	.05491	58.99147	1.00921	10.92484	.01931	.00000	.00000	.00000	.00000	.00000
183.12	825.00	38.15	.05503	61.58592	1.01724	10.74321	.01935	.00000	.00000	.00000	.00000	.00000
183.53	834.12	40.14	.05515	64.17048	1.02523	10.56216	.01940	.00000	.00000	.00000	.00000	.00000
183.97	842.69	42.08	.05529	66.69493	1.03317	10.33903	.01945	.00000	.00000	.00000	.00000	.00000
184.42	850.78	43.99	.05546	69.16143	1.04107	10.07492	.01950	.00000	.00000	.00000	.00000	.00000
184.87	858.44	45.86	.05564	71.58037	1.04893	9.77774	.01955	.00000	.00000	.00000	.00000	.00000
185.33	865.64	47.69	.05585	73.91726	1.05674	9.41789	.01960	.00000	.00000	.00000	.00000	.00000
185.66	872.76	49.55	.05627	78.41421	1.07225	8.56463	.01970	.00000	.00000	.00000	.00000	.00000
186.19	879.08	51.25	.05662	80.29588	1.07991	7.83806	.01975	.00000	.00000	.00000	.00000	.00000
186.81	884.81	52.84	.05662	80.29588	1.07991	7.83806	.01975	.00000	.00000	.00000	.00000	.00000
187.75	889.23	54.09	.05722	81.53481	1.08749	6.57722	.01980	.00000	.00000	.00000	.00000	.00000
189.40	890.79	54.53	.05846	81.25218	1.09489	4.04640	.01985	.00000	.00000	.00000	.00000	.00000
191.05	891.33	54.69	.05992	80.47388	1.10226	1.10371	.01989	.00000	.00000	.00000	.00000	.00000
192.69	890.98	54.59	.06161	79.24133	1.10959	-2.21652	.01994	.00000	.00000	.00000	.00000	.00000
195.61	889.91	54.28	.06350	77.61463	1.11689	-5.86423	.01998	.00000	.00000	.00000	.00000	.00000
197.02	888.42	53.86	.06715	74.78601	1.13157	-12.79984	.02002	.00000	.00000	.00000	.00000	.00000
198.43	886.67	53.36	.06929	72.82623	1.13889	-16.72119	.02011	.00000	.00000	.00000	.00000	.00000
200.14	882.74	52.26	.07212	69.47995	1.14610	-21.79775	.02015	.00000	.00000	.00000	.00000	.00000
201.85	878.43	51.07	.07516	65.88405	1.15332	-27.07970	.02020	.00000	.00000	.00000	.00000	.00000
203.26	875.48	50.28	.07785	63.16618	1.16064	-31.62745	.02024	.00000	.00000	.00000	.00000	.00000
204.88	871.01	49.08	.08113	59.43508	1.16790	-37.01667	.02028	.00000	.00000	.00000	.00000	.00000
206.63	865.50	47.65	.08491	54.99569	1.17514	-42.99195	.02032	.00000	.00000	.00000	.00000	.00000
208.53	858.93	45.99	.08924	49.80880	1.18236	-49.58424	.02036	.00000	.00000	.00000	.00000	.00000
210.54	851.50	44.17	.09413	43.98737	1.18956	-56.69641	.02040	.00000	.00000	.00000	.00000	.00000
213.10	840.62	41.60	.10078	35.72497	1.19663	-65.81870	.02044	.00000	.00000	.00000	.00000	.00000
217.28	820.17	37.17	.11258	21.61096	1.20358	-79.73681	.02047	.00000	.00000	.00000	.00000	.00000
220.00	2442.20	33.93	.12084	10.32239	1.20272	-88.98951	.02046	.00000	.00000	.00000	.00000	.00000

225.00	2353.89	28.79	.13718	-9.32860	1.20120-104.99456	.02045	3.68010	29.95854	12.32495	41.92147	.03005
230.00	2269.63	24.59	.15485	-27.40170	1.19979-119.61203	.02044	3.28215	33.24069	11.33532	53.25679	.02718
235.00	2190.72	21.16	.17366	-43.89931	1.19849-132.88659	.02044	2.94174	36.18243	10.33281	63.58961	.02466
240.00	2117.57	18.36	.19344	-58.92938	1.19728-144.93618	.02044	2.65341	38.83583	9.39619	72.98579	.02247
245.00	2050.30	16.07	.21399	-72.58986	1.19614-155.85922	.02044	2.40984	41.24568	8.51320	81.49899	.02058
250.00	1988.68	14.18	.23513	-84.99645	1.19507-165.76040	.02044	2.20318	43.44985	7.69700	89.19599	.01895
255.00	1932.36	12.62	.25665	-96.26459	1.19404-174.73951	.02044	2.03404	45.47990	6.94907	96.14506	.01753
260.00	1880.90	11.33	.27837	-106.50804	1.19307-182.89198	.02044	1.88197	47.36186	6.27050	102.41556	.01630
265.00	1833.87	10.25	.30012	-115.82815	1.19213-190.30177	.02044	1.75536	49.11722	5.65443	108.07000	.01524
270.00	1790.89	9.33	.32170	-124.31605	1.19123-197.04367	.02044	1.64648	50.76370	5.09542	113.16541	.01431
275.00	1751.57	8.56	.34295	-132.05322	1.19036-203.18417	.02044	1.55233	52.31604	4.58816	117.75357	.01350
280.00	1715.60	7.91	.36372	-139.11178	1.18952-208.78187	.02044	1.47047	53.78650	4.12724	121.88081	.01278
285.00	1682.68	7.35	.38385	-145.55536	1.18870-213.88834	.02044	1.39891	55.18541	3.70756	125.58837	.01216
290.00	1652.54	6.87	.40322	-151.43996	1.18791-218.54886	.02044	1.33607	56.52148	3.32445	128.91282	.01161
295.00	1624.95	6.46	.42170	-156.81517	1.18713-222.80345	.02044	1.28064	57.80211	2.97395	131.88677	.01112
300.00	1599.57	6.11	.43918	-161.74839	1.18637-226.70608	.02044	1.23155	59.03366	2.67108	134.55785	.01070
305.00	1576.35	5.80	.45556	-166.25261	1.18562-230.26752	.02044	1.18772	60.22138	2.37373	136.93158	.01032
310.00	1555.14	5.54	.47076	-170.36157	1.18487-233.51493	.02044	1.14866	61.37004	2.09875	139.03032	.00998
315.00	1535.78	5.31	.48471	-174.10493	1.18415-236.47212	.02044	1.11377	62.48381	1.84343	140.87375	.00969
320.00	1518.15	5.12	.49734	-177.50870	1.18343-239.15999	.02044	1.08255	63.56636	1.60531	142.47906	.00943
325.00	1502.14	4.95	.50859	-180.59560	1.18272-241.59674	.02044	1.05460	64.62095	1.38216	143.86122	.00920
330.00	1487.40	4.81	.51843	-183.38361	1.18205-243.78609	.02044	1.02956	65.65051	1.15979	145.02101	.00900
335.00	1474.26	4.69	.52681	-185.88618	1.18137-245.75503	.02044	1.00673	66.65723	.96222	145.98323	.00883
340.00	1462.49	4.59	.53371	-188.12457	1.18070-247.51564	.02044	.98655	67.64378	.77406	146.75728	.00868
345.00	1452.04	4.52	.53911	-190.11131	1.18004-249.07790	.02044	.96858	68.61236	.59368	147.35097	.00856
350.00	1442.85	4.46	.54298	-191.85675	1.17938-250.45008	.02044	.95263	69.56499	.41955	147.77052	.00845
355.00	1434.88	4.41	.54531	-193.36929	1.17873-251.63891	.02044	.93859	70.50358	.25024	148.02075	.00837
360.00	1428.10	4.38	.54611	-194.65540	1.17808-252.64957	.02044	.92633	71.42992	.08432	148.10508	.00831
360.00	1037.60	1.03	1.68912	-266.69466	1.17808-308.83118	.02044	.00000	71.42992	.00000	148.10508	.00000
540.00	1200.00	1.03	.05461	-6.63366	.03367 -7.99601	.00057	.00000	71.42992	.00000	148.10508	.00000

THETA	T	P	V	H	S	U	N	FLAME SPEED, TEMP., AND EXTENT MASS-FR.	BURN.-VOL.		
166.88	2550.22	17.34	.00002	.00403	.00012	-.00637	.00000	693.568172550.22064	.18768	.00010	.00040
168.96	2550.99	18.22	.00024	.04020	.00124	-.06382	.00002	709.041582548.47394	.39182	.00100	.00393
171.26	2559.60	19.32	.00112	.21093	.00619	-.31093	.00010	732.479902557.99228	.65396	.00500	.01923
172.40	2563.78	20.04	.00215	.43096	.01237	-.61446	.00021	744.816542556.04697	.80083	.01000	.03769
173.97	2574.94	21.19	.00409	.91613	.02472	-1.18414	.00041	769.489082568.44778	1.04218	.02000	.07278
176.25	2582.65	23.13	.00752	1.89394	.04932	-2.31913	.00082	809.479002588.05792	1.35103	.04000	.13617
177.67	2595.70	24.78	.01058	3.02941	.07389	-3.32307	.00123	838.833632591.69345	1.56457	.06000	.19293
178.72	2608.96	26.33	.01335	4.30094	.09842	-4.21394	.00164	866.328712597.07632	1.70534	.08000	.24416
179.55	2621.42	27.81	.01588	5.68592	.12292	-5.01054	.00205	892.046102602.44113	1.83185	.10000	.29069
180.58	2638.80	29.97	.01929	7.95764	.15961	-6.04375	.00267	930.853232617.62723	1.97164	.13000	.35314
181.44	2654.74	32.06	.02233	10.43696	.19625	-6.90396	.00329	965.657972624.73230	2.08377	.16000	.40829
182.09	2669.59	34.12	.02507	13.10873	.23283	-7.60389	.00390	998.632842631.40555	2.17138	.19000	.45746
182.64	2683.42	36.14	.02755	15.95217	.26936	-8.16064	.00452	1030.682372637.98550	2.24425	.22000	.50165
183.12	2696.34	38.15	.02981	18.95150	.30585	-8.58738	.00514	1061.729062644.32162	2.28758	.25000	.54165
183.53	2708.52	40.14	.03188	22.09615	.34229	-8.89111	.00576	1091.958162650.46838	2.34305	.28000	.57807
183.97	2719.59	42.08	.03381	25.34316	.37869	-9.11328	.00638	1120.958542656.11258	2.39998	.31000	.61141
184.42	2729.72	43.99	.03561	28.67371	.41504	-9.26440	.00699	1148.911542661.50839	2.45737	.34000	.64208
184.87	2739.10	45.86	.03730	32.08812	.45136	-9.34580	.00761	1175.885482666.66266	2.51507	.37000	.67042
185.33	2747.55	47.69	.03891	35.54608	.48763	-9.39170	.00823	1201.750432671.48312	2.57337	.40000	.69672
185.66	2756.09	49.55	.04039	39.14792	.52387	-9.31723	.00885	1227.911612676.80453	2.62238	.43000	.72119
186.19	2762.46	51.25	.04187	42.60189	.56005	-9.36893	.00947	1251.541082680.60849	2.68828	.46000	.74404
186.81	2767.41	52.84	.04334	45.95755	.59617	-9.50482	.01009	1273.103962684.13425	2.77031	.49000	.76544
187.75	2768.66	54.09	.04495	48.84111	.63220	-10.04084	.01070	1289.488682685.50375	3.02652	.52000	.78553
189.40	2761.78	54.53	.04703	50.47759	.66806	-11.63083	.01132	1294.783892682.56412	3.24648	.55000	.80445
191.05	2753.06	54.69	.04928	51.71059	.70388	-13.56013	.01193	1296.586872681.49118	3.49778	.58000	.82235
192.69	2742.83	54.59	.05171	52.55651	.73966	-15.81409	.01254	1294.967762679.91635	3.74590	.61000	.83933
194.31	2731.45	54.28	.05432	53.05078	.77543	-18.36207	.01316	1290.568652678.01178	3.98798	.64000	.85545
195.61	2723.56	54.22	.05675	54.01397	.81124	-20.51358	.01377	1290.100052679.15851	4.17768	.67000	.87079
197.02	2713.21	53.86	.05945	54.39514	.84701	-23.15252	.01438	1284.752732676.54221	4.35503	.70000	.88539
198.43	2702.49	53.36	.06231	54.55804	.88279	-25.97280	.01500	1277.838392674.49655	4.53166	.73000	.89930
200.14	2686.03	52.26	.06581	53.40828	.91846	-29.89023	.01561	1262.436292668.42404	4.75110	.76000	.91258
201.85	2669.28	51.07	.06954	51.98136	.95413	-34.03473	.01622	1245.893012664.48178	4.94686	.79000	.92526
203.26	2657.25	50.28	.07298	51.34299	.98991	-37.51546	.01683	1235.185642664.18185	5.10144	.82000	.93738
204.88	2641.38	49.08	.07700	49.70066	1.02563	-41.83038	.01744	1218.002532658.44345	5.25575	.85000	.94898
206.63	2623.12	47.65	.08152	47.32448	1.06132	-46.75149	.01805	1197.309042652.60455	5.41455	.88000	.96007
208.53	2602.39	45.99	.08663	44.15932	1.09699	-52.32226	.01866	1172.750982645.73459	5.57636	.91000	.97071
210.54	2579.70	44.17	.09233	40.29914	1.13265	-58.46044	.01926	1145.804632638.61850	5.73687	.94000	.98089
213.10	2547.47	41.60	.09984	33.93795	1.16818	-66.65591	.01987	1106.353962624.86008	5.92082	.97000	.99065
217.28	2491.14	37.17	.11258	21.61097	1.20358	-79.73681	.02047	1033.459382594.17960	6.18577	1.00000	1.00000

**** ALL INPUT DATA HAVE BEEN PROCESSED.
 AT LOC 42742

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