OPTIMAL DESIGN OF ORGANIC REFRIGERANT REVERSE CYCLE Absorption Air conditioning systems.

A thesis submitted in fulfilment of the requirement for a Master of Engineering Science Degree at the Chemical Engineering Department of the University of Adelaide.

> R.L. O'Loughlin June 1986

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ABSTRACT

Currently there is a small market in absorption refrigeration equipment which is dominated by Lithium Bromide-Water and Water-Ammonia systems. To overcome some of the problems associated this equipment an organic fluid cycle has been proposed.

A computer model for a single stage absorption cycle has been written which provides for optional selection of internal heat exchangers, a choice of working fluid pairs and the selection of system operating parameters. Fluid properties are calculated using the Peng-Robinson cubic equation of state. Tables of pure and mixture fluid properties can be generated as well as system analyses from both the First and Second Laws of thermodynamics.

In the working range of the cycle fluid properties, other than liquid entropy, were generally within 5 percent of published data. The second law analysis revealed that absorber mixing and the distillation column availability losses (irreversibilities) were large and unavoidable.

Coefficient of Performance (COP) was used to assess system performance and 0.35 can be taken as a typical COP value for a cooling cycle under Adelaide conditions. When compared with a commercially available Lithium Bromide system the organic cycle COP was 15 percent lower. In the heat pump mode heating COP's of 1.3 and above were predicted.

These results are considered encouraging enough to warrant investing the additional time required to exhaustively investigate the cycle and optimise its design.

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STATEMENT OF ORIGINALITY

This thesis contains no material which has been accepted for the award of any other degree or diploma in any University and to the best of the author's knowledge and belief contains no material previously published or written by another person, except where due reference is made in the text. The author consents to this thesis being made available for photocopying and loan.

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INTRODUCTION

Currently two types of equipment dominate the established absorption refrigeration technology. For evaporator applications below 0 deg C water ammonia systems are used. Usually, for domestic systems, heat is rejected through a refrigerant to air heat exchanger. Forty years ago in Australia these systems were common as household refrigerators whereas nowadays their application is largely confined to the leisure industry eg gas/electric powered caravan refrigerator where portability and flexibility of energy supply is important. The Lithium Bromide - Water system is used in air conditioning applications where evaporator temperatures are usually in the range 3 deg C to 8 deg C. However, here heat is usually rejected via a water cooled condenser and absorber. Chilled water is produced in the evaporator to effect the air conditioning function and cooling water is required to remove the rejected heat. These requirements generally confine the use of lithium bromide - water equipment to larger commercial systems. Examples include the use of exhaust and cooling water heat from gas and diesel generating sets and the use of off peak or off season steam. A few solar powered systems have been installed (5).

When assessing the viability of this equipment compared with mechanical vapour compression machines it is necessary to account for all costs over a lifespan and not just costs based on coefficient of performance (COP).

For cooling the coefficient of performance (COP) is defined as

COP =

Useful cooling effect = ------Heat input

However, as this report is concerned with fundamental technical aspects of the cycle COP has been chosen as the prime measure for comparison rather than life cycle cost. A typical lithium - bromide based air conditioning system COP is 0.5 (22) whilst less than this can be expected of an ammonia water system operating with a low temperature evaporator eg -10 deg C. Other factors which serve to limit the application of this type of equipment are, in the case of lithium - bromide the possibility of solidification and, for ammonia - water, toxicity, flammability, corrosion and high manufacturing costs.

As early as 1949 (B) researchers were investigating the characteristics of alternative binary mixtures in an effort to improve system performance but despite this the two abovementioned systems remain dominant. During the late 1970's Dr J. R. Roach of the Chemical Engineering Department of the University of Adelaide (17) began experimenting with organic based binary mixture systems because they offered low toxicity, freedom from the crystallization problems of lithium bromide and the possibility of sub-zero operation either for low temperature refrigeration or for use in a heat-pump mode. In addition they can use the existing well established materials and methods used for conventional mechanical vapour compression systems.

Early experimentation demonstrated that these organic systems could be used in an absorption cycle and this project was founded with a view to detailing cycle performance. The aims of the project have been grouped under three broad headings

Fluid property estimation

- a) To represent in a computer thermodynamic and other relevant properties for combinations of organic fluids considered as candidate working fluids for the absorption cycle.
- b) To produce enthalpy concentration diagrams for these fluids.

Absorption system analysis

- a) To create a computer model of the system with the ability to nominate a range of operating parameters, operating fluids, and combinations of internal heat exchangers.
- b) To provide a first law energy balances for the system and to determine performance data such as power input and COP.
- c) To provide second law analyses which illustrate the energy efficiency of the various components.

Optimum design

To establish a system configuration and associated mixture which approaches optimum performance given the

normal constraints applying to such equipment. Three methods were considered for the generation of fluid property data namely, tables, specific empirical equations and a generalized equation approach. The Peng-Robinson cubic equation of state was finally selected because it offered acceptable accuracy and manageable programming for both pure substance and mixture properties. Most pure and mixture

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viz:

property values calculated for vapour were within 5 percent of published data but liquid property values, in particular liquid entropy, sometimes exhibited errors around 10 percent. Both enthalpy and entropy data were generated so that the thermodynamic property of availability (exergy) could be calculated for use in a 'second law analysis'.

The absorption refrigeration cycle computer model was developed and the property data utilized was derived from the Peng-Robinson equation referred to above. Traditionally analysis of such systems has been conducted using the first law of thermodynamics although two references were found which used the second law for a single set of operating conditions (21,7). It was decided that a second law analysis should be a part of the model in an effort to establish which components or aspects of the system were the most wasteful of energy. This was achieved and in addition to being able to select working fluids, heat exchangers and system operating parameters, it is possible to calculate the availability losses through the various cycle elements.

Experimental data from an early model prototype test was used to check the program and, for 1.5kW of cooling, the computed power input to the reboiler was 2.53kW while the measured value was 2.4kW. The resulting COP values of 0.58 (predicted) and 0.61 (measured) respectively were considered an encouraging result.

Using equations developed by Teja and Roach (35) and the first law concept of COP it is possible to derive the expression

$$COP = COP_{CARNOT} - \frac{\frac{D}{Q_G}}{\frac{T_O}{T_E} - 1}$$

where D = Availability loss (or irreversibility)

 Q_G = Absorption system input

 T_O = Temperature of the environment

 T_{E} = Temperature of the evaporator

Using this equation it is possible to calculate the system COP by subtracting the sum of the irreversibility components from the Carnot COP. The smaller the irreversiblity values are, the closer the actual COP is to the Carnot value. Taking this to the limit, if D is zero, then the operating COP equals the Carnot COP. Of course in practice there must always be some availability loss and therefore the COP can never equal the Carnot COP. By separating this availability loss into avoidable and unavoidable components an assessment of the potential for COP improvement is possible. For example, the availability loss which occurs in a heat exchanger due to the temperature difference between the streams is, in theory at least, avoidable by virtue of heat exchanger design whereas the availability loss through mixing of streams is not. For all the systems tested the sum of the distillation column irreversibilities and the irreversibility of mixing in the absorber was always dominant, even though adding internal heat exchangers produced some reductions overall these two losses remained dominant. An extensive investigation of the topology of all possible systems including multistage evaporatorabsorber combinations and multiple column combinations is considered to be beyond the scope of this thesis.

Aside from assessing the importance of system variables such as heat exchanger approaches the computer model offers the possibility of investigating the cycle performance with a large number of working fluid pairs. Only two mixtures have been considered here :-

Dichlorodifluoromethane (R12) -

Trichlorotrifluoroethane (R113) and Chlorodifluoromethane (R22) -

Trichlorofluoromethane (R11). Their performances were found to be significantly different. This suggests that an examination of other pairs could lead to system improvement. The R12-R113 combination consistently produced higher COP values for the same ambient and evaporator conditions because it has a higher Carnot COP capability. Despite this higher value the R12-R113 system cooling COP was still 15% lower than a commercially available lithium bromide-water air conditioning chiller for a typical single stage system.

Both cooling and heat pump systems have been considered and, for the heating mode, if all the heat rejected from the system is used for heating the COP can be significantly greater than one (about 1.3).

It should be noted that the boiler heat input values used throughout this report are nett values and do not include a heating source efficiency (eg a gas combustion efficiency).

It is not possible to optimize the overall system design based on COP alone. Whether or not these absorption systems are economically viable can only be determined by an appraisal which takes into account such factors as fuel costs,

manufacturing costs and the COP variation due to changing load requirements. Ambient temperature changes significantly affect the COP and therefore the importance of temperature variations should not be under-estimated. For an evaporator temperature of approximately 7 deg C the COP for cooling varied from approximately 0.3 for an air cooled condenser operating at a condensing temperature of 48 deg C (ambient air temperature of 38 deg C) to above 0.6 for a system using condenser cooling water at 30 deg C. To make a meaningful assessment of the situation it is proposed that further work include driving the modelling program by some optimization program which takes account of these seasonal variations as well as considering economic criteria.

Even though this work has not conclusively shown the viability or otherwise of the organic absorption system it has demonstrated a method of system analysis and that prototype modelling by computer is feasible. Further work is recommended and it is suggested that it be divided into four broad areas:-

- 1) Investigation of an extended range of binary mixtures.
- Simulation of prototype performance.
- 3) Investigation of alternative system configurations including multistage evaporator/absorber combinations and multiple columns.

Optimization of design by assessing costs and performance.

For the mixtures tested the COP was lower than for existing Lithium Bromide - water systems and therefore, on a COP only basis it is considered unlikely that these would be able to capture the existing cooling market. But it is possible that a market exists, particularly for heating, for systems which can

use low grade heat with refrigerant to air heat exchangers as well as offering low toxicity, subzero operation and construction from components already extensively used in mechanical vapour compression equipment.

CHAPTER 1

Nomenclature

 $\mathbf{A} = \mathbf{Peng}$ Robinson constant

a = Peng Robinson constant

B = Peng Robinson constant

b = Peng Robinson constant

c = constants

 C_p = Isobaric specific heat

f = fugacity

 $\mathbf{H} = \mathbf{enthalpy}$

h = enthalpy

n = number of unknowns

 $\mathbf{P} = \mathbf{pressure}$

R = universal gas constant

S = entropy

T = temperature

 $\mathbf{v} = \mathbf{specific volume}$

 $\mathbf{x} =$ concentration

 $\mathbf{Z} = \text{compressiblity}$

 $\kappa = Peng Robinson constant$

 $\delta = Peng$ Robinson interaction coefficient

 $\omega = Acentric factor$

Subscripts

c = critical

 $\mathbf{R} = \mathbf{reduced}$

i = component

j = component

 $\frac{\text{Superscripts}}{\text{o} = \text{ideal gas state}}$

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FLUID PROPERTIES

1.1 Property calculation methods

Three methods were considered for the generation of the following pure and mixture properties:

Enthalpy Entropy Density Fugacity Availability (exergy) Compressibility

Method 1: Published data and tables

The use of interpolation or curve fitting on published thermodynamic tables was dismissed through lack of comprehensive data. For pure refrigerants some 35 tables were found (3) but only 2 references were found for nonazeotropic binary mixtures. One for refrigerants 13B1& 152a (10) and the other for Ammonia & water (2). A search for equations used to generate data yielded a similar result. A family of equations by Downing (14) included pure refrigerants 11, 12, 13, 14, 21, 22, 23, 113, 114, 500, 502 and C318 but no suitable equations were found for mixtures.

Method 2: Hybrid of published pure substance data and the Van Laar equations for mixture vapour-liquid equilibrium.

This evolved from method 1 in that it was thought that the lack of mixture data could be overcome by using some generalized approach. Using the published pure substance data and the activity coefficient approach discussed by Null (26) vapour-liquid equilibrium data was found. This method was pursued (Appendix B) and some enthalpy concentration diagrams for R12-R113 were produced using the Van Laar equations for liquid phase activity coefficients and the Peng-Robinson equation of state for the other thermodynamic properties based on P-V-T data. This method was abandoned because data was restricted to substances with detailed thermodynamic tables (properties at low pressures were used in Van Laar equations) and the program required inconveniently large input files.

Method 3: Generalized Equation of State

By using an equation of state for all properties including vapour-liquid equilibrium a consistent approach could be obtained for a large range of substances with minimal input Reid et al (31) claimed good accuracy from the data. Benedict-Webb-Rubin equation, however, its use was rejected on the basis of lack of data for some refrigerants and its complexity for programming. Cubic equations of state were considered next because it had been demonstrated by Asselineau et al (4) that the Soave Redlich Kwong (SRK) equation gave acceptable engineering data for refrigerants. This approach was reinforced in an article (11) on cubic equations of state in which it was concluded that " A practical balance must be struck between accuracy on the one hand, and simplicity and generalizability on the other. The cubic equations, despite their short comings, meet this

requirement ".

It was decided to try the newer Peng-Robinson equation (28) rather than the SRK because the authors claimed that " it (the equation) performs as well as or better than the SRK equation in all cases tested and shows its greatest advantages in the prediction of liquid phase densities ". At the time, 1981, no publications could be found on PR generated data for organic refrigerants thus making it an opportunity to test the Peng Robinson claim by checking PR results against the Asselineaus SRK data. It is interesting to note that other researchers had similar ideas. Meskel-Lesavre et al (24) published (before this work was complete) PR data and described the equations performance as "fair". Also Otakake et al (27) claim that of the SRK, Redlich Kwong Soave Fuller, Benidict Webb Rubin Nishiumi, The Combination and the PR equations "the PR is the most suitable for superheated vapours as well as pressurized liquids".

Conclusion

Of the 3 methods the equation of state approach used the least and most general input data, as well as offering an acceptable accuracy overall. Appendix F shows final tables of calculated pure substance data while appendix E shows comparisons with published values. The accuracy is considered to be acceptable and as good as that which Asselineau et al gained from the SRK equation. (See Appendix D for a comparison)

1.2 Pure substance:

Dichlorodifluoromethane (R12) was used to prove the pure substance calculations.

The Peng Robinson equation of state is

$$P = \frac{RT}{v - b(T)} - \frac{a(T)}{v(v + b(T)) + b(T)(v - b(T))}$$
(1.1)

This basic equation (28) expressed in terms of compressibility factor is

$$Z^{3} - (1 - B)Z^{2} + (A - 3B^{2} - 2B)Z - (AB - B^{2} - B^{3}) = 0$$
 (1.2)

where

$$A = \frac{a(T)P}{(RT)^2} \tag{1.3}$$

$$B = \frac{b(T)P}{RT} \tag{1.4}$$

$$Z = \frac{Pv}{RT} \tag{1.5}$$

$$a(T) = a(T_c)\alpha \tag{1.6}$$

$$\alpha = 1 + \kappa \left(1 - \sqrt{T_r} \right)^{\sigma} \tag{1.7}$$

$$b(T) = b(T_c)$$

(1.8)

In equation 1.7 K is a constant for each substance and may be determined from

$$\kappa = 0.37464 + 1.54226\omega - 0.26992\omega^2 \tag{1.7}$$

where $\boldsymbol{\omega}$ is the acentric factor.

This correlation, by Peng & Robinson, between K and acentric factor was determined by establishing vapour-liquid equilibrium using $f_L = \tilde{f}_V$. Initially a range of K was determined for each substance by comparing experimental and calculated saturated pressure values for a given temperature. Feng & Robinson then used equation 1.7 to linearize the relationship. This in turn established K. Finally, to generalize the application of the equation, K was correlated to the acentric factor which yielded equation 1.9 .

Initially equation 1.9 was used to generate saturated data, however, it was reasoned that by using the Peng -Robinson approach and minimising the error between calculated and selected experimental data a value of K could be generated for each substance rather than rely on the acentric factor correlation. (Note Meskel-Leasavre (24) did this too) This was done and equation 1.11 was discarded. Figure 1.1 illustrates how the prediction of saturated data is affected by K. Convergence for the selection of K was based on pressure by minimizing

SD1 =
$$\sqrt{\frac{\sum_{1}^{n} (calcd - publd)^{2}}{(n-1)}}$$
 (1.10)

for a range of temperatures between the higher of -40 deg Cor the freezing point and the lower of T critical or 200 deg C. See Appendix A for detail.

The a(Tc) and b(Tc) values of equations 1.6 and 1.8 are generated by substituting critical conditions in equation 1.1 thus

$$a(T_c) = 0.457252 \frac{R^2 T_c^2}{P_c} \tag{1.11}$$

$$b(T_c) = 0.0778 \frac{RT_c}{P_c}$$
(1.12)
$$Z_c = 0.3074$$
(1.13)



SATURATION PRESSURE VS TEMP FOR R12

TEMPERATURE DEG C Fig 1.1

These critical constants may be checked by equating the second derivative of equation 1.2 (corresponding to the point of inflextion at the critical condition) to zero. From equation 1.2 B=1-3Z (1.14) Substituting Zc = 0.3074 in equation 1.14 yields

$$B = 0.0778$$

Rearranging 1.2

$$A = \frac{-Z^{3} + (1 - B)Z^{2} + Z(3B^{2} + 2B) - (B^{2} + B^{3})}{(Z - B)}$$

Substituting for B and Zc yields

It is worth noting that the set of constants given in the original PR publication of $Z_{\rm C} \approx 0.307$, $B \approx 0.0778$ and A = 0.45724 did not satisfy the above equations and that later publications (11) used the same set as was derived here.

One of the features of cubic equations of state is their fixed critical condition. This PR Z value of 0.3074 seldom coincides with the actual value (viz R12 = 0.276) and therefore must be in error. No attempt was made to substitute the correct value because the small latent heat value near the critical region made refrigeration system operation in this area undesirable. It was essential though that the algorithm operate up to and through this critical region. Occasionally a "guess" generated in an iteration was quite close to the critical point, well away from the system operating condition. This sometimes led to computational difficulties. When the cubic yielded 3 roots but only 1 was real, program execution terminated. This

failure to get both a liquid and a vapour root was overcome by the definition of a "dead zone" around the critical point. Inside this zone, which was specified in terms of Celsius degrees, the critical properties were used. The largest zone width was 0.7 Celsius degrees.

Figure 1.2 illustrates the nature of the equation. A wide flat zone develops as the cubic approaches the point of inflexion. The following points 1.2.1 to 1.2.6 outline the basis used for the calculation of each property.

1.2.1 Saturation pressure

For a given temperature the saturation pressure was determined by the convergence of the vapour and liquid fugacity values.

1.2.2 Compressiblity

Knowing P and T compressibility is evaluated directly from equation 1.5 using the specific volume value from the PR equation.

1.2.3 Fugacity

Appendix A shows the derivation of

$$ln\frac{f}{P} = Z - 1 - ln(Z - B) - \frac{A}{2\sqrt{2}B}ln\left(\frac{Z + 2.414B}{Z - 0.414B}\right) \qquad (1.15)$$

By substituting known values of P and T and a calculated value of Z into equation 1.15 the fugacity can be found.



PENG ROBINSON CUBIC FOR SATURATED R12

z value Fig 1.2

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1.2.4 Enthalpy

Reid et al (31) propose a 3 stage calculation for enthalpy Stage 1

Evaluate enthalpy departure at T_{datum} (equation 1.16) from saturated liquid to ideal gas state.

$$H - H^{o} = RT(Z - 1) + \frac{T\frac{da(T)}{dT} - a(T)}{2\sqrt{2}b(T)}ln\left(\frac{Z + 2.414B}{Z - 0.414B}\right)$$
(1.16)

(See appendix A for all departure derivations)

Stage 2

Integrate, with respect to temperature, the specific heat capacity expression for the ideal gas state. A cubic specific heat capacity equation (equn 1.17) was taken from Reid (31) with constants from the same reference as well as some derived from data in ASHRAE (2).

$$C_{p}^{o} = c4 + c3T + c2T^{2} + c1T^{3}$$

$$\int_{T}^{T_{ref}} C_{p}^{o} dT = \left[\frac{c1T^{4}}{4} + \frac{c2T^{3}}{3} + \frac{c3T^{2}}{2} + c4T\right]_{T_{ref}}^{T}$$
(1.18)

Stage 3

At $T = T_{system}$ evaluate the enthalpy departure from the ideal gas state up to the system pressure. Graphically on a P - T diagram these 3 stages may be represented thus;



1.2.5 Entropy

As with enthalpy a 3 stage process with saturated liquid as the datum was used. The departure was given by

$$S - S^{\circ} = \frac{\frac{da}{dT}}{2\sqrt{2}b(T)} \ln \frac{Z + 2.414 B}{Z - 0.414 B} - R \ln \frac{Z}{Z - B} + R \ln \frac{V}{V^{\circ}}$$
(1.19)

and by the specific heat equation integration for the ideal gas state

$$\Delta S = \int_{T_{ref}}^{T} \frac{C_{p}^{o}}{T} dT \qquad (1.20)$$

$$\Delta S = \left[\frac{c1 T^{3}}{3} + \frac{c2 T^{2}}{2} + c3 T^{2} + c4 \ln T\right]_{T_{ref}}^{T}$$

1.2.6 Latent Heat

For a given temperature the latent heat was taken as the difference between the liquid and vapour isothermal enthalpy departure from the ideal gas state to the pressure under consideration as computed using the PR equation.

1.3 Binary Mixtures

Like most cubic equations of state the PR equation copes

with mixtures via mixing rules and a coefficient which is peculiar to a pair of substances. Mixture properties are sensitive to the value of the coefficient and therefore care should be taken in its evaluation. Values for the Peng-Robinson constants a(T) and b(T) were taken directly from the original publication (28). Applying these rules to a binary mixture yields

$$a_{mis} = x_1^2 a(T)_1 + 2 x_1 x_2 a(T)_{12} + x_2^2 a(T)_2$$

$$b_{mis} = x_1 b(T)_1 + x_2 b(T)_2$$
(1.21)

where

SD2

$$a(T)_{12} = (1 - \delta_{12}) \sqrt{a(T)_1 a(T)_2}$$
(1.23)

(1.22)

For the mixtures listed below the binary interaction $coefficient \delta$ was determined by minimizing the error between calculated and published data

mixture			e	reference
	R1381	-	R152a	(10)
	R11	-	R22	(24)
	R12	-	R152a	(27)
	R12	-	R13	(25)
	R12	-	R22	(16)

No experimental data could be found for an R12 R113 mixture and so the regular solution data shown in Appendix B was used. For all mixtures the saturation pressure data generated from a given temperature and liquid concentration was compared to published data using

$$= \sqrt{\frac{\sum_{1}^{n} \left(\frac{calcd - publd}{publd}\right)^{2}}{(n-1)}}$$

Table 1.1 shows the S values whilst appendix E shows all data.

Mixture		SD2	No Pts	8
R13B1	R152a	0.014	19	0.079
R11	R22	0.016	22	0.0495
R12	R152a	0.011	18	0.081
R12	R13	0.033	10	0.011
R12	R22	0.013	22	0.047
R12	R113	0.069	14	0.04525

Table 1.1

(Note: SD2 is based on saturation pressure)

For saturation pressure, compressibility, enthalpy, entropy and latent heat calculations the pure property equations were used but with mixture values of Z. In the case of the specific heat capacity integration for enthalpy (and entropy) a simple molar proportion of the pure values was used.

$$\int_{T_{ref}}^{T} (C_{p}^{o})_{mis} dt = x_{1} \int_{T_{ref}}^{T} (C_{p}^{o})_{1} dt + x_{2} \int_{T_{ref}}^{T} (C_{p}^{o})_{2} dt$$

The fugacity coefficient was evaluated after configuring the published equation for a binary mixture viz;

$$ln\frac{f_{1}}{x_{1}P} = \frac{b(T)_{1}(Z-1)}{b(T)} - ln(Z-B) - \frac{A}{2\sqrt{2}B}$$

$$\left(\frac{2(x_{1}a(T)_{1} + x_{2}a(T)_{12})}{a(T)} - \frac{b(T)_{1}}{b(T)}\right) ln\left(\frac{Z+2.414B}{Z-0.414B}\right)$$
(1.24)

Note: In equation 1.24 properties without a subscript are mixture properties calculated from the main PR equation but with the pure a and b constant values replaced with the mixture values determined from the mixing rules (equations 1.21 and 1.22).

1.4 Subcooled liquid properties

For subcooled liquid conditions the PR equation sometimes yields a meaningless negative root instead of a valid liquid value. To overcome this discrepancy in the calculation all subcooled liquid calcuations were treated as saturated condition calculations at the temperature in question. The error introduced by this was considered insignificant. For example the error in enthalpy between R12 liquid at saturation pressure and 60 deg C and liquid at the same pressure and 40 deg C is less than one percent.

1.5 Superheated Vapours

A vapour root was found from the PR cubic for all superheated values tested. No detailed check on superheated vapour points was conducted, however, all pure refrigerant tables in Appendix F show these data and spot checks on enthalpy revealed a typical accuracy of around 3 percent. For example Dupont data gives 231 kJ/kg for R12 at 959 kPa and 30 degrees of superheat and the calculated value was 224.6 kJ/kg.

CHAPTER 2

Nomenclature

a = activity B = availability D = irreversibility F = flow f = fugacity H = enthalpy Q = energy n = mols S = entropy

Subscript

A = absorber

- B = bottom
- C = condenser
- D = distillate
- $\mathbf{E} = \mathbf{evaporator}$
- $\mathbf{F} = \mathbf{feed}$
- G = generator (boiler)
- i = component
- i = internal
- L = liquid
- o = datum
- p = pump
- q = energy across boundary
- R = reflux
- V = vapour

y = relates to internal subsystem

z = relates to external system

 $\frac{\text{Superscript}}{-= \text{partial molar property}}$

Availability, the Second Law and the Absorption cycle.

2.1 The second law and the closed system

By using the approach to second law system analysis developed by Teja and Roach (35) it is possible to write an availability balance equation for an absorption cycle where only heat and work cross the system boundary.

Consider the absorption refrigeration cycle shown in figure 2.1;

- Heat input to the boiler ($Q_{\mathbf{G}}$) powers the distillation column.
- Heat is rejected from the condenser (Ω_c) to the surroundings.
- After throttling to a lower pressure heat (Q_E) is transferred from the medium being cooled to the boiling low temperature refrigerant.
- Heat is rejected from the absorber (Q_A) to condense the two phase mixture formed when the refrigerant vapour from the evaporator is mixed with the bottom product from the distillation column.
- A work input (Wp) is required to pump the feed flow into the column.

Several internal heat exchangers may be employed but only one, the preheater, is shown here. If a boundary is drawn around this system the availability balance equation may be written thus;





$$\Sigma Q_s \left(1 - \frac{T_o}{T_s}\right) - W = D \qquad (2.1)$$

- W = work supplied to the system

 $\Sigma Q_s \left(1 - \frac{T_o}{T_s}\right) = \text{the Carnot work equivalent across the}$ boundary due to the transfer of Q_z from an adjacent system at temperature T_z with respect to datum T_o . If the boiler heat Q_G is transferred at boiler temperature T_G and the evaporator heat Q_E at T_E then

$$= Q_E \left(1 - \frac{T_O}{T_E} \right) - Q_G \left(1 - \frac{T_O}{T_G} \right)$$

D = total system irreversibilities.

Note: If heat is transferred across the boundary at the datum temperature there is no Carnot work associated with it. Assuming Ω_{C} and Ω_{A} have been transferred at T_{O} , then

$$Q_C \left(1 - \frac{T_O}{T_O} \right) - Q_G \left(1 - \frac{T_O}{T_O} \right) = 0$$

If irreversibilities are considered as available work which has been degraded to thermal energy at T_0 without producing useful work then equation 2.1 states that the total available work minus the useful work done must equal the total system irreversibility.

2.2 The absorption system and the Carnot cycle

It is possible to represent the ideal absorption cycle as a



According to the second law the maximum possible output of the Carnot engine (W) operating between T_G and T_O is given by (36)

$$W = Q_G \left(1 - \frac{T_O}{T_G} \right) \tag{2.2}$$

Similarly the minimum work required by the heat pump to raise Ω_E from T_E to T_O is the carnot work

$$W = Q_E \left(\frac{T_O}{T_E} - 1\right) \tag{2.3}$$

Defining coefficient of performance as

$$COP = \frac{Energy \ transferred}{Energy \ input} = \frac{Q_E}{Q_G}$$
(2.4)

Substituting equations 2.2 and 2.3 in the COP expression yields

$$COP = \frac{\left(1 - \frac{T_O}{T_G}\right)}{\left(\frac{T_O}{T_E} - 1\right)}$$

$$COP_{CARNOT} = \frac{T_E \left(T_G - T_O\right)}{T_G \left(T_O - T_E\right)}$$
(2.5)
Equation 2.5 represents the maximum COP possible from an absorption system.

2.3 A relationship between Carnot and actual COP

Rewriting equation 2.1

$$Q_E\left(1-\frac{T_O}{T_E}\right)-Q_G\left(1-\frac{T_O}{T_G}\right) -W_p = D$$

Dividing through by $Q_{\mathbf{G}}$ and substituting from 2.4 gives

$$COP\left(1-\frac{T_O}{T_E}\right) - \left(1-\frac{T_O}{T_G}\right) - \frac{W_P}{Q_G} = \frac{D}{Q_G}$$

Therefore

ł

$$COP = \frac{\frac{W_P}{Q_G} + \frac{D}{Q_G} + \left(1 - \frac{T_O}{T_G}\right)}{\left(1 - \frac{T_o}{T_E}\right)}$$
(2.6)

$$COP = \frac{\frac{W_P}{Q_G} + \frac{D}{Q_G}}{\left(1 - \frac{T_o}{T_E}\right)} - \frac{T_E (T_G - T_O)}{T_G (T_O - T_E)}$$

Ignoring the work term because it is typically a second order effect

$$= \frac{T_E (T_G - T_O)}{T_G (T_O - T_E)} - \frac{D}{Q_G \left(\frac{T_O}{T_E} - 1\right)}$$

Substituting from equation 2.5

$$COP = COP_{CARNOT} - \frac{\frac{D}{Q_G}}{\frac{T_O}{T_E} - 1}$$
(2.7)

This means that a system COP may be calculated two ways, either by a first law energy balance approach or by subtracting the system irreversibility from the Carnot COP. The two results may be compared and by examining the individual irreversibility terms in equation 2.7 it is possible to establish which processes causes the largest departure from the Carnot value.

2.4 System irreversibilities

For convenience system irreversibilities may divided into two types, boundary and subsystem.

In 2.1 the overall system was defined such that only thermal energy and work crossed the boundary. If losses associated with the work transfer (the mechanical pump) are ignored then the availability losses associated with the boundary will be Carnot work lost due to thermal energy passing across the boundary from one temperature to another with no useful work output. (see equation 2.1) For this analysis it has been assumed that all boundary heat exchange takes place at constant temperature. If this was not the case an integration along the heat exchanger surface would be necessary.

Irreversibilities occuring internally may be dealt with by defining internal subsystems. Many of these subsystems involve streams as well as heat transfer. Using the same representation as 2.1 and including streams yields the equation



$$-\Delta B + \Sigma Q_Z \left(1 - \frac{T_O}{T_Z}\right) - W - \Delta (KE + PE) = D \qquad (2.8)$$

where B is the stream availability function which may be evaluated by selecting a datum and applying the following (35) equation

 $B\{T, P, x_i\} = (H - H_O) - T_O(S - S_O) + \sum n_i (RT_O \ln a_i^o)$ (2.9)

By replacing the activity term with a fugacity expression it is possible to evaluate 2.9 with data from the Peng-Robinson equation. The following fugacity terms were developed for the 3 possible binary mixture conditions.

Liquid phase only

$$RT_o\left[x_1 ln\left(\frac{\bar{f}_{1L}}{f_{1L}^o}\right) + x_2 ln\left(\frac{\bar{f}_{2L}}{f_{2L}^o}\right)\right]$$

Vapour phase only

$$RT_o\left[x_1 ln\left(\frac{\bar{f}_{1V}}{f_{1V}^o}\right) + x_2 ln\left(\frac{\bar{f}_{2V}}{f_{2V}^o}\right)\right]$$

Two phase

$$RT_o\left[x_1 ln\left(\frac{\bar{f}_{1V}}{f_{1V}^o}\right) + x_2 ln\left(\frac{\bar{f}_{2L}}{f_{2L}^o}\right)\right]$$

Appendix F shows availability values for pure refrigerants. Due to the absence of published data no checking was possible. All that can be said is that accuracy is dependent on the data generated by the Peng-Robinson equation. No doubt the liquid property values would contribute the greatest error, in particular liquid entropy.

It should be noted that availability changes across a given process are dependent on the datum conditions. Consider some arbitrary heat exchange process using a pure substance.



 $\Delta B_{process} = B_2 - B_1$

from equation 2.1

 $B_{1} = (h_{1} - T_{o}S_{1}) - (h_{o} - T_{o}S_{o})$ $B_{2} = (h_{2} - T_{o}S_{2}) - (h_{o} - T_{o}S_{o})$ $\Delta B = (h_{2} - T_{o}S_{2}) - (h_{1} - T_{o}S_{1})$

Here the datum $h_0 - T_0 S_0$ values cancel and ΔB depends only on the enthalpy and entropy at the two conditions and the datum temperature. Therefore if the same process is considered (same h, h₂ viz same temperatures and pressures) for two different availability datum temperatures (same datum pressure) ΔB will have different values. This means that it is not possible to produce a single general table of availability values as it is for say enthalpy; there has to be a separate table for each T₀ value.

An example of a subsystem irreversibility is the loss in availability through the streams entering and leaving a heat exchanger. The contra-flow heat exchanger example is treated in many texts (6)



An irreversibility occurs when the availability lost by the hot stream (B1-B2) is greater than the availability gained

(B4-B3) by the cold stream. In terms of a temperature profile for zero availability loss to occur both profiles would have to be coincident is matching profiles with no approach. In summary all the internal heat exchangers have the possiblity of incurring irreversibilities.

Now applying equation 2.8 to find the individual subsystem irreversibilities:

Evaporator

Consider that the superheating at the evaporator exit is negligible and therefore the refrigerant temperature/pressure will be constant at saturation conditions (T_E). On this basis the Carnot term can be written

$$Q_E\left(1-rac{T_o}{T_E}
ight)$$

This represents the Carnot work required to lift Q_E from T_E to T_O . Applying equation 2.8 to the evaporator in figure 2.1 we get

$$D_{EVAP} = Q_E \left(1 - \frac{T_o}{T_E}\right) - F_D (B_7 - B_8)$$

where D_{EVAP} is the total evaporator irreversibility and for cooling the temperature of the adjacent system T_{Z} is T_{E} .

Condenser

Assume subcooling and superheating to be zero and that the fluid is at the datum temperature.

$$D_{COND} = -(F_D + F_R)(B_5 - B_6))$$

where the subsystem temperature T_y adjacent to the boundary is T_0

Absorber

In the case of the absorber usually a two phase mixture results from the mixing of the bottoms and distillate streams. The feed concentration is far from being a pure substance and therefore a temperature change can be expected through the heat exchanger. However, if the mixing and heat exchange process are combined it can be argued that the heat of mixing may be removed as it forms with an isothermal process resulting. Using this and applying equation 2.8

$$D_{ABS} = -(F_D B_8 + F_B B_1 - F_F B_2)$$

where the subsystem temperature T_y adjacent to the boundary is T_0

Generator

Both the generator temperature and the temperature of the external system supplying heat have been assumed to be the same. This means that the generator Carnot term in equation 2.1 gives the total work available to the fluid in the system. In practice of course substantial irreversibilities may occur during the heat exchange process eg gas flame to fluid, and therefore care should be taken when comparing this data with actual systems. For the generator and column, from equation 2.8

$$D_{GEN} = -Q_G \left(1 - \frac{T_o}{T_G}\right) - (F_F B_4 + F_R B_6 - (F_D + F_R) B_5 - F_B - B_9)$$

where the external adjacent system temperature $T_{m{z}}$ is $T_{m{G}}$.

Pump

If some of the pump work which crossed the boundary was lost in the efficiency of the pump there would be an

irreversibility. For this analysis it will be assumed that all work crossing the boundary is imparted to the fluid and therefore applying 2.8 gives

$$W_p = -\Delta B = -F_F(B_2 - B_3)$$

Distillate throttling

$$D_{DE} = -F_D(B_6 - B_7)$$

Bottoms throttling

$$D_{BE} = -F_B(B_{10} - B_1)$$

Preheater

$$D_{PH} = -(F_F B_3 + F_B B_9 - F_F B_4 - F_B B_{10})$$

These system irreversibilities may be summed and used in equation 2.7 to determine COP or, in equation 2.1, to compile an availability balance around the cycle viz;

$$D = Q_E \left(1 - \frac{T_o}{T_E} \right) - F_D (B_7 - B_8) - (F_D + F_R) (B_5 - B_6)$$

-Q_G $\left(1 - \frac{T_o}{T_G} \right) - (F_F B_4 + F_R B_6 - (F_D + F_R) B_5 - F_B - B_9)$
-(F_D B_8 + F_B B_1 - F_F B_2)
-F_D (B_6 - B_7) - F_B (B_{10} - B_1)
-(F_F B_3 + F_B B_9 - F_4 B_4 - F_B B_{10})

$$= Q_E \left(1 - \frac{T_O}{T_E}\right) + F_F B_2 - F_F B_3 - Q_G \left(1 - \frac{T_O}{T_G}\right)$$

$$D = Q_E \left(1 - \frac{T_O}{T_E}\right) - W_p - Q_G \left(1 - \frac{T_O}{T_G}\right)$$

This checks with equation 2.1 ie the availability changes around the internal circuit sum to zero.

2.5 Avoidable and unavoidable irreversibilities

For this system there is no way of deriving work from the mixing process and therefore there will always be unavoidable mixing irreversibilities.

The only way this loss can be minimised is to change the stream compositions and not the actual mixing process. By keeping the availabilty of the streams being mixed as close as possible the losses are minimized. In the case of mixing the distillate and bottom flows minimising of the mixing loss is done by changing the stream availabilities with internal heat exchangers. (see chapter 3) The column and its mixing may be assessed by taking an overall view. If the minimum work of separation is taken as the increase in availability between the products (12) and the feed then

> Column loss = equivalent Carnot input - minimum work of separation

An example of a theoretically avoidable irreversibility occurs in the contra flow heat exchanger. If there is no approach and the temperature profiles of the two streams are coincident then the loss is zero.

CHAPTER 3

Nomenclature

 $\begin{array}{l} D = \mbox{irreversibility} \\ Q = \mbox{heat} \\ T = \mbox{temperature} \end{array}$

Subscript

E = evaporator C = condenser G = generator (boiler) o = datum

ABSORPTION SYSTEM

3.1 Introduction

Chapter 3 deals with the application of the computer model to an ideal system, a practical system and a prototype system with an emphasis on second law analyses. The sections in the chapter and their objectives are as follows:

Second law and Carnot performance

- To configure the second law COP expression (equation

2.7) for use with the absorption cycle.

The ideal System

- To model an ideal system, ie one in which all avoidable irreversibilites are removed so that the COP predicted is the maximum possible.
- To investigate the effect of internal heat exchangers
 on COP for an ideal system via second law analyses.
- To assess the effect of performance variables such as mixture concentrations and operating temperature on COP for the ideal system.

Practical cycle

- To model an air cooled system and show a typical
 cooling COP using conditions encountered in practice.
- Model a heat pump mode to predict a heating COP for a practical air to air system.
- Model an organic system using published performance data for a commercially available Lithium Bromide plant and compare COP values for the two systems.

Prototype performance

- Model an existing prototype system and compare the measured and predicted COP's.

For this analysis Coefficient of Performance (COP) has been selected as the measure of performance. It is recognised that in the application of this equipment many other factors, in addition to COP, should be considered, however, since this work is confined to the thermodynamics of the internal system COP alone has been used.

3.2 Ideal cycle and second law analysis

By using a second law analysis (equation 2.7) it is possible to assess the effect of the various irreversibilities on COP. Further to this by distinguishing between avoidable and unavoidable irreversibilities it should be possible to establish a maximum achievable COP. Using the computer model it is possible to eliminate some avoidable irreversibilities by

- Selecting a zero approach on all heat exchangers
 which means rejecting all heat at ambient conditions,
 transferring all heat to the evaporator at T_E and
 supplying the boiler heat at T_E.
- Ignoring friction as well as heat and "cold" losses for the system pipework.

System performance may then be analysed by substituting the remaining irreversibilities in equation 2.7 . For example using the cycle shown in figure 3.1 and the conditions listed above equation 2.7 yields

$$COP = Carnot COP - \frac{(D1+D2+D3+D4)}{Q_g \begin{pmatrix} T_{--} \\ -- \\ T_E \end{pmatrix}} \qquad (3.1)$$





BASIC CYCLE

FIGURE 3.1

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where the irreversibilities are

- D1 = column
- D2 = absorber mixing
- D3 = Throttling of bottom flow
- D4 = Throttling of distillate flow

The Carnot COP expression derived in section 2.2 is

Carnot COP =
$$\frac{T_{E}(T_{G} - T_{C})}{(T_{C} - T_{E})}$$
 (3.2)

In equation 3.2 the the boiler temperature (T_G) is dependent on the ambient temperature (T_O) and so this relationship may be plotted for a specified evaporator temperature (T_E). In the system (figure 3.1) the distillation column pressure is determined by the condensing condition, which in turn, is established by the heat rejection temperature. Assuming heat is rejected to the environment and not to some other system then the heat rejection temperature (T_O) determines the generator pressure and therefore its temperature (T_G).

This relationship between T_{O} and T_{G} is determined by both mixture properties and by the system operating parameters. It should be noted that an ideal case will be considered and therefore the condensing temperature (T_{C}) will be assumed to be the ambient temperature (T_{O}) . Figure 3.2 shows T_{C} versus T_{O} for 3 mixtures where the distillate and bottoms mass concentrations are 0.99 and 0.05 respectively. Therefore for a given ambient temperature the Carnot COP can be calculated for a range of T_{E} values (figure 3.3). This then defines the maximum thermodynamically possible COP.

It is interesting to note that even though the ammonia-water cycle (R717 R718) Carnot COP is much higher than the organic

PARAMETERS FOR FIGURE 3.2

Distillate mass concentration 0.99 Bottom mass concentration 0.05 Assume column at constant pressure

188 128 200 48 8 8 8 BOTTOM FLOW SATURATION CONDITION VERSUS CONDENSING TEMPERATURE 1 22 RIZ R1 1 **R717** 1 - R113 - R718 RRS 8 8 \$ **A**

:Saturation temperature of bottom composition Tg Deg C corresponding to saturation pressure of distillate at Tc

Fig 3.2

CONDENSING TEMPERATURE To DEG C

PARAMETERS FOR FIGURE 3.3

Data was derived from equation 3.2

Distillate mass concentration 0.99

Bottom mass concentration 0.05

 T_{G} = Saturation temperature of bottom composition corresponding

to saturation pressure of distillate at 30 deg C.



CARNOT COP

cycles the actual practical COP's are of the same order (7) because of the large heat of mixing value. In terms of equation 3.1 this means a large D2 value.

To establish the importance of the various irreversibilities a cycle with a range of evaporator temperatures will be considered. This range was chosen so as to cover typical comfort air conditioning operation for Adelaide (see section 3.3). Various internal heat exchangers will be added later in an effort to reduce irreversibilities but firstly consider the basic cycle.

3.2.1 Basic Cycle

Equation 3.1 was applied to the basic cycle for a range of evaporator temperatures (table 3.1) so that the effects of the various irreversibilities could be assessed. For example in figure 3.4, which is a plot of the data in table 3.1, the distance between the Carnot COP line and the case A line is an indication of the effect of the column irreversibilities. Similarly the effect of the absorber mixing losses is indicated by the distance between case A and case B. Only the two irreversibilities have been considered here because the effect of both the distillate and bottom flow throttling processes was negligible.

If all of the irreversibilities were unavoidable then the final plot (case B) would represent the maximum COP theoretically possible. For example the absorber mixing irreversibilities are unavoidable because no useful work can be derived from the mixing process. However, this is not the case for the column.

PARAMETERS FOR FIGURE 3.4 AND TABLE 3.1



•	Irreversiblity terms included in equation 3.1
Case A	Column
Case B	Column and absorber mixing

TABLE 3.1: COP VALUES FROM EQUATION 3.1					
evap temp deg C	0	5	10	15	20
Carnot	1.73	2.11	2.69	3.65	5.57
Case A	0.73	0.90	1.14	1.42	1.79
Case B	0.16	0.19	0.26	0.33	0.43



CARNOT CASE A CASE B

EVAPORATOR TEMPERATURE Te deg C

FIGURE 3.4

COP VERSUS EVAPORATOR TEMPERATURE

The column requires a minimum reflux flow to operate and irreversibilities associated with the production of this minimum flow are unavoidable. Irreversibilities created by the production of reflux flow in excess of the minimum are theoretically avoidable in a column of infinite stages. The computer model does not separate the two and therefore it is not possible to quantify the avoidable value but it is possible to estimate it. Later, in section 3.2.7, it is shown that for a 30 deg C ambient a reduction in reflux ratio from around 1.15 to 1.00 results in a COP increase of approximately 7 percent. Since the column irreversibilities account for in excess of 50 percent the avoidables are small in comparison with the unavoidables and therefore case B is within say 10 percent of being the maximum attainable COP.

All the irreversibilities being discussed here are due to the mixing of streams. This is illustrated in the column analysis print out (Appendix H, fig H1) which shows large feed plate and stripping section losses. If the feed stream availability could be brought closer to the feed plate value then at least the feed plate losses would be reduced!

To improve the maximum possible COP a reduction in one or more of the irreversibilities is necessary. Since the column and mixing values together represent approximately 90 percent of the total (fig 3.6) they invite close inspection. Unfortunately, other than the reduction of the reflux flow to the minimum it is not possible to eliminate irreversibilities. A change in the cycle is necessary to realize an improvement and two possibilities are;

 reduction of feed-plate availability loss by increasing the feed availability towards that

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PERCENT

PARAMETERS FOR FIGURE 3.6

Ambient (T_o) 30 deg C Boiler temp (T_G) 101.1 deg C Bottom mass concn. 0.05 Distillate mass concn. 0.99 R11 and R22 mixture

CUMULATIVE PERCENT OF AVAILABILITY VS EVAP TEMP



FIGURE 3.6

of the feed-plate liquid

making use of the availability in the bottoms stream instead of 'losing' it to the mixing process in the absorber.

The common practice of preheating the feed flow with the hot bottoms stream achieves both of the above.

3.2.2 Cycle with preheater

Figure 3.7 shows typical data for this system at a 5 deg evaporator temperature (see appendix H, figure H2 for complete computer model print out) while table 3.2 is a precis of the data for the full range of evaporator temperatures.

For this preheater cycle data has been tabulated to illustrate typical trends in performance. For example the absorber output reflects the evaporator capacity and varies little whilst the condenser kW follow the boiler input. The evaporator temperature determines the low side pressure in the system which, together with the ambient temperature, determines the liquid concentration leaving the absorber. Therefore as the evaporator temperature drops the feed concentration is reduced to achieve the required lower evaporator temperature and, in order to maintain a constant bottom concentration of 0.05, the feed/distillate ratio is increased.

Figure 3.8 includes both first law COP values from table 3.2 as well as a plot of the second law data. Here again the final COP, case D, is not the maximum possible value because the column irreversibility contains an avoidable reflux flow component.



PREHEATER CYCLE

FIGURE 3.7

PARAMETERS FOR FIG 3.7, FIG 3.8 AND TABLE 3.2

Ambient air temperature (To) 30 deg C Boiler temperature (Tg) 101 deg C Bottom mass concentration 0.05 Distillate mass concentration 0.99 R11 & R22 mixture

	Irreversibility terms included in equation 3.1
CASE A	Column
CASE B	Column and absorber mixing
CASE C	Column, absorber mixing and precooler
CASE D	All losses

SYSTEM DATA

No	Temp Deg C	Avbty kJ/kg	
1	101.1	5.8	
2	30.0	2.9	
3	30.0	2.5	
4	30.0	65.6	-
5	5.0	63.8	
6	5.8	50.6	
7	30.0	9.0	
8	29.9	9.4	
9	66.4	11.5	
10	68.3	6.8	liquid
	68.3	40.0	vannur



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EVAPORATOR TEMPERATURE To Deg C

FIGURE 3.8

TABLE 3.2: PREHEATER SYSTEM DATA

evap temp deg C	0	5	10	15	20
Pump kW	0.173	0.107	0.046	0.039	0.021
Boil kW Cond kW	25.19 24.87	21.18 20.91	18.26 18.03	16.17 15.97	14.88 14.75
Evap kW Abs kW	10.00 10.3	10.00 10.26	10.00 10.22	10.00 10.18	10.00 10.13
Reflux [#] ratio	1.17	1.15	1.14	1.14	1.18
Plates Dist kg/s FOD ratio [†]	11 0.059 6.3	12 0.058 4.41	13 0.057 3.17	14 0.057 2.31	14 0.056 1.71
Feed conc	0.199	0.263	0.346	0.457	0.601
÷-			kJ/s		
Boiler Work input	4.788	4.024	3.469	' 3.071	2.827
Losses					
Column Mixing Dist. exp Bottom exp Preheater	4.453 1.248 0.172 0.144 0.032	1.993 1.129 0.101 0.08 0.057	1.773 0.991 0.048 0.043 0.054	1.724 0.831 0.012 0.02 0.047	1.84 0.639 0.007 0.046
Useful Evap work	1.066	0.868	0.677	0.493	0.315
	COP				
Carnot CASE A CASE B CASE C CASE D	1.73 0.843 0.392 0.278 0.266	2.114 1.067 0.474 0.378 0.349	2.69 1.315 0.547 0.476 0.434	3.65 1.601 0.614 0.576 0.519	5.57 1.944 0.685 0.685 0.595
First law	0.394	0.47	0.546	0.617	0.671

Reflux ratio = actual reflux / minimim reflux

↑ FOD = feed flow rate / distillate flow rate

For all these tests a reflux value approximately 20 percent greater than the minimum was used to achieve manageable computing times and to avoid the accuracy problems associated with a large number of plates (see chapter 4). In practice a reflux flow less than this would be desirable but since the effect of reflux can be estimated the value used for calculation is not critical.

Other than this reflux component all the other irreversibilities are considered unavoidable. In the case of the preheater even though a zero approach was used for the controlling end an irreversibility occurred because the stream temperature profiles did not match. The differing stream flow rates and specific heat capacities resulted in a temperature difference at the other end of the heat exchanger. For example the difference for the system shown in figure 3.7 is 101.1 - 66.4 = 34.7 degrees.

Care should be taken when making comparisons between the first and second law COP's. Theoretically the first law COP value should equal the second law calculation with all the irreversibilities taken into account. Unfortunately, due to inaccuracy in the availability values (see chapter1), this is not the case.

For example, in table 3.2 for a 5 deg C evaporator the first and second law calculations yielded COP's of 0.47 and 0.349 respectively and, using the first law value as a basis, this represents a -25% variation. The second law value was arrived at by subtracting all the individual components, each with their own error, from the Carnot value of 2.1 and the accumulation of the errors has resulted in this

variation. Consequently in figure 3.8 the plot D and the first law values do not coincide.

Despite these inaccuracies the improvement in COP of the preheater cycle over the basic cycle is obvious. For example at a 5 deg C evaporator temperature the first law COP value changed from 0.32 (table 3.1) to 0.47 (table 3.2) a 47% increase.

By pursuing the concept of reducing mixing losses a further COP improvement may be had by making as much use as possible of the high availability leaving the evaporator before it is lost during mixing in the absorber. For example in figure 3.7 the distillate vapour leaves the evaporator with an availability of 51 kJ/kg and is mixed in the absorber with a bottom stream having an availability of 2.5 kJ/kg. Since the heat of mixing is rejected this availability is lost as heat in the absorber.

3.2.3 Cycle with preheater and precooler

Figure 3.9 illustrates the cycle and shows data extracted from the 5 deg C evaporator example in Appendix H (figure H3) whilst table 3.3 summarizes the results for a range of evaporator temperatures. This data may be compared with table 3.2 to assess the effect of the precooler. Firstly comparing case B only, the improvement in COP is 0.09 ie 0.474 to 0.564 however, when all the irreversibilities are considered (in case D) the gain is of the order 0.07; from 0.349 to 0.423, a 20% increase. One reason for the reduction from 0.09 to 0.07 is the introduction of an irreversiblity in the new precooler heat exchanger.

PARAMETERS FOR FIGURE 3.9 AND TABLE 3.3

Ambient air temperature 30 deg C Boiler temperature 101 deg C Bottom mass concentration 0.05 Distillate mass concentration 0.99 Reflux nominally 20% above minimim R11&R22 mixture

	Irreversibility terms included in equation 3.1
CASE A	Column
CASE B	Column and absorber mixing

CASE C Column, absorber mixing, precooler and preheater

CASE D All losses

TABLE 3.3	: COP VAL	UES FOR A	RANGE	OF EVAP	TEMPERATURES
evap temp deg C	0	5	10 .	15	20
Carnot	1.73	2.11	2.69	3.65	5.57
Case A	0.84	1.063	1.315	1.6	1.95
Case B	0.475	0,564	0.649	0.723	0.796
Case C	0.396	0.497	0.598	Ŭ.687	0.796
Case D	0.344	0.423	0.508	0.583	0.659
1st law	0.44	0.515	0.588	0.653	0.697



PREHEATER-PRECOOLER CYCLE

FIGURE 3.9

SYSTEM DATA

R	Temp Deg C	Avbty kJ/kg	
1 2 3 4 5 6 7 8 9 10 11 12	101.1 30.0 30.0 5.0 5.8 30.0 29.9 66.4 68.3 30.0 17.2	5.8 2.9 2.5 65.6 66.4 50.6 9.0 9.4 11.5 6.8 40.0 47.6 67.2	liquid vapour

COLUMN DATA

Plate No.	Temp Deg C	Avbty Change kJ/s ≭	Comment
1	30.0	-0,006	Condenser
2	30.7	-0.018	oondensei
3	32.0	-0.026	
4	34.6	-0.043	
5	39.7	-0.078	
6	48.5	-0.105	
7	58.3	-0.069	
8	65	-0.027	
9	68.3	-0.251	Feedplate
10	74.1	-0.175	·
11	85.6	-0.493	
12	101.1	3.137	Boiler 19.3kW

* Negative indicates an availability loss

An inspection of the column data in figure 3.9 reveals that the greatest availability losses occur on the stripping section plates. In figure 3.9 the boiler plate shows a gain in availability but in fact there is an availability loss of 0.534 kJ/s incurred. This loss value is calculated by subtracting the nett increase of 3.137 kJ/s for the plate from the equivalent Carnot work input of 3.671 kJ/s.

The next section demonstrates the effect of an internal column heat exchanger on these stripping section losses.

3.2.4 Cycle with preheater, precooler and column heat exchanger.

For this system no attempt has been made to model a heat exchanger on each individual plate but rather a whole section approach has been taken with equal heat exchange per plate. Figure 3.10 shows COP data extracted from the computer model print out figure H4 (appendix H) while table 3.4 shows COP data for a range of evaporator temperatures.

The addition of the column heat exchanger produces an improvement in the COP which is greatest at the lower evaporator temperatures. As T_E drops the feed/distillate ratio increases and the larger bottom flow contributes more heat to the column. For example at a 5 deg C evaporator temperature the improvement in COP over the precooler/preheater system was 0.029 (0.515 to 0.544) but at a 20 deg C evaporator the increase was only 0.008 (0.697 to 0.705).

PARAMETERS FOR FIGURE 3.10 AND TABLE 3.4

Ambient air temperature (To) 30 deg C Boiler temperature (Tg) 101 deg C Bottom mass concentration 0.05 Distillate mass concentration 0.99 Reflux nominally 20% above the minimum R11 & R22 mixture

	Irreversibility terms included in equation 3	3.1
CASE A	Column -	v
CASE B	Column and absorber mixing	
CASE C	All losses	

TABLE	E 3.4: CO	P VALUES	FROM EQL	JATION 3.1	
evap temp deg C	0	5	10	15	20
Carnot	1.73	2.11	2.69	3.65	5.57
CASE A	0.996	1.172	1.387	1.62	1.88
CASE B	0.582	0.645	0.689	0.721	0.725
CASE C	0.386	0.449	0.527	0.593	0.643
1st law	0.494	0.544	0.615	0.669	0.705



Bottom flow

PREHEATER-PRECOOLER-BOTTOM HX CYCLE

FIGURE 3.10

SYSTEM DATA

R	Temp Deg C	Avbty kJ∕kg	
1 2 3 4 5 6 7 8 7 10* 11 12 13	101.1 29.9 30.1 30.0 5.0 5.8 30.0 29.9 57.9 64.5 64.5 30.0 17.2 69.5	5.8 2.9 2.5 65.6 66.4 50.6 9.0 9.4 8.8 7.8 41.8 47.6 67.2 2.7	liquid vapour

*****Feedplate

COLUMN DATA

Plate	Temp	Avbty Change	Comment	
No.	Deg C	kJ/s		
1 2 3 4 5 6 7 8 9 10 11 12	30.0 30.7 31.9 34.2 38.5 45.7 54.3 61.0 64.5 73.2 86.1 101.1	-0.006 -0.016 -0.02 -0.029 -0.049 -0.069 -0.054 -0.022 -0.133 0.02 -0.119 2.98	Condenser Feedplate 2.79kW 2.79kW Boiler 18.3kW	

Overall the improvement comes about because availability from the bottom stream is being used to vapourize some of the downward liquid flow before it gets to the generator. Therefore the generator heat input is reduced and a COP improvement results.

It may be argued that use of the stripping section heat exchanger is unnecessary since this bottom stream availability can be recovered in the preheater and returned to the column via the feed flow anyway. A comparison of figures 3.9 and 3.10 reveals that the the availability of the bottom flow leaving the preheater is the same (2.9 kJ/kg) for both configurations. However, this is not the case because the column heat exchanger system shows a superior COP. This is because the availability transfer via the preheater and feed flow path incurs greater irreversibilities than the more direct route via the column heat exchanger.

This choice of path is fundamental to optimum design. In the case of the absorption cycle we are seeking to transfer the work potential, which is input as heat to the boiler, to the evaporator via a path of least loss or degredation. Indeed the transfer of work input to the point of application via a path of minimum loss most often determines the design of thermodynamic systems.

Despite the use of the stripping section heat exchanger an irreversibility still remains in the preheater due to the mismatch of temperature profiles. Whilst it is not possible to modify the specific heat capacities of the streams it is possible to change the flow rates by splitting the feed

stream. By reducing the feed flow it is theoretically possible to match the temperature profiles of the two streams. The proportion of the feed flow diverted away from the preheater may be directed to a heat exchanger in the rectification section of the column.

3.2.5 Cycle with preheater/precooler and both stripping and rectification heat exchangers

Initially 2 percent of the feed flow was directed through the rectification heat exchanger (figure 3.11) and at a 5 deg C evaporator temperature this resulted in a COP improvement over the stripping heat exchanger system of 0.07, ie from 0.544 (table 3.4) to 0.616 (table 3.5) ie a 12% increase.

This improvement in COP can be attributed to reduced irreversibilities in the column and in the preheater. Here part of the total reflux requirement is generated (condensed) internally in the column by the cooler feed stream. This incurs a penalty in column size. A comparison of plate numbers in figures 3.10 and 3.11 bears this out; the dual heat exchanger column has 13 plates while the single only 12, an 8% increase in column size. The feed and feed plate availabilities are still close, 8.68 kJ/kg and 9.81 kJ/kg respectively, thus losses here are minimal.

By reducing the flow through the feed side of the preheater the temperature profile of this stream can be brought closer to that of the bottom flow. In an effort to establish an optimum proportion for the feed split tests were conducted for 1,2,4 and 6 percent of the feed flow through the rectification section heat exchanger and the results plotted

PARAMETERS FOR FIGURE 3.11 AND TABLE 3.5

Ambient air temperature 30 deg C Boiler temperature 101 deg C Bottom mass concentration 0.05 Distillate mass concentration 0.99 2 percent feed flow thru rectification heat exchanger R11 & R22 mixture

	Irreversibility terms included in equation 3.1
CASE A	Column
CASE B	Column and absorber mixing

TABLE	3.5: CC	P VALUES	FROM EQ	UATION 3.1	
evap temp deg C	0	5	10	15	20
Carnot	1.73	2.11	2.69	3.65	5.57
CASE A	1.098	1.299	1.468	1.687	÷.:
CASE B	0.644	0.7	0.729	0.75	
ist law	0.541	0.616	0.653	0.696	
8

SYSTEM DATA



Bottom flow

PREMEATER-PRECOOLER-BOTTOM-RECTIFICATION BX CYCLE

FIGURE 3.11

	Temp Deg C	Avbty kJ/kg	
1 2 7	101.1	5.8 2.9	
১ 4 5	30.1 30.0 5.0	2.5 65.6 66.4	
6 7	5.8 30.0	50.6	
8 9	29.9 53.0	9.4 8.7	
10	61.0 61.0	9.2 43.5	liquid vapour
11 12	30.0	47.6	
13	62.2	8.8	

*****Feedplate

Plate No.	Temp Deg C	Avbty Change _ kJ/s	Comment
1	30.0	-0.005	Condenser
2	30.7	-0.007	0.01kW
3	31.7	-0.001	0.01kW
4	33.6	0.006	0.01kW
5	36.8	0.014	0.01kW
6	42.0	0.022	0.01kW
7	49.2	0.031	0.01kW
8	56.4	0.016	0.01kW
9	61.0	-0.107	Feedplate
10	65.9	0.095	2.27kW
11	74.0	-0.011	2.27kW
12	86.4	-0.153	2.27kW
13	101.1	2.598	Boiler 16.1kW

in figure 3.12 indicate an optimum around one percent.

For most of the tests the heat exchanged in the rectification section was approximately one tenth of that transferred in the stripping section.

For the column, even with heat exchangers fitted, the feed plate, boiler and bottom plate irreversibilities remained dominant. At the feed plate the availability loss can be attributed, in part, to the mixing of the column and feed flows which have different temperatures and concentrations. The remaining loss is due to the difference between the availabilities of vapour and liquid in equilibrium. This is the main reason for the large loss in the boiler too.

The total loss can be minimised by

- having a mixture where the equilibrium availability values for liquid and vapour are closer together. Support for this appears towards the top of the column. Condensing is taking place at the datum temperature and therefore the difference in availability between liquid and vapour is due only to the non-datum pressure. (at datum temperature and pressure liquid and vapour availability are equal, see section 2.2) At the condenser and on the top few plates, (see figure 3.11) availability losses are negligible in comparison with the boiler losses.
- reducing the system flow rate. Mixing irreversibilities are a function of both specific availability values of the streams and of the stream flow rates. This explains why the lower flows of a system with minimal reflux and with a high latent heat refrigerant are an advantage.



COP VS HEAT EXCHANGER FLOW FOR A RANGE OF EVAP TEMPERATURES

PERCENT OF FEED FLOW THROUGH COLUMN HEAT EXCHANGER

FIGURE 3.12

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FIRST LAW COP

These low flows minimise the effect of the unavoidable irreversiblities.

Another important factor is the relationship, for a given mixture, between temperature and availability. Consider both the minimum work of separation and the Carnot input for the system shown in figure 3.11. These two are related via the characteristics of the fluid properties. Using data from the column analysis of figure H5 the following may be compiled for the boiler.



5.8 kJ/kg 0.05 (mass concn)

Carnot input =
$$Q_G \begin{pmatrix} 1 - \frac{T_O}{T_O} \\ T_G \end{pmatrix}$$

= 3.477 kJ/s

Therefore availability entering the boiler is

 $3.477 + (0.2655 \times 4.8) = 4.75 \text{ kJ/kg}$

and availability leaving is

 $(0.1804 \times 5.8) + (0.0857 \times 32.9) = 3.87 \text{ kJ/kg}$ Total availability loss is

4.75 - 3.87 = 0.88 kJ/kg

Minimum work of separation = the availability difference of the streams entering and leaving = $3.87 - (0.2655 \times 4.8) =$ 2.59 kJ/s

For these conditions the theoretical minimum work of separation (12) is 2.59 kJ/s but at the same time the

equilibrium temperature of 101.1 deg C requires, via the Carnot term, an input of 3.477 kJ/s. The difference is the 0.88 kJ/s availability loss. If the temperature required had been 89 deg C for example then the Carnot requirement would have equalled the separation work required and the loss would have been zero. Examination of other pairs of working fluids may reveal a combination with more favourable characteristics.

As far as the usefulness of the rectification heat exchanger is concerned it is suggested that the feed split option be tried on a prototype to establish if the magnitude of the COP improvement justifies the additional complexity in control and manufacture.

3.2.6 Variations in ambient conditions.

COP variations due to ambient temperature changes have been considered on the ideal preheater/precooler/bottom system only. Table 3.6 and figure 3.13 show 38 deg C results while figure 3.14 compares the Carnot and first law COP's for ambients of 30 deg C (table 3.4) and 38 deg C (table 3.6).

For example at a 5 deg C evaporator temperature the ambient temperature increase from 30 deg C to 38 deg C caused a reduction in the First Law COP of 0.12 which is significant but small in comparison with the corresponding Carnot COP drop of 0.49. Referring to equation 3.1 if the irreversibility term had decreased at the same rate as the Carnot COP then the first law values would have differed by 0.49 too. The fact that the first law COP variation was smaller indicates that as the ambient temperature increases the irreversibility term in equation 3.1

PARMETERS FOR FIGURE 3.13 AND FIGURE 3.14 PLUS TABLE 3.6

Ambient air temperature (To) 38 deg C Boiler temperature (Tg) 112.4 deg C Bottom mass concentration 0.05 Distillate mass concentration 0.99 Reflux nominally 20 percent above minimum R11 and R22 mixture

	Irreversibility terms included in equation 3.1
CASE A	Column
CASE B	Column and absorber mixing

TABLE	3.6: COP	for a 38	3 deg C a	ambient	
evap temp deg C	0	5	10	15	20
Carnot	1.386	1.626	1.95	2.416	3.143
CASE A	0.78	0.914	1.109	1.312	
CASE B	0.492	0.544	0.61	0.661	
1st law	0.375	0.428	0.505	0.573	







PRSHEATER-PRECOOLER-BOTTOM HX CYCLE

FIGURE 3.13

B	Temp Deg C	A∨bty kJ/kg	
1 2 3 4 5 6 7 8 9 10* 11 12 13	112.4 37.8 38.1 38.0 5.0 5.8 38.0 37.8 79.3 84.8 84.8 38.0 21.4 89.3	11.3 7.9 7.3 72.6 73.1 53.4 10.7 11.3 11.1 11.0 41.9 49.0 74.8 8.5	liquid vapour

* Feedplate

COLUMN DATA

Plate Temp Deg C Avbty Change kJ/s Comment 1 38.0 -0.007 Condenser 2 38.6 -0.025 Condenser 3 40.0 -0.083 -0.083 5 49.0 -0.167 -0.234 7 72.7 -0.153 -0.056				
1 38.0 -0.007 Condenser 2 38.6 -0.025 Condenser 3 40.0 -0.043 -0.083 4 42.8 -0.083 -0.167 6 60.1 -0.234 -0.153 7 72.7 -0.153 -0.056	Platé No.	Temp Deg C	Avbty Change kJ/s	Comment
9 84.8 -0.161 Feedplate 10 91.9 0.23 3.76kW 11 101.1 0.142 3.76kW 12 112.4 3.95 Boiler 23.2k	1 2 3 4 5 6 7 8 9 10 11	38.0 38.4 40.0 42.8 49.0 60.1 72.7 81.0 84.8 91.9 101.1 112.4	-0.007 -0.025 -0.043 -0.083 -0.167 -0.234 -0.153 -0.056 -0.161 0.23 0.142 3.95	Condenser Feedplate 3.76kW 3.76kW Boiler 23.2kW



COP VS EVAPORATOR TEMPERATURE; AMBIENT VARIATION

FIGURE 3.14

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g

$$COP = COP_{CARNOT} - \frac{\frac{D}{Q_G}}{\frac{T_O}{T_E} - 1}$$

diminishes more rapidly than the Carnot term. In terms of figure 3.14 this means that the distance between the Carnot and first law plots is less for 38 Deg C than it is for the 30 deg C ones. Therefore it is the relationship between the irreversibilities and boiler input which is important. A study of the relationship between D and $Q_{\rm G}$ for various mixtures and temperature conditions could be used to establish desirable characteristics for these mixtures.

3.2.7 Variations in Reflux Ratio

Reflux ratio was varied for the precooler/preheater/bottoms cycle at both 30 deg C and 38 deg C ambients for a 5 deg C evaporator. Figure 3.16 shows increasing COP with diminishing reflux while figure 3.15 shows data extracted from appendix H, figure H7.

For a given set of conditions reducing the reflux ratio reduces the availability loss in the column whilst, apart from an insignificant change at the preheater, all the other losses remain constant. Obviously from the point of view of maximizing COP a low reflux ratio is desirable but since the column size/cost increases as reflux ratio decreases economics will determine the optimum.

3.2.8 Variation of bottom concentration

By increasing the ratio of feed flow to distillate flow it is possible to change the concentration of the bottom product. This was done for the preheater/precooler/bottom



Bottom flow

PREHEATER-PRECOOLER-BOTTOM HX CYCLE

FIGURE 3.15

PARMETERS FOR FIGURE 3.15 AND FIGURE 3.16

Bottom mass concentration 0.05 Distillate mass concentration 0.99 Reflux nominally 20 percent above minimum R11 and R22 mixture

SYSTEM DATA

NO .	Temp Deg C	Avbty kJ/kg	
1 2 3 4 5 6 7 8 9 10* 11 12 13	101.1 29.9 30.1 30.0 5.0 5.8 30.0 29.9 58.5 45.3 65.3 30.0 17.2 70.4	5.8 2.9 2.5 65.6 66.4 50.6 9.0 9.4 8.8 7.6 41.4 47.6 67.2 2.7	liquid vapour

*Feedplate

COLUMN DATA

Flate	Temp	Avbty Change	Comment
No.	Deg C	kJ/s	
1	30.0	-0.009	Condenser
2	30.7	-0.039	
3	32.3	-0.085	
4	36.8	-0.219	
5	47.9	-0.475	



COP AND NUMBER OF IDEAL PLATES VS REFLUX RATIO

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C 0 P

Fig 3.16

heat exchanger cycle shown in figure 3.17, a sample program output is shown in Appendix H figure H8.

Since the computer model establishes the number of plates required in the column for each evaporator condition it was difficult to assess the effect of bottom concentration without a change in the number of plates and a variation in reflux ratio. In figure 3.18 the dislocation between FOD values of 5.2 and 5.3 is caused by a change in the number of plates from 12 to 11.

Ignoring the discontinuity for the moment and considering a feed/distillate range from 4.4 to 5 there is an increase in bottom concentration from 0.05 to 0.08 together with a 0.02 COP increase. However, at the same time there was a reflux ratio reduction from 1.24 to 1.155 which, according to section 3.2.7, represents a COP increase of 0.02 approximately. Consequently it is assumed that the effect of bottom concentration on COP is negligible. Of greater significance is the fact that a bottom concentration increase from 0.05 to 0.13 decreased the boiler temperature from 101 deg C to 82 deg C thus making the system more attractive for use with low grade heat.

3.2.9 Variation of distillate concentration

One of the effects of reducing the distillate concentration is to produce a temperature gradient across the evaporator surface. Within reason this is not considered detrimental to performance, in fact Connon et al (10) consider this an advantage for heating situations. For this work a gradient of 5 degrees, approximately half of that allowed by Connon et al, has been selected as tolerable where typical air



PARMETERS FOR FIGURE 3.17 AND FIGURE 3.18

Ambient temperature (To) 30 deg C Evaporator temperature (Te) 5 deg C Distillate mass concentration 0.99 R11 and R22 mixture

Bottom flow

PREHEATER-PRECOOLER-BOTTOM HX CYCLE



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. <u>.</u> . .

temperatures might be 24 deg C approaching the evaporator and 15 deg C leaving.

In terms of a second law consideration a contra flow refrigerant to air heat exchanger could benefit from the closer matching of temperature gradients via a reduction in availability loss. Also the work required to raise the heat from the evaporator condition to the reference condition would be less because part of the surface is warmer. It was thought that this reduction in work as well as less effort required for distillate and absorbant separation would improve the COP.

At ambients of both 30 deg C and 38 deg a 7 percent increase in COP resulted from a reduction in distillate concentration from 0.99 to 0.93 (figure 3.20).

It is suggested that this approach is worth investigating on a prototype. A distillate concentration of 0.93 resulted in a 5.9 degree evaporator gradient. (see figure 3.19 or, for a complete output, appendix H,figure H9)

3.2.10 Mixture variations

Even though an extensive investigation of binary mixtures suitable for use in absorption refrigerators is beyond the scope of this report it was decided to test an R12 R113 mixture as well as R11 R22.

This was prompted by the fact that the former showed Carnot COP values approximately 10 percent higher than the latter. The precooler/preheater/bottom heat exchanger combination (figure 3.21) was tested and the results are shown in table



PARAMETERS FOR FIGURES 3.19 AND 3.20

Ambient temperature (To) 30 deg C Bottom mass concentration 0.05 Reflux nominally 20 percent above minimum Rll and R22 mixture

SYSTEM DATA

R	Temp Deg C	Avbty kJ/kg	
1	99.2	5.5	
2	29.9	2.9	
3	30.1	2.5	
4	30.0	58.0	
5	5.0	58.8	
6	10.9	43.6	
7	30.0	8.2	
8	29.9	8.6	
9	52.9	7.9	
10*	60.7	5.3	liquid
	60.7	36.8	vapour
11	30.0	41.3	
12	19.7	59.4	
13	62.0	2.4	·

* Feedplate

COL	UMN	D	ΑT	A
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		and the second se	
Plate No.	Temp Deg C	Avbty Change kJ/s	Comment
1	30.0	-0.036	Condenser
2	35.1	-0.109	
3	43.8	-0.101	
4	53.8	-0.062	-
5	60.7	-0.150	Feedplate
6	65.4	0.107	2.2kW

PREHEATER-PRECOOLER-BOTTOM HX CYCLE

FIGURE 3.19

.6 .56 30 DEG C AMBIENT .52 .48 .44 38 DEG C AMBIENT .4 .954 .966 .978 .942 .99 .93 DISTILLATE CONCENTRATION

Fig 3.20

EFFECT OF DISTILLATE CONCENTRATION ON COP

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3.7 (Complete print out Appendix H, figure H10).

First law COP values from tables 3.4 and 3.7 plotted in figure 3.22 indicate that the R12 R113 COP values are higher for all conditions tested. Taking a specific case at a 5 deg C evaporator temperature and a 30 deg C ambient temperature the R12 R113 Carnot COP was higher by 0.113 (2.227-2.114) but this dropped to 0.087 (0.536-0.449) for the final COP value. This full Carnot improvement of 0.113 was not carried through to the final value because the R12 R113 system had higher column and absorber irreversibilities than the R11 R22 combination. Nevertheless this 0.087 represents a worthwhile improvement in the maximum achievable COP. It should be noted that this Carnot COP increase was achieved at the expense of a boiler temperature increase from 101.1 deg C to 105.9 deg C.

Examination of a large number of mixtures may reveal a relationship between fluid properties and COP and lead to an even greater improvement in COP than the one shown here.

3.2.11 Summary of the ideal cycle

For all of the system configurations considered the column and absorber mixing irreversibilities dominate and therefore appear to offer the largest potential for COP improvement. However, a closer investigation reveals that these column irreversibilities are unavoidable and therefore, other than with a structual change, system improvement is difficult. Overall these unavoidable irreversibilities result in a maximum attainable COP far below the Carnot value. This is illustrated in figure 3.23 where, other than a minor variation due to reflux in excess of the minimum, the plot

PARAMETERS FOR FIGURES 3.21 AND 3.22 PLUS TABLE 3.7

Ambient air temperature 30 deg C Boiler temperature 105 deg C Bottom mass concentration 0.05 Distillate mass concentration 0.99 Reflux ratio nominally 20% above the minimum Rl2 and Rl13 mixture

	Irreversibility terms included in equation 3.1
CASE A	Column
CASE B	Column and absorber mixing
CASE C	All losses

TABLE	3.7: COP	values	for R12	& R113	
evap temp	0	5	10	16	
deg o		0	10	15	20
Carnot	1.822	2.227	2.833	3.845	5.867
CASE A	1.082	1.265	1.413	1.62	1.95
CASE B	0.653	0.713	0.748	0.773	0.808
CASE C	0.45	0.536	0.602	0.627	0.732
lst law	0.536	0.616	0.672	0.720	0.779



Bottom flow

PREHEATER-PRECOOLER-BOTTON HX CYCLE

FIGURE 3.21

SYSTEM DATA

No	Temp Deg C	Avbty kJ/kg	
1	105.9	3.7	
2	29.9	-1.0	
3	30.1	-1.2	
4	30.0	37.9	
5	5.0	38.8	
6	6.4	26.9	
7	30.0	2.4	
8	29.9	2.6	
9	53.3	2.2	
10*	60.9	2.7	liquid
	60.9	27.7	vapour
11	30.0	24.6	
12	15.2	39.3	
13	61.8	-1.0	

*Feedplate

COLUMN DATA

Plate No.	Temp Deg C	Avbty Change kJ/s	Comment
1	30.0	-0.006	Condenser
2	31.1	-0.018	
3	33.6	-0.024	
4	38.7	-0.042	
5	47.1	-0.049	
6	55.7	-0.028	
7	60.9	-0.122	Feedplate
8	66.8	0.157	3kW .
9	76.0	0.108	3kW
10	89.7	0.048	3kW
11	106.0	2.814	Boiler 16.2kW



COP COMPARISON FOR R11-R22 AND R12-R113 MIXTURES

EVAPORATOR TEMPERATURE To DEG C

Fig 3.22

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g

for each system represents the maximum CDP theoretically attainable for a specified set of parameters.

Primarily the unavoidable minimum reflux losses are brought about by mixing losses which depend on the column flow rates and the difference between liquid and vapour availability values. Therefore not only system configuration but also operating parameters have an effect on COP and two such parameters are reflux ratio and distillate concentration.

The minimum reflux flow is determined by fluid properties and the feed condition but the excess flow above the minimum is determined by allowable column size. As the excess reflux is reduced column size increases. The distillate concentration limit is usually determined by the temperature gradient in the evaporator. For an Rl1 R22 mixture a reflux ratio of 1.15 gave 14 ideal plates whilst a distillate concentration of 0.93 resulted in a 5 degree evaporator gradient. Whilst COP gains from variations in distillate concentration and reflux flow are worthwhile, say of the order of 10 percent, they do not significantly reduce the gap between the attainable COP and the Carnot COP.

From the COP point of view increasing the bottom concentration from 0.05 to 0.13 for the R11 R22 system made no measurable difference and therefore 0.05 was adopted as the standard test concentration. Bottom concentrations below 0.05 were not investigated.

The heat rejection temperature, or in this case the ambient temperature, has a significant effect on performance. For example a change in ambient temperature from 30 deg C to 38

LEGEND FOR FIGURE 3.23

 Carnot
 Basic cycle
 Basic+preheater
 Basic+preheater+precooler
 Basic+preheater+precooler +bottom column
 Basic+preheater+precooler +bottom column+top column

۳.

PARMETERS FOR FIGURE 3.23

30 deg C ambient

101 deg C boiler

0.05 bottom concentration

0.99 distillate concentration

R11/R22



COP VERSUS EVAPORATOR TEMPERATURE

EVAPORATOR TEMPERATURE To deg C

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deg C for an R11 R22 system at an evaporator temperature of 5 deg C reduced the COP from 0.54 to 0.43, a 20 percent reduction.

The 10 percent COP improvement of R12 R113 over R11 R22 suggests that mixture properties are important and that more combinations should be considered.

3.3 A practical cycle

Having considered an ideal cycle it is now opportune to establish both ambient and internal system conditions for a practical cycle.

Using design data from the Australian Institute of Refrigeration Air-conditioning and Heating (AIRAH) for Adelaide the following design conditions have been chosen

Summer outdoor design dry bulb temperature 38 deg C Winter outdoor design dry bulb temperature 3.5 deg C Indoor design air dry bulb temperature 24.0 deg C

3.3.1 Cooling

Ten kilowatts has been arbitrarily selected as the system cooling capacity but the 15 deg C air temperature leaving the cooling coil has been based on AIRAH data (1). Allowing an 8 degree approach between distillate into the evaporator and air leaving (33) results in a distillate evaporator entry temperature of 7 deg C. Often, for mechanical vapour compression systems using the common refigerants, approximately 4 degrees of superheat at the evaporator exit is required by the refrigerant flow control device. However, for this absorption system a more sophisticated

control system is expected and therefore only 1 degree of superheat will be used. This allows maximum use of the high heat transfer coefficients experienced with boiling fluids.

Therefore, by assuming a distillate concentration of 0.99 with 1 degree of superheat, a 1 degree gradient and a 7 deg C entering temperature the vapour leaving the evaporator would be at 9 deg C. This is considered workable for an air temperature entering at 24 deg C and leaving at 15 deg C. (Assuming percent relative humidity requirements can be met by coil selection).

Pressure drop, and for that matter operating temperature, is selected largely on an economic basis and therefore the optimum for mechanical vapour compression systems may not be applicable to absorption systems. Because of this instead of selecting a pressure drop based on mechanical compression systems a range of values was tested.

No facility has been programmed to allow for condenser pressure drop. With regard to condensing temperature a 10 degree approach between air entering and distillate leaving was used. Mechanical vapour compression systems may use a 15 degree condenser approach but in this case, based on prototype performance (see section 3.4), it was limited to 10 degrees with one degree of subcooling.

In an effort to avoid cavitation at the pump 5 degrees of subcooling was allowed at the absorber out and, as a starting point, an initial approach value of 5 degrees was used between air entering and the feed flow. Again this selection was influenced by prototype performance.

Similarly 5 degree approaches were allowed on all other heat

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exchangers. It should be noted that the column heat exchangers are specified by setting values between the end of the heat exchanger and the adjacent column plate and not by an approach setting for each plate.

In the first instance all pressure drops were set to zero and figure 3.24 shows the result for an R11 R22 mixture with preheater, precooler and stripping section heat exchangers. Appendix H, figure H11 shows a complete computer model print out. If the 38 deg C COP value of 0.428 from table 3.6 is used for a comparison between the ideal and practical cycle then there is a 30 percent drop to 0.3 for this practical system.

To test the sensitivity of COP to some of the parameters each was varied with the others constant and the results are presented graphically in figure 3.25. All the temperature variations have been plotted such that the slope represents the COP change per degree.

The variation in the evaporator approach plot is caused by a change in the number of column plates from 14, for the 0 to 4 degrees span, to 13 for the 5 to 8 degree section.

Ambient temperature was found to be the most important variable (steepest slope) and the precooler approach the least. The variation in the evaporator plot makes it difficult to assess its slope, however, it can be said to be approaching that of the ambient. This suggests that variables which change the pressure ratio of the system are of equal importance. eg ambient or evaporator approach. Maintaining the condenser condition and reducing the



PREHEATER-PRECOOLER-BOTTOM HX CYCLE

FIGURE 3.24

PARAMETERS FOR FIGURE 3.24

Ambient temperature 38 deg C Bottom concentration 0.05 Distillate concentration 0.99 Rll and R22 mixture Air entering evaporator 24 deg C Air leaving evaporator 15 deg C Evaporator superheat 1 degree Condenser subcooling 1 degree Absorber subcooling 5 degrees Heat exchanger approaches **Evaporator 8 degrees** Condenser 10 degrees Absorber 5 degrees Preheater 5 degrees Precooler 5 degrees Stripping section 5 degees

SYSTEM DATA

Ng	TEMP DEG C
1	127.9
2	47.7
3	48.1
4	48.0
5	7.0
6	8.7
7	43.0
8	42.7
9	96.7
10	103.9
11	43.0
12	31.3
13	109.0

LEGEND FOR FIGURE 3.25

	Approach evaporator: Air off evaporator to refrigerant in.
	Each degree represents a degree of approach For example at zero degrees, air off and refrigerant in are at the same temperature
	Approach absorber: Air on absorber to feed leaving.
	Each degree represents a degree of approach
·· — ·· — ·· —	Ambient temperature variation:
	The 10 degree value represents a 38 deg C ambient, therefore 4 degree represents 32 deg C.
	Approach Preheater: Feed flow in to bottom flow out.
	Each degree represents a degree of approach.
	Approach Precooler: Liquid in to vapour out.
	Each degree represents a degree of approach.
NOTE: The the p	X axis represents the chosen conditions for ractical cycle viz;
	10 deg approach on evaporator 8 deg approach on absorber 5 deg approach on all other heat exchangers 5 deg C evaporator 38 deg C ambient



PRACTICAL CYCLE: COP VERSUS TEMPERATURE VARIATION

TEMPERATURE CELCIUS DEGREES

FIGURE 3.25

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absorber temperature results in an increase in COP but at a lesser rate than for the overall ambient reduction.

Pressure drops were tested on the evaporator, absorber and feed pipework. Of these the evaporator was the most important with a COP reduction of 0.001 for each 1 kPa drop.

A variation of 0.09 in CDP can be expected per 0.1 change in the reflux ratio (figure 3.26). This is considered significant and highlights the importance of controlling reflux flow.

Using the R12 R113 mixture on the practical cycle gave a COP of 0.327 which is a 9 percent improvement on the 0.3 value of R11 R22.

Use of the rectification heat exchanger with the practical cycle increased the R11 R22 COP from 0.3 to 0.32 and for the R12 R113 value from 0.327 to 0.366, both increases are considered significant. On this basis testing of the rectification heat exchanger is recommended.

In summary the effect of preheater and precooler approaches is small in comparison with temperature variations which cause a change in the difference between condensing and evaporator conditions. For example, with R11 R22 at a given evaporator condition a COP change of approximately 0.015 can be expected for each degree change in ambient. Optimum pressure drops are best established on an economic basis and the same can be said for reflux ratio. Mixture selection is important and use of the rectification heat exchanger is recommended.



Practical system: COP versus Reflux Ratio

FIGURE 3.26

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3.3.2 Heating

Some of the practical cooling cycle parameters were varied for the heating cycle. For the sake of uniformity the evaporator capacity was maintained at 10 kW with one degree of superheating but the approach was reduced because one of the problems encountered with air to air heat pumps is the fouling of the outdoor heat exchanger with ice. Selecting a large heat transfer surface is therefore desirable and so if a 5 degree approach is allowed with an ambient of 3.5 deg c, an air leaving temperature of 0.5 deg C results if the evaporator temperature is -4.5 deg C.



It is envisaged that the heat rejected from both the condenser and the absorber will be used for heating purposes and so the air entering these heat exchangers will be at the return air condition of 24 deg C. If a 10 degree approach is allowed with 1 degree of subcooling a condenser (and absorber) temperature of 35 deg C results. This allows for an air leaving temperature of say 32 deg C which is considered adequate for a comfort heating situation.

In addition to zero pressure drops throughout a 5 degree approach was used for the precooler, preheater, and both column heat exchangers.

For these conditions a heating COP of 1.32 (Cooling COP = 0.336) was obtained for R11 R22 and 1.39 (0.402) for R12

R113. Whether or not these systems are viable at these COP's depends on economic considerations which should include such factors as fuel costs, manufacturing costs, boiler input efficiency and the COP variation due to changing load requirements on the plant. It should be noted that for Adelaide a heating plant would spend a significant proportion of its time operating at less demanding conditions than have been chosen here and therefore an annual average COP could well be higher than the 'spot' values shown.

3.4 Prototype simulation

In the late 1970's Dr J. R. Roach built and tested the first prototype R12 R113 organic absorption cycle with a circuit as shown on the work sheet in figure 3.27. Currently the computer model is not designed to simulate this cycle which included a preabsorber heat exchanger. However, if the mixing/preheater/preabsorber/absorber is considered collectively then an approximate simulation is possible. Data for the program was set up as follows: Pressure drop

> - Evaporator out to absorber out was 35 kPa total. In the absence of detail about the components the following was assumed

Evaporator to precooler	zero
Precooler	6 kPa
Precooler to mixing	10 kPa
Mixing	zero
Mixing out to absorber in	10 kPa
Absorber	9 kpa

(Note: pressures on test sheet are gauge)



Subcooling and superheating

 Values on the actual cycle were estimated by comparing the measured temperatures with saturation temperatures corresponding to the measured pressure.

Absorber 301 kPa at 0.29 gives 24.7 deg C Subcooling = 24.7-22.4 allow 2 degrees Condenser 726 kPa for R12 gives 29.5 deg C Subcooling = 29.7-29.5 allow 0 degrees Evaporator 336 kPa for R12 gives 2.8 deg C Superheating = 24.8 - 3.8 = 21 deg

(allow for a one degree gradient)

Approach values

Absorber 22.5 - 21.4 = 1.1 degree Condenser 29.7 - 21.4 = 8.3 degrees Evaporator 23.3 - 2.8 = 20.5 degrees Precooler 25.7 - 24.8 allow 1 degree Preheater 35.7 - 35.4 allow 1 degree Column, Bottom heat exchanger

Feed plate to heat exchanger out

assume 3 degrees

Bottom plate to heat exchanger in

assume 8 degrees

Evaporator capacity = $(27.5 - 23.3) \times 0.349 = 1.466$ kW Given the above parameters the computer model was run so that the boiler temperatures coincided with the measured value of 115.4 deg C. For ease of comparison the data below have been taken from appendix H, figure H12:
	measured	predicted
Power input kW	2.395	2.528
Condensing pressure kPa	726	733
Feed temperature deg C	50.8	53.5
Bottom Heat Exchanger leaving column deg C	65.3	65.1
Precooler liquid out deg C	25.7	29.3
Bottom flow kg/s	0.0265	0.0312
Feed flow kg/s	0.0367	0.0417
Distillate flow kg/s	0.0102	0.0105
COP	0.606	0.577

Even though these results are considered encouraging checks with other tests have not been conducted because of the different system configuations.

This first prototype has now been dismantled and a second, with the same configuation as the program, has just been completed.

3.5 Comparison with a Lithium Bromide/water system.

Yazaki Australia Pty Ltd market a range of absorption chillers including a gas fired unit model CH600, the typical operating characteristics (37) of which are given in the table below. The R12 R113 system was modelled using all the Yazaki operating conditions except for the preheater and absorber approaches. It was considered reasonable to have an absorber approach of the same order as the condenser and evaporator and so the preheater approach was reduced from 15.1 degrees to 5 degrees. The Yazaki absorber approach was reduced from 6.6 degrees to the prototype value of 2.0 degrees.

	Yazaki	R12-R113
Cooling capacity kW	6.87	6.87
Chilled water in deg C	11.9	11.9
Chilled water out deg C	6.8	6.8
Evaporator temperature deg C	4.5	4.5
Cooling water deg C	29.2	29.2
Condenser approach degrees	2.4	2.4
Absorber approach degrees	6.6	2.0
Evaporator approach degrees	2.3	2.3
Precooler approach degrees	-	2.0
Freheater approach degrees	15.1	5.0
All column HX approach degrees	5.0	5.0
COP	0.745	0.635
Heat input to generator kW	9.22	10.757
Feed flow/distillate flow ratio	14.5	4.42
Distillate flow rate kg/s	0.0029	0.0493
Column feed flow rate kg/s	0.0421	0.2177
Heat rejected in condenser kW	7.45	10.349
Heat rejected in absorber kW	8.64	7.298
Pump power kW	0.0	0.058
Distillate mass concentration	1.0	0.99
Feed mass concentration	0.58	0.263
Bottom mass concentration	0.54	0.05
Feed temperature deg C	70.6	59.1
Bottom temperature deg C	87.0	108.0
Pressure ratio	11.75	2.17
Highside pressure kPa	9.87	770.8
Lowside pressure kPa	0.84	355.0

Compared with the Lithium Bromide/water system the R12 R113 variation is:

> COP 15% lower Distillate flow 1600% higher Feed flow 417% higher

Two factors contributing to the superior performance of Lithium Bromide are

- The high latent heat value of water. It is 1600% higher than R12. (See section 3.2.5)
- Reduced difference between high and low pressure sides of the system.

For the R12 R113 system COP to rival the Lithium Bromide/water system under these conditions a reduction in column and/or absorber irreversibilities is necessary. A preliminary investigation indicates that these reductions may be achieved with a multistage cycle. If the organic system is significantly cheaper to produce it may be competitive from an economic point of view even with the lower COP and/or a more complex cycle.

Space heating for the Lithium Bromide system is achieved not by using the system as a fluid to fluid heat pump but by boiling refrigerant out of solution with the gas flame and then condensing it in what was previously the evaporator. The condenser and absorber are rendered inoperative and the condensed refigerant is absorbed and circulated back to the boiler. Unlike the organic system this mode of operation cannot produce a COP greater than one and therefore no attempt has been made to compare heating cycle performance.

CHAPTER 4

Nomenclature

a,b,c = constants

f = fugacity

O = pressure

 $\mathbf{R} = \mathbf{residual}$

T = temperature

N = number of distillate column plates

Z = compressibility

 $\tau =$ convergence criterion

Subscripts

i and j = component k = iteration number l = liquid v = vapourc = critical

COMPUTER MODEL

To analyse the absorption refrigeration cycle it was decided to write a VAX 11 Fortran computer model which provided the following:

- A comparison between calculated and published fluid properties for a range of working fluids.
- Easy selection of working fluids.
- Tables of fluid properties for both pure substances and binary mixtures.
- Optional use of various internal heat exchangers in the refrigeration cycle.
- Specification of refrigeration cycle operating
 variables such as evaporator temperature.
- A system analysis showing fluid properties at various points in the cycle as well as cycle operating characteristics such as COP.

The program was written in stages, firstly the fluid property calculation section, then the property checking section and finally the absorption cycle model hence the extensive use of subroutines. These subroutines allowed sections of the program to be run independently and also simplified the location of errors. Consequently the main program contains only an input facility and some minor calculations followed by the appropriate subroutine calls.

4.1 Program structure and ancilliary programs

A single input file contains variables for selecting program options such as tables of fluid properties and also for specifying system model values such as temperatures and heat

exchanger options. As shown in the file structure in Appendix C it is in this file that the two operating fluids are specified by ASHRAE number. Once the two fluids are specified their properties are read from individual data files. Therefore, prior to running the system model separate input files must exist for each pure substance and a value for the Peng-Robinson equation interaction coefficient must have been entered into the program.

The program may be imagined to have 3 levels, the first being the main program which reads and checks input data and also calculates various Peng-Robinson constants. Next are the system and property table subroutines and, finally, the individual calculation subroutines. The system and property table subroutines contain the information required for the writing of the output files such as the system layouts shown in Appendix H and, in the case of the mixture properties, a plot file is written for the enthalpy concentration diagrams as shown in appendix G.

In all there are three ancilliary programs which support the main program but these are not essential for its operation viz: a plotting program (32), a batch program and an interactive program for writing the main input file.

The plot program may use the file generated by the main program as an input to generate enthalpy concentration diagrams. All graphics are produced using external IGL subroutines.

No keyboard input is required during execution of the main program and therefore it can be run in the batch processing

mode. In fact most of the system modelling used in chapter 3 was done via an overnight batch program which modified the main input file after each run and then returned the program to the queue for the next execution.

For single runs of the main program an interactive program which reads the current input file, prompts for the changes required and then writes a new version has been created. This avoids having to memorise the input file structure.

For portability the main program was written to stand alone and therefore numerical method subroutines were included. Overall the program occupies 240K bytes and execution on the VAX 11/780 varies between 2 and 5 minutes of processing time depending on the number of heat exchangers in the circuit.

4.2 <u>Numerical Methods</u>

Early in the development of the program it was found to be necessary to write some standard subroutines for numerical methods for the solution of equations and so the solution of the Peng-Robinson cubic was used as a test case. The methods considered were direct substitution, half interval, regula falsi (or linear inverse interpolation) and Newton's method (9). Three were tried and typical results for the cubic solution were

Half interval	20	****	20	iterations
Regula falsi	6		9	iterations
Newton	3		5	iterations

From this a set of priorities for methods was developed which rated Newton's method first, but of course this was

only available for analytically differentiable functions, followed by regula falsi, half interval and direct substitution. It transpired that Newton's method was confined to the solution of the Peng-Robinson cubic because it was the only analytic function and elsewhere, because of speed and reliability, the regula falsi method was used. In some cases two methods were combined. For example it is possible to obtain the two guesses required to start the regula falsi method by supplying one guess and using direct substitution to find the second.

A robust subroutine was written for the regula falsi method by incorporating a similar method called false position and by storing three successive values of the function. False postion was included because the positive and negative residual required by regula falsi for the first two guesses could not always be guaranteed. In the case of false position both residuals can have the same sign but during computation one of the guesses remains stationary while the other is updated. For true regula falsi (9) the two initial guesses must have different signs and then both are updated. The equation used for both methods is:-



Three values of x and their corresponding residual values are required by the subroutine. The flow chart is as follows:



return

return

For the first call, which involved the two initial guesses, values X1 and X2 were made equal to guess one and X3 equal to guess two. This procedure will run for initial pairs of residuals with either the same signs or opposite signs.

4.3 Fluid properties

Peng-Robinson equation subroutines were written to generate basic data such as pure and mixture enthalpy values and then these in turn are called by the subroutines which establish working fluid conditions like vapour-liquid equilibrium.

4.3.1 Peng-Robinson equation

A section in the main program calculates all constants for the Peng-Robinson (chapter 1) equation of state, including those sections of the enthalpy, entropy and availability computation which remain constant throughout. These values are transmitted via a common block.

The solution of the Peng-Robinson cubic in Z (eqn 1.2) involves three meaningful possibilities :-

vapour only one real root

liquid only one real root

Two phase three real roots, the smallest for liquid and the largest for vapour.

For a given temperature and pressure the number of roots and the stationary points are determined by setting the first derivative of equation 1.2 to zero. The form of the equation is

 $aZ^2 + bZ + c = 0$

The number of real roots can be obtained by evaluating the discriminant of the quadratic formula

$$Z = \frac{-b \pm \sqrt{b^2 - 4ac}}{2a}$$

discriminant = $\sqrt{b^2 - 4ac}$

If the discriminant is

Knowing the stationary points and the number of solutions a single half interval calculation using two limiting values is used to find a starting value for Newton's method. The limiting values are:-

One root



Two roots

Liquid: zero and the lower stationary point Vapour: Upper stationary point and 1.1



For the solution of the cubic a residual of less than -12 1 x 10 is required. The program terminates if the number of iterations exceeds 10 or if Newton's method yields a guess outside the starting limits. Generally this method was found reliable although even with calculations in double precision it can fail near the critical point if the value of the slope approaches the accuracy limit of the machine. However, this is not a problem because failure always occurs inside the 'dead zone' (see chapter 1, end of section 1.2).

This solution of the cubic is the most frequently used section of the program and therefore a reduction in its computation time could represent a significant reduction overall. Further development of the program should include a comparison with an analytical method.

4.3.2 Pure Substance Vapour-liquid equilibrium

For a given temperature and substance the saturation pressure is established by the convergence of

absolute =
$$|P_{guess}\left(1-\frac{f_v}{f_L}\right)| < 5Pa$$



RETURN all thermodynamic properties:

RETURN all thermodynamic properties



4.3.3 Mixture vapour-liquid equilibrium

Given two properties the vapour liquid equilibrium data for a mixture can be established. Several subroutines have been written which establish mixture vapour-liquid equilibrium and for these the input variable combinations are:-

- temperature and pressure
- pressure and liquid mole concentration
- temperature and liquid mole concentration
- pressure and vapour mole concentration
- temperature and vapour mole concentration

These subroutines all follow a similar pattern and are based on the equilibrium requirement that the fugacity of a component in the liquid phase must be equal to the fugacity of that component in the vapour phase. Typically the convergence criterion used in these subroutines is based on

$$1 = x_1 \frac{\bar{f}_{v1}}{f_{L1}} + x_2 \frac{\bar{f}_{v2}}{f_{L2}}$$
(4.2)

ie. when the pure liquid and vapour fugacities are equal then the sum of the mole fractions of the components equals one. Only the flow diagram for the temperature/pressure subroutine is shown in figure 4.3 because the other combinations follow a similar pattern. In this case the convergence criterion for the liquid mixture is based on the difference between the guessed and calculated mole fractions of the components. Equation 4.3 forms the basis of the liquid mole fraction calculation:



$$\frac{y_i}{x_i} = \frac{\frac{f_{iL}}{x_i}}{\frac{f_{iV}}{y_i}}$$

Checks, not shown in the flow chart, include

- that the temperature specified lies between the pure component saturation temperature for that pressure.
- that the temperature does not exceed the critical temperature of either of the components.
- the number of iterations does not exceed 30.

4.3.4 General mixture subroutines

Two subroutines have been written which generate mixture properties for any state given three variables. One requires inputs of pressure, concentration and enthalpy and the other pressure, concentration and availability. These two subroutines call various combinations of the mixture subroutines mentioned in section 4.3.3 and also follow the same flow pattern of an initial guess from the program, a second guess from direct substitution and all subsequent predictions from regula falsi.

4.4 Fluid property tables

Since the refrigeration system model's accuracy depended on fluid property data the production of fluid property data tables was a secondary consideration to the checking of calculated data against published values.

(4.3)

4.4.1 Fure property tables

Saturated temperature values are calculated at 10 degree intervals between the nominated enthalpy datum and the critical condition. Variables to specify are

- Substance: nominated by refrigerant number eg R12

- Units : SI or Imperial

and the output is in tabular form as shown in appendix F. Options include superheated vapour data at 10,20,30 and 40 degrees of superheat from each of the saturated values and a comparison between calculated and published values from the input file (appendix E).

4.4.2 Mixture properties and enthalpy concentration diagrams

Calculation of mixture data is centred around the production of enthalpy concentration diagrams. At the pressure in question equilibrium data is calculated at 10 degree intervals between the saturation temperatures of the two pure components. Superheated values are calculated along isotherms using the vapour solution of the Peng-Robinson equation at the temperature and pressure in question. No subcooled solutions for the Peng-Robinson equation are used. All subcooled calculations are treated as saturated ones at the temperature and concentration in question but at a different pressure. It has been assumed that liquid properties are independent of pressure because the Peng-Robinson equation fails at varying distances (depending on temperature) out into the subcooled region. ie the effect of small pressure changes on liquid enthalpy is ignored. Variables to specify:-

- Components : by refigerant number

- Pressure : either by a pressure value or by the saturation temperature for the most volatile component in its pure state.
- Concentration : Intervals for calculation, down to 0.01

 Number of isotherms : Number of isotherms above and below the equilibrium lines.

Output is in tabular and graphic form although only the graphics of the enthalpy concentration diagram are shown in appendix G.

4.5 Absorption system

4.5.1 Overall sequence

A sequential modular approach (9) was used to analyse the absorption cycle and the sequence used along with the corresponding listing is shown in figures 4.4 and 4.5 respectively. Iterative procedures were necessary in the evaporator and column sequences as shown.

Consider the evaporator. Here two factors interacting make a loop necessary. The pressure in this section of the system is determined by the saturation condition in the evaporator. Guessing the evaporator outlet condition allows the evaporator inlet condition to be calculated via the precooler energy balance. This inlet condition must then agree with the temperature calculated from air leaving the evaporator minus the evaporator approach value.

The column loop is necessary because at the outset the program has too many constraints. It was decided to adopt the approach of specifying too many variables and then modifying one to suit so that program would yield the nearest possible solution to the overall requirement. In



FIG 4.4

Sec	uence mber -	Condition	
	1	Condenser out	
	2	Condenser in	
	3	Precooler liquid in	
	4	Evaporator out	
	5	Precooler vapour in	
	6	Precooler vapour out	
	7	Mixing in NEW APPROACH	
	8	Precooler liquid out	
	9	Distillate throttle in	
	10	Evaporator in Check calculated evaporator inlet value with value given by T _{air} leaving minus evaporator approach.	
	11	Absorber out : This gives feed concentration Calculate bottom concentration	
	12	Pump in	
	13	Pump out	
	14	Feed split	
	15	Preheater feed in	
	16	Rectification Heat exchanger in	
	17	Boiler out	
	18	Bottom heat exchanger in	
	19	Guess feed plate temperature	
	20	Bottom heat exchanger out	
	21	Bottom flow into preheater	
	22	Bottom flow out of preheater	
	23	Feed out of preheater	
	24	Feed out of rectification heat exchanger	PLATE
	25	Feed join	IEMPERATURE
	26	Feed condition	
		Column : No heat exchangers	Î
		Calculate condenser down to feed plate	
		(on first run only check feed	
		temperature against 19, loop back if	
		necessary) NEW R	
		Feed plate to boiler (gives number of plates = N) RAT	
		Use plate temperatures to locate column heat exchangers	
		Column : With heat exchangers NEW HEAT	
		Calculate condenser down N plates	
		Check location of neat exchangers NU	
		Column concentration against II : NU	
		Column energy Dalance	
		Check feedplate temperature against 19 ; ND	
	27	Bottom flow into throttle	
	28	Bottom flow into mixing	
	29	Mixing out Fig 4.5	
	20	Absorber in	

keeping with this the bottom concentration was calculated via a specified feed flow/distillate flow ratio and then again from the reflux flow via the column analysis. The reflux flow was modified until the two bottom concentration values converged and generally this modified the original value of the order of five percent. The explanatory notes relate to the sequence numbers in figure 4.5:

No	comment		
10	Required residual is < 0.1 degrees. The first guess		
	for the approach is a fixed value, the second is from		
	direct substitution and finally regula falsi is used.		
13	Constant enthalpy assumed across the pump.		
19	Initial feed temperature guess uses a simple linear		
	interpolation between the saturation temperatures of		
	the two pure components and is based on the bottom,		
	feed and distillate mass concentrations. viz		
	feed concentration - bottom concentration		
	distillate concentration - bottom concentration		
26	First time through the calculated feedplate temperature		
	is checked against the first guess. If the difference		
	is > 0.5 degrees control is returned to 19 and the new		
	calculated value is used. From then on only the full		
	column loop can modify the feedplate temperature value.		
	Reflux ratio is varied so that the bottom concentration		
	calculated from the column coincides with the		
	concentration from 11 by <= 0.0005 . Plate		
	temperatures determine column heat exchanger location		
	and heat transfer. Adding the heat exchanger modifies		
	the plate temperatures making a check of new plate		
	conditions necessary.		

All input variables required for modelling are contained in a single input file (appendix C). Data required includes

- Substances

- Enthalpy temperature datum
- Availability pressure datum
- Availability temperature datum
- Cooling fluid temperature onto condenser
- Condenser subcooling
- Approach: distillate out to fluid temperature on
- Cooling fluid temperature onto absorber
- Absorber subcooling
- Approach: feed out to cooling fluid temperature entering
- Evaporator capacity
- Fluid temperature leaving evaporator
- Evaporator superheating of distillate
- Approach: Fluid temp leaving to distillate entering
- Distillate mass concentration
- Feed flow/distillate flow ratio
- Ratio of actual to minimum reflux flow
- Pressure drops (various)
- Precooler approach: liquid in to vapour out
- Preheater approach : feed flow in to bottom flow out
- Bottom heat exchanger approach: HXr out to feed plate
- Bottom heat exchanger approach: Bottom plate to HXr out
- Rectification HXr approach: HXr out to feed plate
- Rectification HXr approach: Top plate to HXr
- Percentage of feed flow through preheater

Program control data specifies system configuration by allowing independent selection of heat exchangers *:

Preheater

Frecooler

Column heat exchanger, stripping section Column heat exchanger, rectification section and also output by controlling the amount of data in the print out.

* A feed boost heat exchanger was programmed but not tested.

Typical print outs are shown in Appendix H.

4.5.2 Column calculations

Two methods were tried for the solution of the column, a rigorous approach by Henly et al (19) and a computer model of the Ponchon Savarit construction. Henly's method for column analysis is based on equation partitioning and tearing methods. 'This method requires an initial guess of temperature and vapour flow rate for each plate and then, for subsequent guesses, Henley recommends direct substitution with damping. With regard to damping a 10 percent maximum variation in temperature and flow was suggested. The method was easy to program but it was too sensitive to the initial guesses to be reliable.

The initial guess for the vapour flow rates on each stage was set equal to the flow entering the condenser (as suggested in the text). Linear interpolation between condenser and boiler temperatures was used for the plate temperature guesses but after unreliable performance these were replaced with Ponchon Savarit values. Even then the

method occasionally failed. Damping did not guarantee success and when regula falsi was used to predict subsequent plate temperatures the method failed completely.

Difficulty was also experienced with the convergence criterion. Henley suggested that a criterion based on temperature and vapour flow was unnecessarily complicated and that the following expression, based on temperature alone, would suffice.

$$\tau = \sum_{j=1}^{N} \left[T_j^k - T_j^{k-1} \right]^2 \le 0.01N$$
(4.4)

Indeed this was found to be the case. The vapour flow values did converge more quickly than the plate temperatures and so equation 4.4 was used. However, it was discovered that this limit of 0.01N demanded a convergence between subsequent temperatures calculated for each plate of less than one degree. Assuming the number of plates (N) is 12 then the sum of the squares of the temperature differences between iterations k and k-1 must be ≤ 0.12 . If this difference is assumed equal for all plates then this criterion requires a difference in temperature between successive guesses of 0.1 degrees. This means that loops used to find fluid properties must have a convergence criterion less than 0.1 otherwise variations in fluid properties cause unstable operation. The overall accuracy of the program does not warrant such a tolerance on temperature and so the criterion was increased to 1.0N. However, by relaxing the value to 1.0N (a 1 degree difference per plate) allowed a convergence situation where values were within a fraction of one degree on most plates

but 2 or 3 degrees for another. It was not possible to embark on an detailed investigation of this method and so it was replaced with the Ponchon Savarit method.

Despite the failure of this method two benefits did result from this work. One was a check on the Ponchon Savarit column calculations. When plate temperatures from successful runs of Henley's method were checked against Ponchon Savarit values they were at most two degrees different. Secondly it was found that by using Ponchon Savarit temperatures and enthalpies in an adapted version of Henley's method plate flow rates could be easily calculated.

Column temperatures and concentrations are determined using a Ponchon Savarit construction procedure (figs 4.6 and 4.7). In the case of the column with heat exchange the same procedure is used except that the difference point c (and d) is progressively stepped down (d goes up) for each plate with a heat exchanger. Protection in the procedure includes checks for

- Excessive number of plates (100 maximum)

- Azeotrope

- Excessive heat exchange in the column

- Excessive number of iterations (most are set at 30) By knowing plate temperatures as well as liquid and vapour concentrations it is possible to calculate plate flow rates. As mentioned Henley's rigorous method required the adjustment of two variables, temperature and vapour flow rate. It was found that by using Ponchon Savarit temperatures and varying only vapour flow rate all the flows could be determined. The sequence starts with all the vapour flows being set at the condenser flow

PONCHON SAVARIT PROCEDURE

Number	Operation	Known values	method
1	Calculate minimum difference point (b)	Feed concentration Feed enthalpy Distillate concn	Guess temperatures and compare slopes F-b1 and b2-b1
2	Actual difference point (c)	Ratio of actual to minimum reflux flow	
2	Stripping section difference point (d) (Principle operating line)	Bottom concentration difference point Feed point	
4	Locate isotherm e-f	Distillate concn Pressure Temperature	
5	Locate g (vapour leaving stage)	difference point Liquid condition (f)	Guess temperature & use ratio of sides of triangles to compare c-f1/c-g1 to f-f1/g-g1
6	Repeat 4 & 5 until the isotherm crosses the principle operating line g1-h1 (No attempt has been made to find the optimum feed plate)		
7	Repeat 4 & 5 using d instead of c until j concentration is less than or equal to bottom concentration.	,	

FIGURE 4.6



CONCENTRATION

FIGURE 4.7

value, then uses direct substitution and finally terminates when the largest difference between successive vapour flow calculations for any plate is less than one percent.

CHAPTER 5

•

GENERAL CONCLUSIONS

This chapter is divided in the same way that the project aims were grouped in the introduction; viz fluid property estimation, absorption system analysis and optimisation of system design. The requirements of the aims for the first two headings were fulfilled but the system optimization aspect was not pursued. After a substantial time investment in the areas of fluid properties and system analysis it was realized that the course requirements could be fulfilled on this basis and that an effort of similar magnitude in the optimization area could not be justified. Consequently this work provides a basis for an exhaustive investigation of absorption refrigeration cycles and their working fluids by supplying the method, in terms of the second law analysis, and the means, by way of the computer model.

Information contained in this chapter is a collection of the conclusions stated in various parts of this report.

5.1 Mixture properties

Both pure and mixture fluid properties have been predicted satisfactorily for a range of organic fluids. From this work it can be concluded that

a) The Peng-Robinson equation of state (section 1.2) can be used to predict fluid properties with sufficient accuracy for engineering calculations. Typically calculated values were within 15 percent of the published data although liquid entropy values did show larger deviations. Appendix E shows a comparison between Peng-Robinson and published data while appendix D illustrates that the Peng-

Robinson equation performs as well as, and in some cases better, than the widely used Soave-Redlich-Kwong cubic equation.

- b) Property prediction can be improved by fitting the Peng-Robinson constant K value to published data for each substance rather than using the generalised Peng-Robinson correlation based on acentric factor.
- c) For the mixtures tested the effect of the Peng-Robinson binary interaction coefficient on properties was significant. Assuming zero for this constant is not recommended as errors between calculated and published values of between 20 and 40 percent may result. This coefficient was determined by substituting published vapour-liquid equilibrium data in the Peng-Robinson equation and then working "backwards" through the calculation. In the absence of published data for mixtures the activity coefficient approach of Van Laar can be used to estimate vapour-liquid data. These values can then be substitued in the Peng-Robinson equation to establish the interaction coefficient. This was done for the R12 R113 mixture and the method is discussed in appendix B.

5.2 Absorption refrigeration cycle analysis

A computer model has been written which executes successfully for evaporator temperatures between 0 deg C and 20 deg C and for ambient temperatures between 25 deg C and 38 deg C. The model has facilities to predict and check fluid properties as well as model an organic absorption refrigeration cycle.

5.2.1 Computer model options and performance

Input variables allow selection of heat exchanger approaches, ambient temperatures, mixture concentrations, evaporator capacity and working fluids as well as individual selection of four internal heat exchangers. Output from the program includes fluid property tables and pressure enthalpy diagrams as well refrigeration cycle details. Modelling of the refrigeration cycle has produced:

- a) A simulation of an R12 R113 system prototype which gives comparable results to experimental test data. For example the measured and predicted COP values were 0.606 and 0.577 respectively.
- c) A Second Law of Thermodynamics analysis of the system (chapter 3) which shows that
 - the column and absorber mixing processes are the least energy efficient.
 - energy utilization may be improved by the use of internal heat exchangers.
 - After fitting internal heat exchangers the column irreversibilities mainly occur on the feed plate and in the boiler. For the boiler it is due to the difference in availabilities of the vapour and liquid streams in equilibrium and in the case of the feed plate there is an added loss due to the introduction of another stream with a different availability. Overall the greater this vapour liquid difference, and the higher the column flow rates, the greater is the irreversibility loss. For this reason external reflux flow to the column should be minimised to the

extent justified by economics of distillation column size. These unavoidable irreversibilities due to mixing set the maximum achievable COP at a value well below the Carnot value. These achievable COP's are closest to the Carnot values at the lower evaporator temperatures: for example for an R11 R22 system with all the internal heat exchangers, ideal conditions and a 0 deg C evaporator the achievable COP is approximately one third of the Carnot value.

Fluid selection is important. Both the Carnot
 COP and the irreversibilities depend on this.
 The R12-R113 mixture consistently produced COP
 values 10% higher than those for R11-R22,
 mainly because of its higher Carnot COP.

d) A First Law of Thermodynamics analysis shows that

- The system COP is most sensitive to variables which affect the pressure ratio (such as ambient temperature) and least sensitive to preheater and precooler heat exchanger approaches. For example with an R11 R22 mixture and a constant evaporator temperature of 5 deg C a change of 0.015 in COP can be expected for each degree change in ambient.
- As the column reflux flow reduces, the COP increases,
 but with a corresponding increase in column size.
- Reducing distillate concentration provides an increase in COP whilst introducting a temperature gradient across the evaporator. For R11-R22 reducing the concentration from 0.99 to 0.93 introduced a 6 degree gradient and a 7% COP increase.

- Increasing the bottom concentration significantly reduces the generator temperature but has little effect on COP and pump power. For an R12 R113 system of 10kW capacity an increase in bottom concentration from 0.05 to 0.13 reduced the generator temperature from 101 deg C to 82 deg C with a corresponding pump power change from 0.1kW to 0.14kW.
- e) A comparison with a commercially available Yazaki gas fired Lithium Bromide water system commonly used for air conditioning revealed a 15% lower COP for the R12-R113 system. This comparison was made at the Yazaki rating conditions for cooling which included heat rejection to cooling water at 29.2 deg C and an evaporator temperature of 4.5 deg C.
- f) Modelling of a practical heating cycle revealed COP's of the order of 1.3 for R11 R22 system operating with an evaporator temperature of -4.5 deg C. This suggests that further work is worthwhile.

5.2.2 Computer model methods

With regard to computational procedures (section 4.2) Newton's method has been used successfully to solve for the Peng-Robinson equation cubic roots. For non-differentiable functions the regula falsi method has proved to be far more robust and faster than both the direct substitution and the half interval methods.

For the distillation column analysis (section 4.5.2) a programmed representation of the Ponchion Savarit method has been used in preference to the equation partitioning and tearing variable method proposed by Henley et al (19).

Henley's method failed often because it was too sensitive to initial guesses and also it required a fluid property accuracy greater than that which was possible with the Peng-Robinson equation.

Execution for one set of refrigerant cycle conditions required 2 to 5 minutes of central processing unit time on the VAX 11/780 computer. These relatively long run times are often inconvenient when using the computer interactively and so the input was configured to enable multiple runs in the batch processing mode.

5.3 Optimum system design

An optimum design procedure has not been established because to effectively optimise absorption system design both cost and performance must be considered, not just COP as has been the case here. For example even though the organic unit COP is 15% less than for the Yazaki design it may still be cost competitive because of lower production costs. Also, unlike the Yazaki units, the organic system is capable of heat pump operation and therefore has a wider market potential. It is considered that the organic COP is sufficiently close to the already commercial Yazaki equipment to warrant further investigation

5.4 Future work

Further work is recommended and it is suggested that it be divided into four broad areas:-

1) Investigation of a range of binary mixtures with a

view to establishing relationships between fluid properties and COP. This should include a comparison with existing systems.

- 2) Simulation of prototype performance. Checks should be conducted against test measurements to establish if trends predicted by the program occur in pratice.
- 3) Investigation of multistage evaporator/absorber combinations in conjunction with two or more separate column sections is warranted. These systems offer the possiblity of reducing the column and absorber mixing irreversibilities and therefore may offer significant improvements.
- Optimization of design by assessing both costs and performance.

APPENDIX A

PENG-ROBINSON EQUATION OF STATE

Nomenclature

 $\overline{\mathbf{A}} = \operatorname{Peng} \operatorname{Robinson} \operatorname{constant}$

a = Peng Robinson constant

B = Peng Robinson constant

b = Peng Robinson constant

 $\mathbf{f} = \mathbf{fucacity}$

 $\mathbf{P} = \mathbf{pressure}$

R = universal gas constant

S = entropy

T = temperature

U = Helmholtz function

 $\mathbf{v} = \mathbf{v}$ olume

 $\mathbf{Z} = \text{compressibility}$

Subscript

 $\begin{array}{l} L = liquid \\ V = vapour \end{array}$

 $\frac{\text{Superscript}}{\text{o} = \text{ideal gas state}}$

PENG ROBINSON EQUATION

A.1 Optimization of K equation 1.7

To optimize K a decision had to be made on the comparison with experimental data with respect to :

a) Temperature range.

b) Convergence criterion for f and f.

c) Deviation parameter.

d) Property to be used as a basis for selection.

A.1.1 Temperatures

For the absorbtion cycle under consideration temperatures can range from subzero in the evaporator to in excess of 100 deg C in the distillation column. As most thermodynamic tables (3) use saturated liquid at -40deg C as the enthalpy and entropy datum condition this was chosen as the lower limit. Also it was below the lowest evaporator temperature expected. From ambient temperature considerations it was not unreasonable to expect 120-150 deg C in the distillation column. Using this data a range for the selection of K was chosen from the higher of -40 deg C and the freezing point to the lower of 200 deg C and the critical temperature. It should be noted that property data computation is not
restricted to this range.

A.1.2 Convergence

A saturated liquid-vapour fugacity convergence criterion of 5x10-3 kPa was selected because further reduction increased computing time with no change in the property values generated.

A.1.3 Deviation

Figure A1 illustrates that, based on pressure, minimums occur for equation A1 at 0.6352 and for equation A2 at 0.6381. This variation is typical for these refrigerants.

SD1 =
$$\sqrt{\frac{\sum_{1}^{n} (calcd - publd)^{2}}{(n-1)}}$$
 (A.1)

$$SD2 = \sqrt{\frac{\sum_{1}^{n} \left(\frac{calcd-publd}{publd}\right)^{2}}{(n-1)}}$$
(A.2)

Equation A1 was selected as the optimization criterion on the basis that it was biased toward the higher working end, and not the low useage end, of the temperature range. At the low temperature even though the absolute errors are small the relative error is large which increases the importance of this section for equation A2. For example, compare the two pressure percentage error curves in figure A2. Equation A1 allows the greater



EQUATIONS A1 & A2 VERSUS PR KAPPA FOR R12

PENG ROBINSON KAPPA VALUE

Fig A1



PRESSURE PERCENT ERROR VS TEMP FOR SATD R12

PERCENT ERROR IN SATD PRESSURE

error at the lower end but maintains better accuracy for Tr greater than approximately 0.7 . Appendix E2 shows, at 10 Fahrenheit degree intervals, both the values calculated from equation A.1 and the published (3) R12 saturation pressure data.

A denominator of n-1 was chosen for equation A1 to allow for the smaller sample sizes which occured when the temperature range was small.

A.1.4 Property

Equation A.3 is an indication of the relative deviation between published and calculated values (Average absolute value of relative percent deviation) making it an ideal basis for a direct comparison between properties.

$$AAD = \frac{\sum_{1}^{n} \left| \left(\frac{calcd - publd}{publd} \right) \right| 100}{n}$$
 (A.3)

Figure A3 ((a)=liquid (b)= vapour) shows R12 saturated propety AAD values for a range of K. Typically the largest errors occured in the liquid properties. With Kat 0.6352 the largest deviation of 6.1 was for liquid entropy. To improve accuracy it was proposed that liquid entropy be used for optimization. However, to optimize on a minimum for this property causes unacceptable errors. Even at the graph lower limit of K = 0.58 the error in pressure is in excess of the original maximum of 6.1 and the liquid entropy minimum has not yet been reached. Also entropy (and enthalpy) accuracy is



AND VERSUS KAPPA FOR SATURATED R12





influenced by two features:- liquid compressiblity and Specific Heat Capacity data. Since liquid compressiblity is determined by the fundamental accuracy of the PR equation any change in liquid entropy would have to come from the Specific Heat Capacity equation constants. It was not possible to embark on an investigation into the accuracy of this data and so the constants were used as published. (31)

Since enthalpy is the basis for the first law analysis of systems it was given a higher priority than entropy. Liquid and vapour enthalpy minima occured either side of the pressure minimum making pressure an acceptable comprimise. On this basis pressure was retained as the K selection property.

Table A.2 is a resume of the SD1 and AAD values. A full data set are given in appendix E.

Reft	Formula	к	SD1		AAD				Tr 1	No.				
no.			Pressure psi	Density liq	Compres liq	siblity vap	Enth	vap	Entr liq	vap	Latent Heat	Lowe:	upper	Points
11	C CI ₃ F	8.6654	0.5112	5.65	5.99	1.07	3.4	8.59	3.70	2.36	2.15	0.495	8.998	43
12	C C1 ₂ F	0.6352	0.1826	5.11	4.89	0.91	2.15	0.39	6.11	2.93	1.76	8.585	0.995	28
13	C C1 F ₃	0.6321	0.4361	4.95	5.25	1.47	3.45	0.51	10.5	2.69	3.72	0.772	0.993	13
1381	C Br F3	0.6319	8.3541	5.03	5.37	2.24	2.18	8.99	7.15	1.74	4.42	0.685	8.996	29
22	CH C1 F2	8.7072	1.25	3.98	4.48	9.88	2.37	0.64	8.41	3.37	2.45	0.632	0.993	25
113	C CI2 F-C CL F2	8.7514	8.4879	5.76	5.2	1.17	5.43	8.94	2.33	1.94	2.18	0.479	0.992	46
152a	CH3 CH F2	8.7688	.552	8.74	9.27	1.40	4.73	2.55	7.53	2.72	2.13	0.603	0.962	26
			<u>kPa</u>			÷				Ð	1			
718	^H 2 ⁸	8.8592	111.7									0.422	8.994	38
-	C ₂ C1 ₄	0.7511	78.4		No th	ermodynan	nic data	for Per	 chloroet 	hlvene		8.376	1.000	40

Table A2

A.2 Departure function and fugacity derivation

Reid et al (1977) expresses all the departure functions in terms of the Helmholtz energy (U).

$$U - U^{\circ} = -\int_{\infty}^{V} \left(P - \frac{RT}{V}\right) dV - RT \ln \frac{V}{V^{\circ}} \qquad (A4)$$

$$S - S^{\circ}_{\downarrow} = -\frac{\partial (U - U^{\circ})_{\bullet}}{\partial T}$$
(A5)

$$H - H^{\circ} = (U - U^{\circ}) + T(S - S^{\circ}) + RT(Z - 1)$$
 (A6)

$$ln\frac{f}{P} = \frac{U - U^{\circ}}{RT} + ln\frac{V}{V^{\circ}} + (Z - 1) - ln Z$$
 (A7)

substituting equation 1.1 into equation A4

$$U - U^{o} = \int_{\infty}^{V} \frac{RT}{V - b(T)} - \frac{a(T)}{V(V + b(T)) + b(T)(V - b(T))} - \frac{RT}{V} dV$$
$$= -\int_{\infty}^{V} \frac{RT \ b(T)}{V^{2} - V \ b(T)} - \frac{a(T)}{-(b(T))^{2} + 2 \ b(T)V + V^{2}}$$

(Note; The integral of the second term was taken from Hodgeman) (20)

$$= RT \ln \frac{V}{V - b(T)} + \frac{a(T)}{2\sqrt{2} b(T)} \ln \frac{V - 0.414 b(T)}{V + 2.414 b(T)} - RT \ln \frac{V}{V^{o}}$$

substituting in equation A5

using

$$b(T) = \frac{BRT}{P}$$
$$a(T) = \frac{AR^2T^2}{P}$$
$$V = \frac{RTZ}{P}$$

rives

$$S - S^{\circ} = \frac{\frac{da}{dT}}{2\sqrt{2} b(T)} \ln \frac{Z + 2.414 B}{Z - 0.414 B} - R \ln \frac{Z}{Z - B} + R \ln \frac{V}{V^{\circ}}$$

substituting in equation A6

$$H - H^{\circ} = RT \ln \frac{V}{V - b(T)} + \frac{a(T)}{2\sqrt{2} b(T)} \ln \frac{V - 0.414 \ b(T)}{V + 2.414 \ b(T)} - RT \ln \frac{V}{V^{\circ}}$$

$$+RT \ln \frac{V}{V^{o}} - RT \ln \frac{V}{V - b(T)} - \frac{T \frac{da}{dT}}{2\sqrt{2} b(T)} \ln \frac{V - 0.414 b(T)}{V + 2.414 b(T)} + RT(Z - 1)$$

Therefore

$$H - H^{o} = RT(Z - 1) + \frac{T\frac{da}{dT} - a(T)}{2\sqrt{2}b(T)} \ln \frac{Z + 2.414 B}{Z - 0.414 B}$$

Substituting in equation A7

.

$$ln\frac{f}{P} = ln\frac{V}{V-b(T)} + \frac{a(T)}{2\sqrt{2}b(T)} ln\frac{V-0.414b(T)}{V+2.414b(T)} - ln\frac{V}{V^{o}} + ln\frac{V}{V^{o}} + (Z-1)$$
$$= Z - 1 - ln(Z-B) - \frac{A}{2\sqrt{2}B} ln\frac{Z+2.414B}{Z-0.414B}$$

.

APPENDIX B

EGULAR SOLUTION THEORY CALCULATION USING THE VAN LAAR EQUATIONS

Nomenclature

 μ = free energy γ = acitivity coefficient η = fugacity coefficient

 $\delta =$ solubility coefficient

 $\frac{Subscript}{i = component}$

 $\frac{\text{Superscript}}{\text{o} = \text{datum}}$ - = partial molar

R113 ENTHALPY CONCENTRATION DIAGRAM theory and R12

calculations using liquid activity coefficients from the Van Laar equations and some thermodynamic properties from the PR equation.

B.1 Vapour Liquid equilibrium

Null (26) states that three conditions are required for vapour liquid equilibrium.

$$P_{iL} = P_{iV} \tag{B.1}$$

$$T_{iL} = T_{iV}$$

$$\mu_{iL} = \mu_{iV} \tag{B.2}$$

 $\mu_i = \mu_i^o + \Delta \mu_i$ (B.4)

where $\Delta \mu_i$ is the partial molar free energy of mixing

 μ^o_i is the pure substance free energy for that state at standard conditions.

Standard conditions for all subsequent calculations are system pressure and the temperature under consideration. Using $\Delta \mu_i = RTln(\gamma_i x_i)$ (B.5)

Null shows that

$$\frac{\gamma_{iL} \boldsymbol{x}_{iL}}{\gamma_{iV} \boldsymbol{x}_{iV}} = e \boldsymbol{z} \boldsymbol{p} \left(\frac{\boldsymbol{\mu}_{iV}^{o} - \boldsymbol{\mu}_{iL}^{o}}{RT} \right)$$
(B.6)

where

$$\gamma_i = \frac{f_i}{x_i f_i^o} \tag{B.7}$$

If

Null's equation for fugacity of a component in a mixture is

$$f_i = f_i^o x_i exp\left[\frac{1}{RT} \int_0^p (\bar{v}_i - v_i^o) dp\right]$$
(B.8)

The affect of the mixture is related through concentration and the fugacity at the reference state. Null evaluates the chemical potential difference term using a 3 step path between liquid and vapour based on

$$d\mu_i = -s_i dT + v_i dP \tag{B.9}$$

Step 1: Increase pressure from P to P on an psat incompressible liquid at constant temperature.

Step 2: Change of state at constant temperature and pressure.

Step 3: Compression of vapour at constant temperature. Summing these steps and substituting in equation B6 gives

$$\frac{y_i}{z_i} = \frac{\gamma_{iL} P_{isat}}{\gamma_{iV} P} exp\left[\frac{v_{iL} (P - P_{isat})}{RT}\right] exp\left[\frac{1}{RT} \int_{P_{isat}}^{P} \left(\frac{RT}{P} - v_{iL}\right)\right] \quad (B.10)$$

B.1.1 Activity Coefficients

B1.1.1 Vapour

Null suggests that assuming an ideal gas phase for a mixture is often an acceptable assumption. To test this the PR equation with a mixture interaction coefficient of 0.0 was used to generate fugacity values for the components in both the pure state and in the mixture. It was considered that if the variation in the evaluation of equation B10 attributable to the vapour activity coefficients was less that 1% then the coefficient could be considered to be unity. Table B1 shows that with the exception of the two lower mol fractions of R12 (0.4 and 0.5) which had temperatures close to the critical point of R12 (112 deg C), all variations were less than 1%. Based on this, and the fact that for the diagram at 423.3 kPa all R12 vapour activity coefficients were between 1.0 and 1.01, Null's assumption was considered acceptable.

Temp deg C	50	60	70	80	90	100	110
Mol fraction of R12				3			
0.4							1.017
0.5						1.012	1.011
0.6					1.008	1.008	
0.7			1.006	1.005	1.005		
0.8		1.003	1.003	1.002			0
0.9	1.0008	1.0007					
1.0	1.0						

lable Bl	Typical activity coefficients for R12 vapour	in
	and R12 R113 mixture at 1009 kPa.	

B.1.1.2 Liquid

Null expresses reservations about the ability of an equation of state and its mixing rules to accurately predict liquid molar volumes. He suggests that some of the "empirical or, at best semitheoretical approaches " available might be adequate. Of the methods discussed by him it was decided that the two constant binary equations of Van Laar were a good comprimise between accuracy and complexity. Edmister (15) gives the Van Laar equations in the form

$$ln\gamma_{1} = \frac{A}{\left[\frac{Az_{1}}{Bz_{2}} + 1\right]^{2}}$$
(B.11)
$$ln\gamma_{2} = \frac{B}{\left[\frac{Bz_{1}}{Az_{2}} + 1\right]^{2}}$$
(B.14)

where

$$A = \frac{v_1}{RT} (\delta_1 - \delta_2)^2$$
(B.13)
$$B = \frac{v_2}{RT} (\delta_1 - \delta_2)^2$$
(B.14)

δ	internal	energy	change	from	saturated	liquid	lo	ideal	ga .
v - V				molar	volume				

$$=\sqrt{\left[\frac{(h^o-RT)-(h_{sat}-RTZ_{sat})}{v_s}\right]}$$
 (B.15)

Note that this evaluation of the solubility coefficient δ assumes a constant temperature. Edmister discusses this and concludes "A constant value of the solubility

coefficient § at all temperatures is satisfactory and much simpler". In this case the reference state, ideal gas, was assumed to be vapour at zero pressure and the temperature under consideration. Reference state enthalpy was taken at the lowest pressure shown (for the required temperature) in published data. For example Dupont R12 (14) shows data down to 965 Pa. By calculating the molar volumes for A and B from the data, finding the solubility coefficient as above and knowing X1 and X2 the liquid activity coefficients were evaluated. Figure B1 shows that as the temperature approaches the critical value for R12 (112 deg C) the activity coefficient increases rapidly.

B.1.2 Evaluation of equation B10

Using published data and calculating activity coefficients as above all of equation B10 may be evaluated except the expression

$$exp\left[\frac{1}{RT}\int_{P_{iast}}^{P}\left(\frac{RT}{P-V_{iV}}\right)dP\right]$$
(B.16)

This may be done by rewriting this as two integrations

$$exp\left[\frac{1}{RT}\int_{0}^{P}\left(\frac{RT}{P-V_{iV}}\right)dP - \frac{1}{RT}\int_{0}^{P_{iast}}\left(\frac{RT}{P-V_{iV}}\right)dP\right]$$
$$= \frac{exp\left[\frac{1}{RT}\int_{0}^{P}\left(\frac{RT}{P-V_{iV}}\right)dP\right]}{exp\left[\frac{1}{RT}\int_{0}^{P_{iast}}\left(\frac{RT}{P-V_{iV}}\right)dP\right]}$$



ACTIVITY COEFFICIENT VS MOL CONC FOR SATURATED CONDITIONS

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ACTIVITY COEFFICIENT

Fig B1

And expressing the fugacity coefficient of a pure substance as

$$\eta_i = \frac{f_i}{P} = exp\left[\frac{1}{RT}\int_0^P v_i^o - \frac{RT}{P}dp\right]$$

Equation B16 becomes

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$$\frac{\eta_{iPsat}}{\eta_{iP}} = \frac{fugacily \ coeff \ at \ saturation \ pressure}{fugacily \ coeff \ at \ system \ pressure} \tag{B.17}$$

These fugacities were obtained from the PR equation. Table B2 shows some typical values.

Temperature		R1	.2	R113		
С	ĸ	system	satd	system	satd	
		*:				
50	323.16	0.8497	0.8187	0.596	0.9605	
60	333.16	0.8621	0.7928	0.64	0.9501	
70	343.16	0.8732	0.765	0.675	0.9379	
80	353.16	0.8832	0.738	0.7118	0.9246	
90	363.16	0.8923	0.7086	0.7367	0.9097	
100	373.16	0.9009	0.6798	0.7576	0.8939	
110	383.16	0.9080	0.6496	0.7760	0.8752	

TABLE B2:

Fugacity coefficients for pure vapour

In summary a combination of pure published data, the Van Laar equations and the PR equation was used to evaluate equation B10. B.1.3 Total system pressure calculation

Rewriting equation B.10 using equation B.17 gives

$$y_{i} = \frac{x_{i}\gamma_{iL}P_{isat}\eta_{isat}}{\gamma_{iV}P\eta}exp\left[\frac{v_{iL}(P-P_{isat})}{RT}\right]$$

A mass balance for a binary system yields

$$1 = y_1 + y_2$$

$$P=P(y_1+y_2)$$

Substituting for Y1 and Y2

$$P = \frac{x_1 \gamma_{1L} P_{1sat} \eta_{1sat}}{\gamma_{1V} P \eta} exp \left[\frac{v_{1L} (P - P_{1sat})}{RT} \right]$$
$$+ \frac{x_2 \gamma_{2L} P_{2sat} \eta_{2sat}}{\gamma_{2V} P \eta} exp \left[\frac{v_{2L} (P - P_{2sat})}{RT} \right] \quad (B. 18)$$

Knowing P and assuming vapour activity coefficients to be 1.0 the system equilibrium may be found by



B.1.4 Sample calculation for R12 - R113 mixture $F = 1.009 \times 10^{-6}$ Pa (Satn pressure of R12 at 42 deg C) T = 50 deg C

Evaluating the terms in equation A10

	R12	R113
$Part1 = exp\left[\frac{v_{iL}(P - P_{isat})}{RT}\right]$	0.9922	1.0427
$Part2 = \frac{P}{P_{isat}}$	1.2091	0.109
$Part3 = \frac{\eta_{isat}}{\eta_P}$	0.9635	1.612

Substituting in B10 for R12

$$y = x \frac{\gamma_L}{\gamma_V} 1.2091 \quad 0.9922 \quad 0.9635$$

Assuming vapour activity coefficient to be one then

$$y = x \gamma_L \quad 1.1559$$

Try X1 = 0.8

Using the Van Laar equations

$$\delta_1 = 11939$$

 $\delta_2 = 14130$
 $A = 0.1784$
 $B = 0.223$
 $\gamma_L = 1.01$

Check system pressure using equation B1B

$$P = 0.8 1.01 1.22 \times 10^{6} 0.9635 0.9922 + 1.00^{5} 0.2 1.138 1.1 \times 10^{5} 1.612 1.0427 - 1.00^{6}$$

$$P = 0.984 \times 10^{6} Pa$$
Residual = -2.4%

Using a second guess and a graphical approach the final result was

X1 = 0.83 and Y1 = 0.97 with a residual of -0.3%

B.2 Subcooled liquid mixture enthalpies

An ideal liquid mixture was assumed, the effects of pressure were ignored and mixture enthalpies were taken as molar proportions of the pure saturated values for that temperature.

$h = x_1 h_{1sal} + x_2 h_{2sal}$

B.3 Superheated vapour enthalpies

Instead of assuming an ideal mixture for the vapour phase it was decided to try a 4 part process which included enthalpy departure values from the PR equations. The process parts were: Part1: Evaluate latent heat of mixture at datum temperature based on molar proportions of the pure latent heats.

Part2: Calculate enthalpy departure at datum temperature, from pressure corresponding to reference condition to zero pressure.

Part3: At zero pressure integrate the specific heat capacity function from T to T. datum

Part4: Calculate enthalpy departure, at T from zero

pressure to system pressure.

Temp	emp State kJ/kg						Percent	
deg C		Part1	Part2	Part3	Part4	Calcd	Publd	Error
42 70 100	satd s/heat s/heat	169.6 169.6 169.6	0.98 0.98 0.98	47.4 64.9 84.3	-11.7 -9.8 -8.3	206.3 225.7 246.6	204.3 225.4 247.5	0.9 0.1 -0.4

Table B3 shows some typical values for pure R12

Table B3 Enthalpy calculation data

B.4 Enthalpy concentration diagram

Note that the rapid change in the near critical activity coefficients for R12 produce a steep slope in the boiling point equilibrium line in Figure B3.

280 230 180 80.0 60.0 40.0 130 20 deg C 80 30 .2 . 8 .6 <u>,</u>4 0

ENTHALPY CONC DIAG FOR R12 - R113 AT P=432.3 KPA

Fig B2

MASS CONC OF R12

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ENTHALPY KJ/KG



ENTHALPY CONC DIAGRAM FOR R12 R113 AT P=1009 KPA

MASS CONCENTRATION OF R12

Fig B3

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ENTHALPY KJ/KG

This method of producing enthalpy concentration data was abandoned because it relied too much on published thermodynamic data and was difficult to program.

APPENDIX C

COMPUTER MODEL INPUT FILES

C.1 Main input file

One input file was used for both program control and for the refrigeration system model input data. The following file structure layout illustrates that the variables ME to JC inclusive relate to program options and their associated data while variables RN(1) to TD(14) detail data for working fluids and the refrigeration system model.

VARIABLE	COMMENT				
ME	O Interactive mode 2 Batch mode				

MF	O Supresses all program comments to screen other than error messages				
	1 During run gives indication of progress				
*****	***************************************				
пн	lable of pure properties. O No table				
	1 Saturated values only				
	2 Saturated and superheat				
мни	company with published data 4 pressure				
	compare with published used i pressure				
	Z all properties				
	A optimize kappa op liguid - value				
	S optimize kappa on indulu z value				
	S Optimize kappa on pressure				
MHI	0 51				
	1 Imperial				
****	******				
МК	Mixture data O No table				
	1 calculate for mixture data given				
	2 compare with data from mixture property file				
MKK	Compare single calc with 1 Pressure (PMK)				
	2 Mass concentration (XMK)				
0.000.000.	3 Mol concentration (XMMK)				
TMK	Temperature in deg C for single calc				
РМК	Pressure in Pa for single calc				
ХМК	Mass cone of component 1 for single cale				
хммк	Mol conc of component 1 for single calc				
*********	****				
mJ	Enthalpy concentration diagram O No diagram				
	1 Equilbrium only				
MA	2 Complete diagram				
nu	Plot files U Table only				
THY	1 Table plus plot file				
1111	temperature in deg C. Saturation temperature for most				
DHT	volatile component				
PHI	Pressure in Pa. Diagram pressure				
	NOIL to I fill and PHI the one not used must be negative				
	Ini (deg C) here diag pressure is sath pressure				
	corresponding to TMI for component 1				
ADD	Mala (Palls pressure of diagram				
VALUE	How have descent (minimum U.01)				
VALUE	now many degrees of isotherm above and below saturation				

MG	Pure data detail O No data
114	2 Data for olot of fugacity vs pressure
	3 Data for Peng-Robinson cubic plot
NEA	Level for Perg-Rebinson cubic plot
MG1	Input for pang-domant 4
	2 Use reirigerand 2
	3 Use mixture
TEMPMG	Temperature for Peng-Robinson cubic pict
PRESSMG	Pressure for Peng-Robinson cubic plot
CONCMG	Mass concentration for Peng-Robinson cubic plot
**********	***************************************
ML	Refrigeration system calculation
1	D No system calculation
	1 System calc
1 11	Saturation pressure calculation method
55	a Dense Poblisson equation
	2 Clausius Clapeyrun eduation
JL	Pressure calculation at enthalpy used
1	O Simple molar proportion at datum temperature
	1 Does equilibrium calc at datum temperature
JC	Peng Robinson Kappa value calo method
	O Standard K calc based single K for whole program
	1 Uses oubic in TR for K values
*********	********
RN(1)	Refrigerant number
RN(2)	Refrigerant number
	Order does not matter. Program will put most volatile
	in position 1
	Refrigerants available are
	R11, R12, R13, R13B1, R22, R23, R113, RPERC, R718, R152A
	PEND
	Transitive datum for enthalow and entropy deg C
	Temperature datum for endispy and the of the second s
BT	Jemperature datum for availability day o
BP	Pressure datum for availability re

	and the second sec
TONE	Fluid temperature entering evaporator deg C
TONE TOFFE	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C
TONE TOFFE SUPHE	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate superheat at evaporator out Celcius degrees
TONE TOFFE Suphe Evapcap	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate superheat at evaporator out Celcius degrees Evaporator capacity kW
TONE TOFFE SUPHE Evapcap Tona	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate superheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C
TONE TOFFE SUPHE Evapcap Tona Subca	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate superheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees
TONE TOFFE SUPHE Evapcap Tona Subca Tonc	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate superheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C
TONE TOFFE SUPHE EVAPCAP TONA SUBCA TONC SUBCC	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate suberheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C Distillate subcooling at evaporator out Celcius degrees
TONE TOFFE SUPHE EVAPCAP TONA SUBCA TONC SUBCC DCONCMASE	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate suberheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C Distillate subcooling at evaporator out Celcius degrees Mass concentration of distillate
TONE TOFFE SUPHE EVAPCAP TONA SUBCA TONC SUBCC DCONCMASE FOD	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate suberheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C Distillate subcooling at evaporator out Celcius degrees Mass concentration of distillate Mass ratio of feed to distillate flow
TONE TOFFE SUPHE EVAPCAP TONA SUBCA TONC SUBCC DCONCMASS FOD TEFED	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate superheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C Distillate subcooling at evaporator out Celcius degrees Mass concentration of distillate Mass ratio of feed to distillate flow Temperature after feed boost heater
TONE TOFFE SUPHE EVAPCAP TONA SUBCA TONC SUBCC DCONCMASS FOD TFEED MTEFED	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate suberheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C Distillate subcooling at evaporator out Celcius degrees Mass concentration of distillate Mass ratio of feed to distillate flow Temperature after feed boost heater Feed boost heater selection
TONE TOFFE SUPHE EVAPCAP TONA SUBCA TONC SUBCC DCONCMASS FOD TFEED MTFEED	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate superheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C Distillate subcooling at evaporator out Celcius degrees Mass concentration of distillate Mass ratio of feed to distillate flow Temperature after feed boost heater Feed boost heater selection O No
TONE TOFFE SUPHE EVAPCAP TONA SUBCA TONC SUBCC DCONCMASS FOD TFEED MTFEED	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate suberheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C Distillate subcooling at evaporator out Celcius degrees Mass concentration of distillate Mass ratio of feed to distillate flow Temperature after feed boost heater Feed boost heater selection O No 4 Yee
TONE TOFFE SUPHE EVAPCAP TONA SUBCA TONC SUBCC DCONCMASS FOD TFEED MTFEED	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate suberheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C Distillate subcooling at evaporator out Celcius degrees Mass concentration of distillate Mass ratio of feed to distillate flow Temperature after feed boost heater Feed boost heater selection O No 1 Yes Proceed beat exchanger
TONE TOFFE SUPHE EVAPCAP TONA SUBCA TONC SUBCC DCONCMASE FOD TFEED MTFEED MPRECOOL	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate superheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C Distillate subcooling at evaporator out Celcius degrees Mass concentration of distillate Mass ratio of feed to distillate flow Temperature after feed boost heater Feed boost heater selection O No 1 Yes Precooler heat exchanger
TONE TOFFE SUPHE EVAPCAP TONA SUBCA TONC SUBCC DCONCMASE FOD TFEED MTFEED MPRECOOL	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate superheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C Distillate subcooling at evaporator out Celcius degrees Mass concentration of distillate Mass ratio of feed to distillate flow Temperature after feed boost heater Feed boost heater selection 0 No 1 Yes Precooler heat exchanger 0 No
TONE TOFFE SUPHE EVAPCAP TONA SUBCA TONC SUBCC DCONCMASE FOD TFEED MTFEED MPRECOOL	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate superheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C Distillate subcooling at evaporator out Celcius degrees Hass concentration of distillate Mass ratio of feed to distillate flow Temperature after feed boost heater Feed boost heater selection O No 1 Yes Precooler heat exchanger O No 1 Yes
TONE TOFFE SUPHE EVAPCAP TONA SUBCA TONC SUBCC DCONCMASE FOD TFEED MTFEED MPRECOOL MRECTHX	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate suberheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C Distillate subcooling at evaporator out Celcius degrees Mass concentration of distillate Mass ratio of feed to distillate flow Temperature after feed boost heater Feed boost heater selection O No 1 Yes Precooler heat exchanger O No 1 Yes Rectification heat exchanger
TONE TOFFE SUPHE EVAPCAP TONA SUBCA TONC SUBCC DCONCMASE FOD TFEED MTFEED MPRECOOL MRECTHX	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate suberheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C Distillate subcooling at evaporator out Celcius degrees Mass concentration of distillate Mass ratio of feed to distillate flow Temperature after feed boost heater Feed boost heater selection O No 1 Yes Precooler heat exchanger O No 1 Yes Rectification heat exchanger O No
TONE TOFFE SUPHE EVAPCAP TONA SUBCA TONC SUBCC DCONCMASS FOD TFEED MTFEED MPRECOOL MRECTHX	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate suberheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C Distillate subcooling at evaporator out Celcius degrees Mass concentration of distillate Mass ratio of feed to distillate flow Temperature after feed boost heater Feed boost heater selection O No 1 Yes Precooler heat exchanger O No 1 Yes Rectification heat exchanger O No 1 Yes
TONE TOFFE SUPHE EVAPCAP TONA SUBCA TONC SUBCC DCONCMASE FOD TFEED MTFEED MPRECOOL MRECTHX MPREHEAT	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate suberheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C Distillate subcooling at evaporator out Celcius degrees Mass concentration of distillate Mass ratio of feed to distillate flow Temperature after feed boost heater Feed boost heater selection O No 1 Yes Precooler heat exchanger O No 1 Yes Rectification heat exchanger O No 1 Yes Preheater heat exchanger
TONE TOFFE SUPHE EVAPCAP TONA SUBCA TONC SUBCC DCONCMASE FOD TFEED MTFEED MPRECOOL MRECTHX MPREHEAT	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate suberheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C Distillate subcooling at evaporator out Celcius degrees Mass concentration of distillate Mass ratio of feed to distillate flow Temperature after feed boost heater Feed boost heater selection O No 1 Yes Precooler heat exchanger O No 1 Yes Preheater heat exchanger O No 1 Yes
TONE TOFFE SUPHE EVAPCAP TONA SUBCA TONC SUBCC DCONCMASE FOD TFEED MTFEED MPRECOOL MRECTHX MPREHEAT	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate suberheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C Distillate subcooling at evaporator out Celcius degrees Mass concentration of distillate Mass ratio of feed to distillate flow Temperature after feed boost heater Feed boost heater selection O No 1 Yes Precooler heat exchanger O No 1 Yes Preheater heat exchanger O No 1 Yes Preheater heat exchanger O No 1 Yes
TONE TOFFE SUPHE EVAPCAP TONA SUBCA TONC SUBCC DCONCMASE FOD TFEED MTFEED MPRECOOL MRECTHX MPREHEAT PHPROP	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate suberheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C Distillate subcooling at evaporator out Celcius degrees Hass concentration of distillate Mass ratio of feed to distillate flow Temperature after feed boost heater Feed boost heater selection O No 1 Yes Precooler heat exchanger O No 1 Yes Rectification heat exchanger O No 1 Yes Preheater heat exchanger O No 1 Yes Proportion of feed flow through preheater ie O to 1.0
TONE TOFFE SUPHE EVAPCAP TONA SUBCA TONC SUBCC DCONCMASE FOD TFEED MPRECOOL MRECTHX MPREHEAT PHPROP MBOTHX	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate suberheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C Distillate subcooling at evaporator out Celcius degrees Mass concentration of distillate Mass ratio of feed to distillate flow Temperature after feed boost heater Feed boost heater selection O No 1 Yes Preccoler heat exchanger O No 1 Yes Preheater heat exchanger O No 1 Yes Preheater heat exchanger O No 1 Yes Preheater heat exchanger O No 1 Yes Preheater heat exchanger O No 1 Yes Proportion of feed flow through preheater is O to 1.0 Bottoms heat exchanger
TONE TOFFE SUPHE EVAPCAP TONA SUBCA TONC SUBCC DCONCMASE FOD TFEED MPRECOOL MRECTHX MPREHEAT PHPROP MBOTHX	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate suberheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C Distillate subcooling at evaporator out Celcius degrees Hass concentration of distillate Mass ratio of feed to distillate flow Temperature after feed boost heater Feed boost heater selection O No 1 Yes Precooler heat exchanger O No 1 Yes Preheater heat exchanger O No 1 Yes Preheater heat exchanger O No 1 Yes Preheater heat exchanger O No 1 Yes Proportion of feed flow through preheater ie O to 1.0 Bottoms heat exchanger O No
TONE TOFFE SUPHE EVAPCAP TONA SUBCA TONC SUBCC DCONCMASE FOD TFEED MTFEED MPRECOOL MRECTHX MPREHEAT PHPROP MBOTHX	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate suberheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C Distillate subcooling at evaporator out Celcius degrees Hass concentration of distillate Mass concentration of distillate flow Temperature after feed boost heater Feed boost heater selection O No 1 Yes Precooler heat exchanger O No 1 Yes Rectification heat exchanger O No 1 Yes Preheater heat exchanger O No 1 Yes Proportion of feed flow through preheater ie O to 1.0 Bottoms heat exchanger O No 1 Yes
TONE TOFFE SUPHE EVAPCAP TONA SUBCA TONC SUBCC DCONCMASE FOD TFEED MPRECOOL MRECTHX MPREHEAT PHPROP MBOTHX RR	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate suberheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C Distillate subcooling at evaporator out Celcius degrees Mass concentration of distillate Mass ratio of feed to distillate flow Temperature after feed boost heater Feed boost heater selection O No 1 Yes Precooler heat exchanger O No 1 Yes Preheater heat exchanger O No 1 Yes Preheater heat exchanger O No 1 Yes Proportion of feed flow through preheater ie O to 1.0 Bottoms heat exchanger O No 1 Yes Proportion of feed flow through preheater ie O to 1.0 Bottoms heat exchanger O No 1 Yes Proportion of feed flow through preheater ie O to 1.0 Bottoms heat exchanger O No 1 Yes Proportion of feed flow through preheater ie O to 1.0 Bottoms heat exchanger O No 1 Yes Proportion of feed flow through preheater ie O to 1.0 Bottoms heat exchanger O No 1 Yes Preflux ratio (Ratio of actual reflux flow to minimum flow)
TONE TOFFE SUPHE EVAPCAP TONA SUBCA TONC SUBCC DCONCMASE FOD TFEED MPRECOOL MRECTHX MPREHEAT PHPROP MBOTHX RR MPCONTPO	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate suberheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C Distillate subcooling at evaporator out Celcius degrees Hass concentration of distillate Mass ratio of feed to distillate flow Temperature after feed boost heater Feed boost heater selection O No 1 Yes Precooler heat exchanger O No 1 Yes Preheater heat exchanger O No 1 Yes Preheater heat exchanger O No 1 Yes Preheater heat exchanger O No 1 Yes Proportion of feed flow through preheater ie O to 1.0 Bottoms heat exchanger O No 1 Yes Prototion of feed flow through preheater ie O to 1.0 Bottoms heat exchanger O No 1 Yes Prototion of feed flow through preheater ie O to 1.0 Bottoms heat exchanger O No 1 Yes Prefux ratio (Ratio of actual reflux flow to minimum flow) Print out control
TONE TOFFE SUPHE EVAPCAP TONA SUBCA TONC SUBCC DCONCMASE FOD TFEED MPRECOOL MRECTHX MPREHEAT PHPROP MBOTHX RR MPCONTRO	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate suberheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C Distillate subcooling at evaporator out Celcius degrees Mass concentration of distillate Mass ratio of feed to distillate flow Temperature after feed boost heater Feed boost heater selection O No 1 Yes Precooler heat exchanger O No 1 Yes Preheater heat exchanger O No 1 Yes Preheater heat exchanger O No 1 Yes Proportion of feed flow through preheater ie O to 1.0 Bottoms heat exchanger O No 1 Yes Proportion of feed flow through preheater ie O to 1.0 Bottoms heat exchanger O No 1 Yes Protout control 1 System layout only
TONE TOFFE SUPHE EVAPCAP TONA SUBCA TONC SUBCC DCONCMASE FOD TFEED MPRECOOL MRECTHX MPREHEAT PHPROP MBOTHX RR MPCONTRO	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate suberheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C Distillate subcooling at evaporator out Celcius degrees Hass concentration of distillate Mass ratio of feed to distillate flow Temperature after feed boost heater Feed boost heater selection O No 1 Yes Precooler heat exchanger O No 1 Yes Preheater heat exchanger O No 1 Yes Preheater heat exchanger O No 1 Yes Proportion of feed flow through preheater is O to 1.0 Bottoms heat exchanger O No 1 Yes Reflux ratio (Ratio of actual reflux flow to minimum flow) Print out control 1 System layout only 2 System layout only 2 System layout only
TONE TOFFE SUPHE EVAPCAP TONA SUBCA TONC SUBCC DCONCMASE FOD TFEED MPRECOOL MRECTHX MPREHEAT PHPROP MBOTHX RR MPCONTRO	Fluid temperature entering evaporator deg C Fluid temperature leaving evaporator deg C Distillate suberheat at evaporator out Celcius degrees Evaporator capacity kW Fluid temperature onto absorber deg C Feed subcooling at absorber out Celcius degrees Fluid temperature onto condenser deg C Distillate subcooling at evaporator out Celcius degrees Hass concentration of distillate Mass ratio of feed to distillate flow Temperature after feed boost heater Feed boost heater selection O No 1 Yes Precooler heat exchanger O No 1 Yes Preheater heat exchanger O No 1 Yes Preheater heat exchanger O No 1 Yes Proportion of feed flow through preheater is O to 1.0 Bottoms heat exchanger O No 1 Yes Print out control System layout only 2 System plus column analysis 3 System plus column analysis plus Second Law analysis

********	***************************************
	Refrigeration system approach values
A(1)	Evap out to fluid on evap
2	Condenser out to fluid onto absorber
3	Absorber out to fluid onto absorber
4	Precooler liquid in to precooler vapour out
5	Preheater feed in to preheater bottoms out (mixture end)
6	Preheater feed out to preheater bottoms in (column end)
7	Rectifier HX out to feed plate
8	Stripping HX out to feed plate
9	Rectifier HX in to plate temperature
10	Stripping HX in to plate temperature
*********	***************************************
	Refrigeration system pressure drops
PD(1)	Absorber out to pump in
2	Pump out to feed split
3	Feed split to feed PH in
4	Feed solit to Rectification HX in
5	Rectification HX in to out
6	Rectification HX out to feed to n
7	Feed feed PH in to out
8	Feed PH out to feed join
9	Feed join to column in
10	Bottoms out to bottom HX in
11	Bottom HX in to out
12	Bottom HX out to preheater in
13	Bottom feed preheat out to expansion in
14	Condernser out to precover liquid in
15	Precoler liquid in to out
16	Precoder liquid out to expansion in
17	Evaporator in to out
18	Evaporator out to precoder vapour in
19	Precoler vacuur in to out
20	Precoler vapour out to mix in
21	Mix out to absorber in
22	Absorber in to out
23	Bottom feed preheat in to out
24	Condenser in to out
********	*****
	Refrigeration system temperature changes
TD(1)	Absorber out to pump in
2	Pump out to feed split
3	Feed split to feed HX in
4	Feed split to Rectification HX in
5	Rectification HX out to feed mix
6	Feed PH out to feed mix
7	Feed boost HX to column in
8	Bottoms out to bottom HX in
9	Bottom HX out to feed PH in
10	Feed PH out to bottom exp in
11	Condenser out to precoler liquid in
12	Precoler liquid out to distillate evnansion
13	Evaporator out to PC vapour in
14	Precooler vapour out to mixing in

NOTE: The following abbreviations have been used

Heat exchanger to HX Precooler to PC Preheater to PH Peng-Robinson to PR

C.2 Pure substance input files

C.2.1 File structure and units

In addition to tabulating the values of constants used for all the pure substance data derivation this section shows the layout of the program data files.

Item	Symbol	Units	comment
name			
formula			
Critical pressure	Pc	Pa	
Critical temp	Tc	Deg K	
Molecular Wt	MW		
Pr K constant	л К		Chap. 1 & appendix A
Dead zone		Deg	Chap. 1 section 1.2
Specific heat capacity const.			
	³ C4		Chap. 1 equ 1.17
4 lipes	С3		н 🐤
, Trues	C2		
	Ci		т. н.
Clausius equn const.		(H	
2 lines	AR		Clausius Clapeyron
L TINES	BR		equation constants
4 lines (See Note	1 below)	
7 lines (See Note	2 below	>	
n lines (See note	3 below	>	

These constants are for a cubic equation which fits the Peng-Robinson constant K as a function of temperature.

This feature, which was suggested by Dr J. R. Roach, was programmed as an alternative to the single K value approach discussed in chapter 1. It was hoped that fitting K as a function of temperature would improve the Peng-Robinson equation's accuracy. Time did not permit the calculation of these constants.

Note 2:

These constants are for the Dupont vapour pressure equation. If these constants were available this equation was used to check the published data.

Note 3:

Each of the next n lines contains published saturation values of temperature, pressure, liquid density, liquid enthalpy, latent heat, vapour enthalpy, liquid entropy, vapour entropy, liquid compressiblity and vapour compressiblity. These values appear as the published values of the respective properties shown in Appendix E. All compressiblity values shown were calculated from the published tables.

C.2.2 File content

The ASHRAE refrigerant number shown in the following table is not part of the data file it has just been included for convenience.

R12	R22	R23
'DICHLORODIFLUOROMETHANE' 'CCL2F2' 4.1155E+06 385.16 120.9 0.6352 0.6 7.547 0.04257 -0.3603E-04 0.1037E-07 9.36195982 1.0581307E+03 0 0 0 39.88381727 -3.43663223E+03 -12.47152228 4.73044244E-03 0.0 0.0 459.7	<pre>'CHLORODIFLUOROMETHANE' 'CHCLF2' 4.9773128E+06 369.16 86.48 0.7020 0.7 6.035167 2.128E-02 2.2E-05 -3.739705E-08 9.567632 1.060095E+03 0 0 0 29.35754453 -38.45193152E+02 -7.86103122 2.190939E-03 4.45746403E-01 6.861E+02 459.69</pre>	'TRIFLUOROMETHANE' 'CHF3' 4.836013E+06 298.77 70.00 0.7822 0.4 5.339472 0.020029 1.8E-05 -2.443922E-08 9.62552 881.3117 0 0 0 0 328.90853 -7952.76913 -144.51423 0.24211502 -2.1280665E-04 9.434955E-08 459.69
R11	R13B1	R718
<pre>'TRICHLOROFLUOROMETHANE' 'CCL3F' 4.409199E+06 471.16 137.38 0.6654 0.7 9.789 3.893E-02 -3.383E-05 9.903E-09 9.51930112 1.35459E+03 0 0 0 0 42.14702865 -4344.343807 -12.84596753 0.0040083725 0.0313605356 862.07 459.67</pre>	<pre>'BROMOTRIFLUOROMETHANE' 'CBRF3' 3.964487E+06 340.16 148.93 0.6319 0.4 5.22741862 0.04851168 -3.557E-05 7.14402592E-10 9.32427949 928.499988 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0</pre>	<pre>'WATER' 'H20' 22048000.0 647.3 18.015 0.8508 0.9 7.701 4.595E-04 2.521E-06 -8.59E-10 10.461 2020.097 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0</pre>

.

R13	*		R152a
<pre>' CHLOROTRIFLUOROMETHANE' ' CCLF3' 3.867983E+06 301.99 104.47 0.6321 0.4 -4.662728E+01 0.54353 -1.62E-03 1.6836E-06 9.35722 8.3659361E+02 0 0 0 0 25.967975 -2.70953822E+03 -7.17234391 2.545154E-03 2.80301091E-01 5.46E+02 459.67</pre>	<pre>, 4610010-2910000000000000000000000000000000</pre>	TETRACHLOROETHYLENE' C2CL4' .7640E+06 20.0 65.83 .7511 .8000 0.98 .05387 5.478E-05 .002E-08 .8098831 912.83397	'DIFLUOROETHANE' 'CH3CHF2' 4.49537050E+06 386.66 66.05 0.7608 0.5 -6.37682450E+01 6.67859200E-01 -2.11450000E-03 2.4000000E-06 9.60311 1.14085525E+03 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0
R113		R500	
<pre>'TRICHLOROTRIFLUOROETHANE 'CCL2F-CCLF2' 3.4397858E+06 487.27 187.39 0.751400 0.6 2.4401721 1.182996E-01 -1.243E-04 4.1417751E-08 9.48006088 1.43562224E+03 0 0 0 0 33.0655 -4.33098E+03 -9.2635 2.0539E-03 0.0 0.0 459.6</pre>		<pre>'73.8% CCL2F2 26.2% 'R12 & R152A' 4425707.9 378.66 99.31 0.6887 0.6 33.462211 -0.189143 0.000652294 -6.66E-07 9.435097 1058.3573 0 0 17.780935 -3422.69717 -3.63691 0.00050272207 0.4629401 695.57 459.67</pre>	снзснг2' No number

APPENDIX D

COMPARISON WITH SIMILAR DATA DERIVED FROM

SOAVE-REDLICH-KWONG (SRK) EQUATION

Comparison with Asselineau et al (4)

Asselineau used 21 points to check the SRK equation saturation data with published data. Five points were checked in each of the intervals

> $0.6 \le Tr \le 0.7$ $0.7 \le Tr \le 0.8$ $0.8 \le Tr \le 0.9$

and three in each of the ranges

$$0.9 \le Tr \le 0.95$$

The deviation parameter used was

SD2 =
$$\sqrt{\frac{\sum_{1}^{n} \left(\frac{calcd-publd}{publd}\right)^{2}}{(n-1)}}$$

Table D1 shows the SD2 values calculated here and those for Asselineau's method three.

		This wo	Asselineau		
Reft	PR	Tr		No.	SRK
No	SD2	lower	upper	pts	SD2
11	0.0077	0.495	0.99	43	0.0126
12	0.0066	0.605	0.995	28	0.0069
13	0.002	0.772	0.993	13	0.0078
22	0.0037	0.632	0.993	25	0.0103
113	0.040	0.479	0.992	46	0.0078
152a	0.0141	0.603	0.962	26	0.0224

Table D1: Saturation Pressure data

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Even though the distribution of points is not the same for each case, with the exception of R13, the range considered was approximately equal to or larger than that of Assenleau. Other than for R113 the PR equation values showed the lesser deviation from the published data.

APPENDIX E

CALCULATED VERSUS PUBLISHED

REFRIGERANT DATA

Table	Fluid	Data
E.1	R11	abridged
E.2	R12	complete
E.3	R13	abridged
E.4	R13B1	н
E.5	R22	л ,
E.6	R113	11
E.7	R152a	н
-E.8	R718	н
E.9	R500	e ^d u
E.10	Perchloroethylene	н
E.11	R13B1 - R152a	complete
E.12	R11 - R22	11
E.13	R12 - R152a	u.
E.14	R12 - R13	п
E.15	R12 - R22	n
E.16	R12 - R113	н

Equations (for definitions see Appendix A)

$$AAD = \frac{\sum_{1}^{n} \left| \left(\frac{calcd - publd}{publd} \right) \right| 100}{n}$$

SD1 = $\sqrt{\frac{\sum_{1}^{n} (calcd - publd)^{2}}{(n-1)}}$
SD2 = $\sqrt{\frac{\sum_{1}^{n} \left(\frac{calcd - publd}{publd} \right)^{2}}{(n-1)}}$
PRESSURE COMPARISON

TABLE E1

TEMP Deg f	PRESSUR	E PSIA PUBLD	ABS Error	PERCENT Error	N	TR	PR
-40.00	0.752E+00	0.743E+00	0.915E-02	1.23	1	0.4949	0.0012
-30.00	0.105E+01	0.104E+01	0.635E-02	0.61	2	0.5067	0.0016
-20.00	0.143E+01	0.143E+01	0.304E-02	0.21	3	0.5184	0.0022
-10.00	0.1928+01	0.192E+01	-0.209E-02	-0.11	4	0.5302	0.0030
40.00	0.255E+U1	U.256E+U1	-0.112E-01	-0.44	5	0.5420	0.0040
20.00	0.3342+01	U.336E+U1	-U.228E-U1	-0.68 0	6	0.5538	0.0052
30.00	0.4312401	0.4305701	-0.54775-04	-1.10		U.5656	0.0067
40.00	0.5302+01	0.3302+01	-0.3032-01	-1.01	8	0.5000	0.0088
50.00	0.868F+01	0.7022+01	-0.1045+00	-1+11	10	0.3072	0.0109
60.00	0.107E+02	0.109E+02	-0.136E+00	-1.25	11	0.6128	0.0158
70.00	0.132E+02	0.133E+02	-0.171E+00	-1.28	12	0.6246	0.0206
80.00	0.160E+02	0.162E+02	-0.207E+00	-1.29	13	0.6364	0.0251
90.00	0.193E+02	0.196E+02	-0.251E+00	-1.28	14	0.6481	0.0302
100.00	0.232E+02	0.235E+02	-D.294E+00	-1.25	15	0.6599	0.0362
110.00	0.276E+02	0.279E+02	-0.338E+00	-1.21	16	0.6717	0.0431
120.00	0.326E+02	0.329E+02	-0.384E+00	-1.16	17	0.6835	0.0509
130.00	0.382E+02	0.387E+D2	-0.427E+00	-1.10	18	0.6953	0.0598
140.00	0.447E+02	0.451E+02	-0.468E+00	-1.04	19	0.7071	0.0698
150.00	0.519E+02	0.524E+02	-0.505E+00	-0.96	20	0.7189	0.0811
160.00	0.599E+02	0.605E+02	-0.534E+00	-0.88	21	0.7307	0.0937
180.00	U.687E+U2	0.694E+02	-0.556E+00	-0.80	22	0.7425	0.1077 -
	U./00E+UZ	U.794E+U2	-0.5682+00	-0.71	23	0.7543	0.1233
200 00	0.070E+UZ	0.4075.07	-0.567E+00	-0.63	24	0.7661	0.1405
210.00	0.1155+03	0.1032703	-0.5522700	-0.45	25	0.7494	0.1574
220.00	0.1305+03	0.1102+03	-0.5100+00	-0.45	20	0.001/	0.1003
230.00	0.146E+03	0.146F+03	-0.4155+00	-0.37	28	0.8132	0.2030
240.00	0.163Ef03	0.163E+03	-0.3296+00	-0.20	20	0.8250	0.2549
250.00	0.182E+03	0.182E+03	-0.231E+00	-0.13	30	0.0250	0.2843
260.00	0.202E+03	0.202E+03	-0.104E+00	-0.05	31	0.8486	0.3161
270.00	0.224E+03	D.224E+03	0.228E-01	0.01	32	0.8604	0.3504
280.00	0.248E+03	0.248E+03	0.181E+00	0.07	33	0.8722	0.3875
290.00	0.273E+03	0.273E+03	0.335E+00	0.12	34	0.8840	0.4274
300.00	0.301E+03	0.300E+03	0.500E+00	0.17	35	0.8958	0.4702
310.00	0.330E+03	0.329E+03	0.665E+00	0.20	36	0.9076	0.5162
320.00	0.362E+D3	0.361E+03	0.821E+00	0.23	37	0.9193	0.5653
330.00	0.395E+03	0.374E+03	0.951E+00	0.24	38	0.9311	0.6179
340.00	0.431E+03	0.430E+03	0.105E+01	0.24	39	0.9429	0.6740
350.00	0.469E+03	0.468E+03	0.110E+01	0.24	40	0.9547	0.7338
560.00	0.510E+03	0.509E+03	0.109E+01	0.21	41	0.9665	0.7974
\$70.00	0.553E+03	0.552E+03	0.980E+00	0.18	42	0.9783	0.8650
580.00	0.599E+03	0.598E+03	0.725E+00	0.12	43	0.9901	0.9369

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AAD= 0.6190972 3D1= 0.5112267 3D2= 7.7368235E-03

********** VAPOUR ENTHALPY ****

TABLE E1 CONTINUED

-40.00 0 -30.00 0 -20.00 0 -10.00 0 0.00 0).859E+02).871E+02).883E+02).896E+02).908E+02	0.872E+02 0.884E+02 0.896E+02 0.908E+02	-0.134E+01 -0.128E+01	-1.53	1	Π. 4949	0.0041
-30.00 C -20.00 C -10.00 C 0.00 C	0.871E+02 0.883E+02 0.876E+02 0.908E+02	0.884E+02 0.896E+02 0.908E+02	-0.128E+01	-4 18	-		0.001/
-20.00 0 -10.00 0 0.00 0	.883E+02 .896E+02 .908E+02	0.896E+02 0.908E+02	0 4076 04	-1.40	2	0.5067	0.0014
-10.00 0	.896E+02 .908E+02	0.908E+02	-U.123E+U1	-1.37	3	0.5184	0.0021
0.00 0	.908E+02		-0.118E+01	-1.30	4	0.5302	0.003(
	0045100	0.920E+02	-0.113E+01	-1.23	5	0.5420	0.0040
10.00 0	1.7216702	0.932E+02	-0.108E+01	-1.16	6	0.5538	0.005:
20.00 0	.934E+02	0,944E+02	-0.103E+01	-1.09	7	0.5656	0.0067
30.00 0	946E+02	0.956E+02	-0.976E+00	-1.02	8	0.5774	0.0084
40,00 0	.959E+02	0.968E+02	-0.924E+00	-0.95	x 9	0.5892	0.0104
.50.00 0	971E+02	0.980E+02	-0.872E+00	-0.89	10	0.6010	0.013(
60.00 0	984E+02	0.992E+02	-0.818E+00	-0.82	11	0.6128	0.016
70.00 (996E+02	0.100E+03	-0.763E+00	-0.76	12	0.6246	0.020(
80.00 0	0.101E+03	0.102E+03	-0.709E+00	-0.70	13	0.6364	0.025
90.00 0	102E+03	0.103E+03	-0.654E+00	-0.64	14	0.6481	0.030;
100.00 0	103E+03	0.104E+03	-0.599E+00	-0.58	15	0.6599	0.036
110.00 0	105E+03	0.105E+03	-0.545E+00	-0.52	16	0.6717	0.043
120.00 r	1.106E+03	D.106E+03	-0.4915+00	-0.46	17	0.6835	0.050
130.00 0	107E+03	0.107E+03	-0.440E+00	-0.41	18	0.6953	0.059
140.00 0	1.108E+03	0.109E+03	-0.390E+00	-0.36	19	0.7071	0.069
150.00 0	109E+03	0.110F+03	-0.344E+00	-0.31	² 20	0.7189	0.081
160.00 0	110E+03	0.111E+03	-0.2995+00	-0.27	21	0.7307	0.093
170.00 0	112E+03	0.112E+03	-0.259E+00	-0.23	22	0.7425	0.107
180.00 0	0.113E+03	0.113E+03	-0.2235+00	-0.20	23	0.7543	0.123
190.00 0	0.114E+03	0.114E+03	-0.193E+00	-0.17	24	0.7661	0.140
200.00 0	1.115E+03	0.115E+03	-0.1692+00	-0.15	25	0.7779	0.159
210.00 0	0.116E+03	D.116E+03	-0.150E+00	-0.13	26	0.7896	0.180
220.00 0	0.117E+03	0.117E+03	-0.139E+00	-0.12	27	0.8014	0.203
230.00 0	118E+03	0.118E+03	-0.136E+00	-0.11	28	0.8132	0.227
240.00 0	.119E+03	0.119E+03	-0.141E+00	-0.12	29	0.8250	0.254
250.00 0	120E+03	0.120E+03	-0.155E+00	-0.13	30	0.8368	D. 284
260.00 0	120E+03	0,121E+03	-0.179E+00	-0.15	31	0.8486	0.316
270.00 0	.121E+03	0 - 121E + 03	-0.213E+00	-0.18	32	0.8606	0.350
280.00 r	1.122E+03	0.122E+03	-0.2585+00	-0.21	77	0.0007	0 397
290.00 C	1.122E+03	0.1235+03	-0 3145+00	-0.21	34	0.0722	0.007
300.00 r	1.1235+03	0 1235+03	-0 3815+00	-0.31	76	0.0040	0.427
310.00 r	1.1235+03	0.1246+03	-0.3012+00	-0.37	× 33	0.0730	D 514
320.00 r	1.1245+03	0.1246+03	-0 5525+00	-0.37	30	0.7070	0.515
330.00 0	1.124E+03	0.1255+03	-0.3526+00	-0.44	79	0.7173	0.505
340.00 0	1.124E+03	0.1255-03	-0.7675-00	-0.55	20	D 9499	0.017
350.00 r	1.124E+03	0.1255+03	-0.8855100	-0.01	40	0.7447	0.0/4
360,00 . (1.123E+03	n.1245103	-0.10052+00	·-Ω.81	40 64	0.7547	0.733
370.00 r	1.1226+03	0.127ETU3	-0.1005101	-0.01	71 40	U.7003 N 0717	0./7/
380.00 1	1.120E+03	0.1215+03	-0.1056+01	-0.86	72	0.9703	0.005

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AAD= 0.5929967 SD1= 0.7286715 SD2= 7.2999690E-03

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**************** PRESSURE COMPARISON *****

FEMP Deg f	PRESSUR CALCD	E PSIA PUBLD	ABS Error	PERCENT Error	N	TR	PR
+0.00	0.951E+01	0.7312+01	0.201E+00	2.16	1	0.6054	0.0159
50.00	0.122E+02	0.120E+02	0.208E+00	1.73	2	0.6198	0.0205
20.00	0.155E+02	0.153E+02	0.207E+00	1.36	3	0.6342	0.0259
0.00	0.194E+02	0.192E+02	0.200E+00	1.04	4	0.6486	0.0325
0.00	0.240E+02	0.238E+02	0.186E+00	0.78	5	0.6631	0.0403
0.00	0.295E+02	0.293E+02	0.164E+00	0.56	6	0.6775	0.0494
20.00	0.359E+02	0.357E+02	0.136E+00	0.38	7	0.6919	0.0601
50.00	0.432E+02	0.431E+02	0.101E+00	0.23	8	0.7063	0.0725
0.00	0.517E+02	0.517E+02	0.617E-01	0.12	9	0.7208	0.0867
;0.00	0.614E+02	0.614E+02	0.193E-01	0.03	10	0.7352	0.1029
50.00	0.724E+02	0.724E+02	-0.253E-01	-0.03	s 11	0.7496	0.1213
70.00	0.848E+02	0.849E+02	~0.674E-01	-0.08	12	0.7640	0.1421
30.00	0.988E+02	0.989E+02	-0.107E+00	-0.11	13	0.7784	0.1655
70.00	0.114E+03	0.114E+03	-0.140E+00	-0.12	14	0.7929	0.1916
00.00	0.132E+03	0.132E+03	-0.161E+00	-0.12	15	0.8073	0.2206
10.00	0.151E+03	0.151E+03	-0.180E+00	-0.12	16	0.8217	0.2529
20.00	0.172E+03	0.172E+03	-0.182E+00	-0.11	17	0.8361	0.2884
30.00	0.196E+03	0.196E+03	-0.171E+00	-0.09	18	0.8506	0.3276
40.00	0.221E+03	0.221E+03	-0.145E+00	-0.07	19	0.8650	0.3705
30.00	0.249E+03	0.249E+03	-0.102E+00	-0.04	20	0.8794	0.4175
50.00	0.280E+03	0.280E+03	-0.415E-01	-0.01	21	0.8938	0.4687
70.00	0.313E+03	0.313E+03	0.280E-01	0.01	22	0.9083	0.5244
30.00	0.349E+03	0.349E+03	0.103E+00	0.03	23	0.9227	0.5849
70.00	0.388E+03	0.388E+03	C.176E+00	0.05	24	0.9371	0.6503
00.00	0.4302+03	0.430E+03	0.254E+00	0.06	25	0.9515	0.7210
10.00	0.476E+03	0.476E+03	0.309E+00	0.07	25	0.9660	0,7972
20.00	0.525E+03	0,524E+03	0.352E+00	0.07	27	0.9804	0.8792
30.00	0.577E+03	0.577E+03	0.351E+00	0.06	28	0.9948	0.9673
).D=	0 3439414						
10-	0.3420415				1 C C C C C C C C C C C C C C C C C C C		

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LIQUID DENSITY COMPARISON ***************

TABLE E2 CONTINUED

EMP	DENS IN Calcd	LB/FT3 PUBLD	ABS Error	PERCENT Error	N	TR	PR
0.00	0.101E+03	0.947E+02	0.605E+01	6.39	1	0.6054	0.0159
10.00	0.997E+02	0.937E+02	0.598E+01	6.38	2	0.6198	0.0205
0.00	0.986E+02	0.927E+02	0.590E+01	6.36	3	0.6342	0.0259
0.00	0.975E+02	0.917E+02	0.580E+01	6.32	4	0.6486	0.0325
0.00	0.963E+02	0.907E+02	0.567E+01	6.26	5	.6631	0.0403
0.00	0.951E+D2	0.896E+02	0.553E+01	6.17	6	0.6775	0.0494
0.00	0.939E+02	0.885E+02	⇒0.536E+D1	6.06	7	0.6919	0.0601
0.00	0.926E+02	0.874E+02	0.517E+01	5.91	8	0.7063	0.0725
0.00	0.912E+02	0.863E+02	0.495E+01	5.74	9	0.7208	0.0867
0.00	0.898E+02	0.851E+02	0.471E+01	5.53	10	0.7352	0.1029
0.00	0.884E+02	0.840E+02	0.439E+01	5.22	11	0.7496	0.1213
0.00	0.869E+02	0.827E+02	0.414E+01	5.00	12	0.7640	0.1421
0.00	0.853E+02	0.814E+02	0.380E+01	4.67	13	0.7784	0.1655
0.00	0.836E+02	0.801E+02	0.343E+01	4.28	14	0.7929	0.1916
0.00	0.818E+02	0.788E+02	0.303E+01	3.84	15	0.8073	0,2206
0.00	0.800E+02	0.774E+02	0.258E+01	3.34	16	0.8217	0.2529
0.00	0.78CE+02	0.759E+02	0.210E+01	2.76	17	0.8361	0.2884
0.00	0.759E+02	0.744E+02	0.157E+01	2,11	18	0.8506	0.3276
0.00	0.737E+02	0.727E+02	0.990E+00	1.36	19	0,8650	0.3705
0.00	0.714E+02	0.710E+02	0.360E+00	0.51	20	0.8794	Π.4175
0.00	0.689E+02	0.692E+02	-0.324E+00	-0.47	21	0.8938	0.4687
0.00	0.662E+02	0.672E+02	-0.107E+01	-1.59	22	0.9083	0.5244
0.00	0.632E+02	0.651E+02	-0.183E+01	-2.89	23	0.9227	0.5849
0.00	0.600E+02	0.627E+02	-0.276E+D1	-4.40	24	0.9371	0.6503
0.00	0.543E+02	0.600E+02	-0.371E+01	-6.18	25	0.9515	0.7210
0.00	0.521E+02	0.568E+D2	-0.473E+01	-8.32	26	0.9660	0.7972
0.00	0.469E+02	0:527E+02	-0.577E+01	-10.96	27	0.9804	0 4702
0.00	0.373E+02	0.458E+02	-0.644E+01	-14.07	28	.0.9948	0.9673

D =

1=

5.111201 4.369181 5.9726264E-02 2=

^{)1= 0.1826097})2= 6.6197840E-03

TEMP	H IN E	BTU/LB	 A8S	PERCENT	 N	TR	 P P
DEG F	CALCD	PUBLD	ERROR	ERROR	N		
40.00	0.000E+00	0.00CE+00	0.000E+00	0.00	1	0.6054	0.0159
30.00	0.201E+01	0.211E+01	-0.107E+00	-5.07	2	0.6198	0.0205
20.00	0.403E+01	0.424E+01	-0.202E+00	-4.76	3	0.6342	0.0259
10.00	0.609E+01	0.637E+01	-0.283E+00	-4.45	4	0.6486	0.0325
0.00	0.817E+01	0.852E+01	-0.352E+00	-4.13	5	0.6631	0.0403
10.00	0.103E+02	0.107E+02	-0.406E+00	-3.80	6	0.6775	0.0494
20.00	0.124E+02	0.129E+02	-0.447E+00	-3.48	7	0.6919	D.0601
30.00	0.146E+02	0.151E+02	-0.473E+00	-3.14	8	0.7063	0.0725
40.00	0.168E+02	0.173E+02	-0.486E+00	-2.81	9	0.7208	0.0867
20.00	0.190E+02	0.195E+02	-0.482E+00	-2.47	10	0.7352	0.1029
50.00	0.213E+02	0.218E+02	-0.466E+00	-2.14	a = 11	0.7496	0.1213
70.00	0.236E+02	0.240E+02	-0.434E+00	-1.80	12	0.7640	0.1421
30.00	0.260E+02	0.264E+02	-0.389E+00	-1.48	13	0.7784	0.1655
70.00	0.284E+02	0.287E+02	-0.330E+00	-1.15	14	0.7929	0.1916
00.00	0.308E+02	0.311E+02	-0.258E+00	-0.83	15	0.8073	0.2206
10.00	0.334E+02	0.335E+02	-0.172E+00	-0.51	16	0.8217	0.2529
20.00	0.359E+02	0.360E+02	-0.7462-01	-0.21	17	0.8361	0,2884
30.00	0.386E+02	0.386E+02	0.358E-01	0.09	18	0.8506	0.3276
+0.00	0.413E+02	0.412E+02	0.157E+00	0.38	19	0.8650	0.3705
30.00	0.441E+02	0.438E+02	0.292E+00	0.67	20	0.8794	D.4175
20.00	0.471E+02	0.466E+02	0.4392+00	0.94	21	0.8938	0.4687
10.00	0.501E+02	0.495E+02	0.601E+00	1.21	22	0.9083	0.5244
30.00	0.533E÷02	0.5262+02	0.784E+00	1.49	23	0.9227	0.5847
10.00	0.568E+02	0.5586+02	0.992E+00	1.78	24	0,9371	0.6503
10.00	0.604E+02	0.5922+02	0.124E+01	2.10	25	0.9515	0.7210
0.00	0.645E+02	0-630E+02	0.156E+01	2.48	26	0,9660	0.7973
20.00	0.693E+02	0.672E+02	0.202E+01	3.01	27	0.9804	0.8792
10.00	0.757E+02	0.729E+02	0.282E+01	3.88	28	0.9948	0.9573
)D=	2.152249				(14)		
11=	0.8692907						

- 12= 2.6591945E-02
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- APOUR ENTHALPY
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TABLE E2 CONTINUED

EMP Eg F	BTU/LB Calcd	PUBLD	ABS Error	PERCENT	N	TR	PR
	0 7225+02	. 7005.00	0.0000.00				
	0.7222702	0.729E+U2	~0.725E+00	-0.99	1	0.6054	0.0159
0.00	0.7545+02	0.740E+02	-0.833E+00	-0.86	2	0.6198	0.0205
0.00	0.7575+02	0.7516+02	-0.541E+00	-0.72	3	0.6342	0.0259
	0.7572+02	0.7022+02	-U.449E+UU	-0.59	4	0.6486	0.0325
	0.7076+02	0.773E+02	-0.357E+00	-0.46	5	0.6631	0.0403
	0.701E+U2	0.783E+02	-0.268E+00	-0.34	6	0.6775	0.0494
	U.792E+U2	0.794E+02	-0.180E+00	-0.23	7	0.6919	0.0601
0.00	0.803E+02	0.804E+02	-0.955E-01	-0.12	8	0.7063	0.0725
0.00	0.814E+02	0.814E+02	-0.156E-01	-0.02	9	0.7208	0.0867
00.0	U.825E+02	0.824E+02	0.597E-01	0.07	10	0.7352	0.1029
0.00	0.835E+02	0.834E+02	0.128E+00	0.15	11	0.7496	0,1213
1.00	0.845E+02	0.844E+02	0.191E+00	0.23	12	0.7640	0.1421
0.00	0.855E+02	0.853E+02	0.244E+00	0.29	13	0.7784	0.1655
1.00	0.865E+02	0.862E+02	0.289E+00	0.34	14	0,7929	0.1916
3.00	0.874E+02	0.870E+02	0.326E+00	0.37	15	0.8073	0 2204
0.00	0.882E+02	0.878E+02	0.351E+00	0.40	16	0.8217	0.2525
.00	0.890E+02	0.886E+02	0.367E+00	0.41	17	0.8361	D. 2884
0.00	0.897E+02	0.893E+02	0.371E+00	0.42	18	0.8504	0.2004
).00	0.903E+02	0.900E+02	0.365E+00	0.41	19	0.8650	0.3276
0.00	0.909E+02	0.905E+02	0.348E+00	0.38	20	0.8794	0.3703
0.00	0.913E+02	0.910E+02	0.321E+00	0.35	21	0.8938	0.4175
0.00	0.916E+02	0.914E+02	0.287E+00	0.31	22	0.0700	0.4007
00.00	0.918E#02	0.916E+02	0.247E+00	0.27	23	0.9227	0.5247
0.00	0.918E+D2	0.916E+02	0.209E+00	0.23	24	0.9371	0.3047
).00	0.915E+02	0,913E+02	0.183E+00	0.20	25	0 9515	0.0000
.00	0.908E÷02	0.906E+02	0.195E+00	0.22	26	0.7313	0.7210
1.00	0.894E+02	0.89CE+02	0.320E+00	0.36	27 8	b.7000	0.7772
.00	0.861E+02	0.851E+02	D.969E+00	1.14	22	0.7004	0.8792

0.3883371

≈ 0.3858528

= 4.7423360E-03

***** LATENT HEAT COMPARISON

TABLE E2 CONTINUED

TEMP Deg f	LH IN B Calcd	TU/LB PUBLD	ABS Error	PERCENT Error	N	TR	PR
40,00	0.722E+02	0.729E+02	-0.725E+00	-0.99		0.6054	0.0159
30.00	0.714E+02	0.719E+02	-0.526E+00	-0.73	2	0.6198	0.0205
20.00	0.705E+02	0.709E+02	-0.339E+00	-0.48	3	0.6342	0.0259
10.00	0,697E+02	0.698E+02	-0.165E+00	-0.24	4	0.6486	0.0325
0.00	0.687E+02	0.688E+02	-0.500E-02	-0.01	5	0.6631	0.0403
10.00	0.678E+02	0.677E+02	0.139E+00	0.21	6	0.6775	0.0494
20.00	0.668E+02	0,665E+02	0.267E+00	0.40	7	0.6919	0.0601
30.00	0.657E+02	0.654E+02	0.378E+00	0.58	8	0.7063	0.0725
40.00	0.646E+02	0.642E+02	0.470E+00	0.73	9	0.7208	0.0867
50.00	0.635E+02	0.629E+02	0.542E+00	0.86	10	0.7352	0.1029
60.00	0.622E+02	0.616E+02	0.594E+00	0.96	11	0.7496	0.1213
70.00	0.609E+02	0.603E+02	0.625E+00	1.04	12	0.7640	0.1421
80.00	0.596E+02	0.5892+02	0.634E+00	1.08	13	0.7784	0.1655
90.00	0.581E+02	0.575E+02	D.619E+00	1.08	14	0.7929	0.1916
00.00	0.565E+02	0.559E+02	0.583E+00	1.04	15	0.8073	0.2206
10.00	0.548E+02	0.543E+02	0.523E+00	0.96	16	0.8217	0.2529
20.00	0.5302+02	0.526E+02	0.441E+00	0.84	17	0.8361	0.2884
30,00	0.511E+02	0.508E+02	0.336E+00	0.66	18	0.8506	0.3276
40.00	0.490E+02	0.488E+02	0.207E+00	0.42	19	0.8650	0.3705
50,00	D.467E+02	0.467E+02	0.558E-01	0.12	20	0.8794	0.4175
60.00	0,443E+02	0.444E+02	-0.118E+00	-0.27	21	0.8938	0.4687
70.00	0.415E+02	0.418E+02	-0.315E+00	-0.75	22	0.9083	0.5244
80.00	0.385E+02	0.370E+02	-0.537E+00	-1.38	23	0.9227	0.5849
90.00	0.350E+02	0.358E+02	-0.783E+00	-2.19	. 24	0.9371	0.6503
00.00	0.310E+02	0.321E+02	-0.106E+01	-3.30	25	0.9515	0.7210
10.00	0,262E+D2	0.276E+02	-0.137E+01	-4.95	25	0.9660	0.7972
20.00	0.201E+02	0.218E+02	-0.170E+01	-7.82	27	0.9804	0.8792
30.00	0.104E+02	0.122E+02	-0,186E+D1	-15.17	28	0.9948	0.9673

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1.759213 D.7337487 3.5766512E-02 AD=

01= D2=

TEMP DEG F	BTU/ Calcd	LB F PUBLD	ABS Error	PERCENT Error	N	TR	PR
40.00	0.000E+00	0.000E+00	0.000£+00	0.00	1	0.6054	0.0159
30.00	0.510E-02	0.496E-02	0.140E-03	2.82	2	0.6198	0.0205
20.00	0.101E-01	0.983E-02	0.306E-03	3.11	3	0.6342	0.0259
10.00	0.151E-01	0.146E-01	0.491E-03	3.36	4	0.6486	0.0325
0.00	0.200E-01	0.193E-01	0.713E-03	3.69	5	0.6631	0.0403
10.00	0.249E-01	0.239E-01	D.955E-03	3.99	6	0.6775	0.0494
20.00	0.297E-01	0.285E-01	0.121E-02	4.25	7	0.6919	0.0601
30.00	0.345E-01	0.3306-01	0.151E-02	4.57	8	0.7063	0.0725
40.00	0.393E-01	0.375E-01	0.182E-02	4.87	9	0.7208	0.0867
50.00	0.440E-01	0.418E-01	D.216E-02	5.15	10	0.7352	0.1029
60.00	0.487E-01	0.462E-01	0.252E-02	5.45	11	0.7496	0.1213
70.00	0.534E-01	0.505E-01	0.290E-02	5.74	12	0.7640	0.1421
80.00	0.580E-01	0.547E-01	0.329E-02	6.02	13	0.7784	0.1655
90.00	0.627E-01	0.590E-01	0.372E-02	6.30	14	0.7929	0.1916
00.00	0.674E-01	0.632E-01	0.414E-02	6.55	15	0.8073	0.2206
10.00	0.720E-01	0.675E-01	D.459E-02	6.81	16	0.8217	0.2529
20.00	0.7676-01	0.717E-01	0.505E-02	7.05	17	0.8361	0.2884
30.00	0.815E-01	0.759E-01	0.552E-02	7.28	18	0.8506	0.3276
40.00	0.862E-01	0.802E-01	0.601E-02	7.49	19	0.8650	0.3705
50.00	0.910E-01	0.845E-01	0.651E-02	7.71	20	0.8794	0.4175
60.00	0.960E-01	0.889E-01	0.702E-02	7.89	21	0.8938	0.4687
70.00	0.101E+00	0.934E-01	0.755E-02	8.08	22	0.9083	0.5244
80.00	0.106E+00	0,980E-01	0.809E-02	8.25	23	0.9227	0.5849
90.00	0.112E+00	0.103E+00	0.867E-02	8.43	24	0.9371	0.6503
00.00	0.117E+00	0.108E+00	0.929E-02	8.61	25	0.9515	0.7210
10.00	0.123E+00	0:113E+00	0.100E-01	8.84	26	0.9660	0.7972
20.00	0.130E+00	0.1192+00	0.109E-01	9.14	27	0.7804	0.8792
30.00	0.140E+CO	0.127E+00	0.123E-01	9.65	28	0.9948	0.9673
40-	1 440447				der i		

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TABLE E2 CONTINUED

TEMP	BTU/	LB F	ABS	PERCENT	N	TR	PR
DEG F	CALCD	PUBLD	ERROR	ERROR			
-40.00	0.172E+00	0.174E+00	-0.161E-02	-0.93	1	0.6054	0.0159
-30.00	0.171E+00	0.172E+00	-0.965E-03	-0.56	2	0.6198	0.0205
-20.00	0.171E+00	0.171E+00	-0.356E-03	-0.21	3	0.6342	0.0259
.10.00	0.170E+00	0.170E+00	0.231E-03	0.14	4	0.6486	0.0325
0.00	0.170E+00	0.169E+00	0.801E-03	0.47	5	0.6631	0.0403
10.00	0.169E+00	0.168E+00	0.135E-02	0.80	6	0.6775	0.0494
20.00	0.169E+00	0.167E+00	0.187E-02	1.12	7	0.6919	0.0601
30.00	0.169E+00	0.166E+00	0.238E-02	1.43	8	0.7063	0.0725
40.00	0.169E+00	0.166E+00	0.285E-02	1.72	9	0.7208	0.0867
50.00	0.169E+00	0.165E+00	0.330E-02	2.00	10	0.7352	0.1029
60.00	0.169E+00	0.165E+00	0.374E-02	2.27	11	0.7496	0.1213
70.00	0.168E+00	0.164E+00	0.415E-02	2.53	12	0.7640	0.1421
80.00	0.168E+00	0.164E+00	0.454E-02	2.77	13	0.7784	0.1655
90.00	0.168E+00	0.164E+00	0.491E-02	3.00	14	0.7929	0.1916
.00.00	0.168E+00	0.163E+00	0.526E-02	3.22	15	0.8073	0.2206
.10.00	0.168E+00	0.163E+00	0.557E-02	3.42	16	0.8217	0.2529
.20.00	0.168E+00	0.162E+00	0.588E-02	3.62	17	0.8361	0.2884
130.00	0.168E+00	0.162E+00	0.616E-02	3.80	18	0.8506	0.3276
40.00	0.168E+00	0.162E+00	0.641E-02	3.97	19	0.8650	0,3705
.50.00	0.168E+00	0.161E+00	0.666E-02	4.13	20	0.8794	0.4175
.60.00	0.167E+00	0.161E+00	0.6886-02	4.29	21	0.8938	0.4687
70.00	0.167E+00	0.160E+00	0.709E-02	4.44	22	0.9083	0.5244
80.00	0.166E+00	0.160E+00	0.677E-02	4.24	23	0.9227	0.5849
.90.00	0.165E 1 00	0.158E+00	0.750E-02	4.75	24	0.9371	0.6503
200.00	0.164E+00	0.157E+00	0.772E-02	4.93	25	0.9515	0.7210
10.00	0.163E+00	0,155E+00	0.800E-02	5.18	26	0.9660	0.7972
120.00	0.160E+00	0.1512+00	0.843E-02	5.56	27	0.9804	0.8792
130.00	0.155E+00	0.145E+00	0.961E-02	6.62	28	0.9948	0.9673

AD= 2.932905 D1= 5.4896330E-03

1

102= 3.4833507E-02

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TEMP	Ζ		ABS	PERCENT	N	TR	PR
DEGF	CALCD	PUBLD	ERROR	ERROŖ			
40 00	0 2535-02	0 2445-02	-0 1045-07	-4 00		0 405/	0.0460
30.00	0.2332-02	0.7745-02	-0.1082-03	-4.00	2	0.6034	0.0157
20.00	0.5216.02	0.3386-02	-0.147E-03	-4 70	4	0,0170	0.0205
10.00	0.402E-02	0.9226-02	-0.1772-03	-4 92	3	0.6342	0.0257
0.00	0.4702 02	0.5276-02	-0.2302-03	-5 24		0.0400	0.0325
10.00	B.744E-02	0.0456-02	-0.3382-03	-3:41	3	0 4775	0.0403
20 00	0.1992 02	0.7032-02	-0.4136-03	-5,20	7	0.0773	0.0474
	0.0772-02	0.4465-02	-0.5002-03	-5.30	/	0.8717	0.0601
40.00	0,1072-01	0.1142-01	-0.0152-03	-2.42		0.7083	0.0725
50.00	0.1202-01	0.1352-01	~0.7202-03	-5.33	9	0.7208	0.0867
40.00		0.1572-01	-0.0332-03	-5.22	10	0.7352	0.1029
		0.18/2-01	-U.951E-U3	-5.08	11	0.7496	0.1213
70.00	U.208E-01	U.218E-01	-0.106E-02	-4.86	12	0.7640	0.1421
00.00	U.242E-01	0.253E-01	-0.116E-02	-4.57	13	0.7784	0.1655
90.00	0.280E-01	0.293E-01	-0.124E-02	-4.23	14	0.7929	0.1916
00.00	0.324E-01	0.337E-01	-0.129E-02	-3.83	15	0.8073	0.2206
10.00	0.373E-01	0.386E-01	-0.130E-02	-3.38	16	0.8217	0.2529
20.00	0.429E-01	0.441E-01	-0.124E-02	-2.81	17	0.8361	0.2884
30.00	0.492E-01	0.503E-01	-0.110E-02	-2.18	18	0.8506	0.3276
40.00	0.563E-01	0.572E-01	-0,815E-03	-1.43	19	0.8650	0.3705
50.00	0.645E-01	0.649E-01	-0.366E-03	-0.56	20	0.8794	0.4175
60.00	0.738E-01	0.735E-01	0.324E-03	0.44	21	0.8938	0.4687
70.00	0.846E-01	0.833E-01	0.134E-02	1.61	22	0.9083	0.5244
80.00	0.972E-01	0.944E-01	0.282E-02	2.99	23	0.9227	0.5849
90.00	0.1125+00	0.107E+00	0.494E-02	4.61	24	0.9371	0.6503
00.00	0.13CE+00	0.122E+00	0.811E-02	6.63	25	0.7515	0.7210
10.00	0.154E+00	0.141E+00	0.128E-01	9.12	26	0,9660	0.7972
20.00	0.1855+00	0.165E+00	0.204E-01	12.36	27	0.9804	0.8792
30.00	0.240E+00	0.206E+00	0.338E-01	16.42	28 8	0.7748	0,9673
4							
AD=	4.892305		*				
01=	8.2559334E-0	3					
02=	5.9710901E-0	2					
*****	**********	****					
VAPOU	R COMPRESSIBI	LITY				TABLE E 2 CON	TINUED
*****	********	****					

18

TEMP		Ζ	ABS	PERCENT	N	TR	PR
JEG F	CALCU	PUBLD	ERRUR	ERNUR			
+0.00	0.976E+00	0.968E+00	0.7396-02	0.76	1	0.6054	0.0159
30.00	0.971E+00	0.962E+00	0.816E-02	0.85	2	0.6198	0.0205
20.00	0.965E+00	0.956E+00	0.885E-02	0.93	3	0.6342	0.0259
10.00	0.958E+00	0.948E+00	0.945E-02	1.00	4	0.6486	0.0325
0.00	0.950E+00	0.941E+00	0.9966-02	1.06	5	0.6631	0.0403
10.00	0.942E+00	0.932E+00	0.104E-01	1.11	6	0.6775	0.0494
20.00	0.933E+00	0.922E+00	0.106E-01	1.15	7	0.6919	0.0601
30.00	0.923E+00	0.912E+00	0.107E-01	1.18	8	0.7063	0.0725
40.00	0.912E+00	0.901E+00	0.107E-01	1.18	9	0.7208	0.0867
50.00	0.900E+00	0.889E+00	0.104E-01	1.17	10	0.7352	0.1029
50.00	0.887E+00	0.877E+00	0.999E-02	1.14	11	0.7496	0.1213
70.00	0.873E+00	0.863E+00	0.938E-02	1.09	12	0.7640	0.1421
30.00	0.858E+00	0.849E+00	0.859E-02	1.01	13	0.7784	0.1655
70.00	0.841E+00	0.834E+00	0.761E-02	0.91	14	0.7929	0.1916
00.00	0.824E+00	0.817E+00	0.647E-02	0.79	15	0.8073	0.2206
10.00	0.805E+00	0.800E+00	0.513E-02	0.64	16	0.8217	0.2529
20.00	0.785E+00	0.781E+00	0.368E~02	0.47	17	0.8361	0.2884
30.00	0.764E+00	0.761E+00	0.217E-02	0.28	18	0.8506	0.3276
40.00	0.741E+00	0.740E+00	0.389E-03	0.05	19	0.8650	0.3705
50.00	0.716E+00	0.717E+00	-0.130E-02	-0.18	20	0.8794	0.4175
50.00	0.689E+00	0.692E+00	-0.298E-02	-0.43	21	0.8938	0.4687
70.00	0.660E+00	0.665E+00	-0.459E-02	-0.69	22	0.9083	0.5244
30.00	0.629E+00	0.635E+00	-0.593E-02	~0.93	23	0.9227	0.5849
70.00	0.595E+00	0.602E+00	-0.696E-02	-1.16	24	0.9371	0.6503
10.00	0.556E+00	0.564E+00	-0.722E-02	-1.28	25	0.9515	0.7210
10.00	0.512E+00	0.519E+00	-0.655E-02	-1.26	26	0.9660	0.7972
20.00	0.459E+00	0:462E+00	-0.340E-02	-0.74	27	0.9804	0.8792
30.00	0.379E+00	D.372E+00	0.750E-02	2.04	28 🔬	0.9948	0.9673

AD= 0.9106935)1= 7.7746701E-03

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)2= 1.0100145E-02

***************** PRESSURE COMPARISON

a.

TEMP DEG F	PRESSURE PSI Calco Publ	LA _D	ABS Error	PERCENT Error	N	TR	PR
						0.7721	0.1553
-40.00	0.871E+02 0.874	+E+02 -	0.291E+00	-0.35	2	0.7905	0.1876
-30.00	0.105E+03 0.104	5E+03 -	0.366E+00	-0.35	<u>-</u> 7	0.8089	0.2246
-20.00	0.126E+03 0.124	6E+03 -	-0,410E+00	-0.32	5	0.8273	0.2667
-10.00	0.150E+03 0.150	JE+03 -	0.46UE+UU	-0.31	5	0.8457	0.3145
0.00	0.176E+03 0.17	/E+U3 -	-U.3/4E+UU	-0.21	4	0.8641	0.3683
10.00	0.207E+03 0.20	/E+U3 ~	0.2012700	-0.10	7	0.8825	0.4285
20.00	0.2402+03 0.240	UE+U3	0.1725-01	0.01	8	0.9009	0.4958
30.00	0.2782+03 0.276	DETU3 DE107	0.2302+00	0.15	9	0.9193	0.5705
40.00	0.320E+03 0.320	15403 15403	0.4782+00	0.16	10	0.9376	D.6533
50.00	0.3662+03 0.360	0E+U3 7E+07	0.5852,00	0.16	11	0.9560	0.7445
60.00	0.4102+03 0.41	76703	0.5886+00	0.12	12	0.9744	0.8449
/0.00	0.4/42103 0.4/	15102	0.2345+00	0.04	13	0.9928	0.9549
80.00	0.5362703 0.93	02703	0.2046.00				
AAD=	0.1813645						
SD1= SD2=	2.2072275E-03						
*****	****	**	ē))				
LIQUID	DENSITY COMPARIS	ON				TABLE E3	CONTINUED
*****	***************	**					
	DENS IN LB/F		ABS	PERCENT	N	TR	PR
DEG F	CALCD PUB	BLD	ERROR	ERROR			
	0 890E+02 0.84		0.490E+01	5.83	· 1	0.7721	0.1553
	0.8702702 0.87	24E+02	0.438E+01	5.31	' 2	0.7905	0.1876
-20.00	0.8455+02 0.80	17E+02	0.379E+01	4.69	3	0.8089	0.2246
~10.00	0.8205+02 0.00	9E+02	0.316E+01	4.00	4	0.8273	0.2667
-10.00	0.7945+02 0.77	70E+02	0.238E+01	3.09	5	0.8457	0.3145
10.00	0.765E+02 0.74	9E+02	0.158E+01	2.11	6	0.8641	0.3683
20.00	0.734E+02 0.72	27E+02	0.652E+00	0.90	7	0.8825	0.4285
30.00	0.700E+02 0.70	3E+02	-0.342E+00	-0.49	8	0.9009	0.4958
40.00	0.662E+02 0.67	77E+02	-0.149E+01	-2.20	9	0.9193	0.5705
50.00	0.620E+02 0.64	47E+02	-0.273E+01	-4.21	10	0.9376	0.6533
60.00	0.5706+02 0.61	11E+02	-0.408E+01	-6.67	11	0.9560	0.7445
70.00	0.5076+02 0.56	55E+02	-0.552E+01	-9.78	12	0.9744	0.8449
80.00	0.421E+02 0.48	88E+02	-0.674E+01	-13.79	13	0.9928	0.9549
AAD≖	4.852096						
SD1=	3.855265						
SD2=	6.2464975E-02						
	*************			State of states and			
LATEN	T HEAT COMPARISON					TABLE E3	CONTINUED
*****	*****						
DEG F	CALCD PU	B Bld	ABS Error	PERCENT	N	TR	РЯ
	0.545E+UZ 0.54	4UE+02	U.437E+00	0.81	1	U.7721	0.1553
-30.00	0.5282+02 0.5	24E+U2	0.380E+00	0.73	2	0.7905	U.18/6
-10.00	0.510E+02 0.50	0/E+U2	0.321E+00	0.63	3	0.8089	0.2248
-10.00	0 4495400 0 4	0027UZ	0.3712400	0.80	* E	0.8273	0.74/5
10.00	0.707E7U2 U.4	006+U2	U.22/E+UU	0.47	5	0.045/	U.3145
20.00	1 6485102 0.44	736702	U:720C-U2	0.02	5	0.0041	0.3003
20.00	0.3805102 U.4	216402	-U.253E+UU	-0.60	. /	0.0025	U.4285 0 /050
20.00	0,0076702 0.3 0 3656102 0.3	1251UZ	-0.3712700	-2 44	0	0,7007	U.1730 N E705
50.00	0.314F+02 0.3	30E+05	-0.7506700	-4.01	10	0.7173 N 0774	0.3703
20.00	0.241E102 0.3	875102	-0 1975-01	- 7:47 _1 E4	44	0.73/0 0 0610	0.0333 N 7445
70 00	0.2005102 0.2	0/ETU2	-0.10/2701	-40 4/	1-1	0.7300	U:/**5
20.00	0 4406+02 0 41	346+00	-U.231E+U1	-10.04	12	0.7/44	0.05/0
00.00	0.1106702 0.1	JOETUZ	-0.2532+01	-10.62	1.3	0.7728	0.7547
AAD=	3.722181				2. 2		2
301-	4+473004						

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SD2= 6.6852726E-02

***** PRESSURE COMPARISON *****

TABLE E4

TABLE E4 CONTINUED

TEMP DEG F	PRESSUR	E PSIA PUBLD	ABS Error	PERCENT Error	N	TR	PR
							0 0554
40.00	0.319E+02	0.319E+02	-0.591E-02	-0.02	1	0.0054	0.0337
30.00	0.395E+02	0.396E+02	-0.541E-01	-0.14	2	0.7018	0.0007
20.00	0.485E+02	0.486E+02	-0.988E-01	-0.20	3	0.7181	0.0843
10.00	0.589E+02	0.591E+02	-0.159E+00	-0.27	4	0.7344	0.1025
0.00	0 7105+02	0.712E+02	-0.209E+00	-0.29	5	0.7508	0.1234
40.00	0.8485+02	0 8505+02	-0.24AE+00	-0.29	6	0.7671	0.1474
10.00	0.0905+02	0.0000002	-0 3275+00	-0.32	7	0.7834	0.1747
20.00	0.1002403	0.1012+03	-0.3205+00	-0.28	8	0.7998	0.2057
30.00	U.118E+U3	0.1172703	0.3302+00	-0.20	ä	0.8161	0.2405
40.00	0.138E+03	U.137E+U3	-0.2076+00	-0.45	10	0.8324	0.2796
50.00	0.161E+03	0.161E+03	-U.234ETUU	-0.15	10	<u>п 8488</u>	0.3231
60.00	0.186E+03	0.186E+03	-0.190E+00	-0.10	11	0.0400	0 3715
70.00	0.214E+03	0.214E+03	-0.755E-D1	-0.04	12	0.0001	0.2715
80.00	0.244E+03	0.244E+03	-0.465E-02	0.00	13	0.8814	0.4230
90.00	0.278E+03	0.278E+03	0.215E+00	0.08	14	0.8978	0.4040
00.00	0.316E+03	0.315E+03	0.383E+00	0.12	15	0.9141	0.5488
10.00	0.356E+03	0.3566+03	0.505E+00	0.14	16	0.9304	0.6198
20 00	0 4015+03	0.4005+03	0.597E+00	0.15	17	0.9468	0.6974
70.00	0.4012.03	0 44002+03	D. 682E+00	0.15	18	0.9631 -	0.7819
30.00	0.4302+03	0.5025+03	0.0012.00	0.12	19	D.9794	0.8737
40.00	0.5026703	0.5026703	0.3726+00	0.08	20	0.9958	0.9733
50.00	0.5602+03	0.5572+03	0.4/32700	0100	20	01//02	
AD=	0.1575302	•			2		

0.3540626 ;D1=

102= 1.8763094E-03

***** LIQUID DENSITY COMPARISON *************************

TEMP	DENS IN	LB/FT3 PUBLD	ABS Error	PERCENT Error	 N	TR	PR
							0.0554
40.00	0.127E+03	0.119E+03	0.7502+01	6.29	-	0.0034	0.0487
30.00	0.125E+03	0.118E+03	0.721E+01	6.13		0.7010	0.0007
20.00	0.123E+03	0.116E+03	0.688E+01	5.93	3	0./181	0.0043
10.00	0.121E+03	0.114E+03	0.649E+01	5.68	- 4	0.7344	0.1025
0.00	0.119E+03	0.112E+03	0.605E+01	5.38	5	0.7508	0.1234
10.00	0.116E+03	0.111E+03	0.556E+01	5.03	6	0.7671	0.1474
20 00	0.1146+03	0.109E+03	0.501E+01	4.61	7	0.7834	0.1747
30.00	0.111E+03	0.107E+03	0.439E+01	4.11	8	0.7998	0.2057
40.00	0.10AE+03	0.105E+03	0.373E+01	3.56	9	0.8161	0.2405
50.00	0.105E+03	0.102E+03	0.295E+01	2.88	10	0.8324	0.2796
60.00	0.102E+03	0.100E+03	0.213E+01	2.13	11	0.8488	0.3231
70.00	D.990E+02	0.978E+02	0.121E+01	1.24	12	0.8651	0.3715
80.00	0.954E+02	0.952E+02	0.202E+00	0.21	13	0.8814	0.4250
90.00	0.915E+02	0.924E+02	-0.905E+00	-0.98	14	0.8978	0.4840
00.00	0.873E+02	0.894F+02	-0.211E+01	-2.36	15	0.9141	0.5488
10.00	0.826F+02	0.861E+02	-0.344E+D1	-4.00	16	0.9304	0.6198
20.00	0.773E+02	0.822E+02	-0.489E+01	-5.95	17	0.9468	0.6974
30.00	0.712E+02	D.776E+02	-0.645E+01	-8.30	18	0.9631	0.7819
40.00	0 4755+02	0 7145+02	-0 8055+01	-11 25	19	0.9794	0.8737
50.00	0.519E+02	0.607E+02	-0.886E+01	-14.59	20	0.9958	0.9733

3D2=

6.2005714E-02

TABLE E5

TEMP DEG F	PRESSUR Calcd	E PSIA PUBLD	ABS Error	PERCENT Error	N	TR	PR
-40 00	n 1545+02	0 1525+02	0 4765+00	0 89		0 4744	0 0213
-30.00	0.1978+02	0.1946+02	0.135E+00	0.64	2	0.0310 N 4444	0.0213
-20.00	0.250E+02	0.248E+02	0.1075+00	0.43	3	0.6617	0.0346
-10.00	0.312E+02	0.312E+02	0.773E-01	0.25	4	0.6767	0.0433
0.00	0.387E+02	0.387E+02	0.355E-01	0.09	5	0.6918	0.0536
10.00	0.474E+02	0.475E+02	-0.151E-01	-0.03	6	0.7068	0.0657
20.00	0.577E+02	0.577E+02	-0.747E-01	-0.13	7	0.7219	0.0799
30.00	0.695E+02	0.696E+02	-0.137E+00	-0.20	8	0.7369	0.0962
40.00	0.830E+02	0.832E+02	-0.198E+00	-0.24	9	0.7520	0.1150
50.00	0.985E+02	0.987E+02	-0.250E+00	-0.25	10	0.7670	0.1364
60.00	0.116E+03	0.116E+03	-0.282E+00	-0.24	11	0.7821	0.1607
70.00	0.136E+03	0.136E+03	-0.286E+00	-0.21	12	0.7971	0.1882
	U.150E+U3	U.158E+U3	-0.259E+00	-0.16	13	0.8122	0.2190
100.00	0.1032703	0.2445+03	-0.165E+00	-0.09	14	0.82/2	0.2534
110.00	0.241E+03	0.241E+03	0.2045+00	0.01	15	0.0423	0.2717
120.00	0.275E+03	0.275E+03	0.2042+00	0.08	17	0.8373	0.3342
130.00	0.312E+03	0.312E+03	0.877E+00	0.28	18	0.8874	0.0011
140.00	0.353E+03	0.352E+03	0.134E+01	0.38	19	0.9025	0.4894
150.00	0.398E+03	0.396E+03	0.183E+01	0.46	20	0.9175	0.5513
160.00	0.447E+03	0.445E+03	0.232E+01	0.52	21	0.9326	0.6190
170.00	0.500E+03	0.497E+03	0.274E+01	0.55	22	0.9476	0.6926
180.00	0.558E+03	0.555E+03	0.294E+01	0.53	23	0.9627	0.7726
190.00	0.620E+03	0.618E+03	0.268E+01	0.43	24	0.9777	0.8592
200.00	0.688E+03	0.686E+C3	0.157E+01	0.23	25	0.9928	0.9529
AAD= 5D1= 5D2=	0.3008717 1.254800 3.7347944E-03	3 *					
: *****	***********	******					
LIQUIE	DENSITY COM	PARISON	×			TABLE E5	CONTINUED
LIQUIC ******	D DENSITY COM	PARISON	¥			TABLE E5	CONTINUED
LIQUIC ****** TEMP DEG F	DENSITY COM Dens in Calcd	PARISON ******* LB/FT3 PUBLD	ABS Error	PERCENT Error	 N	TABLE E5	CONTINUED
TEMP DEG F	DENSITY COM DENS IN CALCD 0.899E+02	PARISON ****** LB/FT3 PUBLD 0.880E+02	ABS Error 0.187E+01	PERCENT ERROR 2.12	 N 1	TABLE E5	CONTINUED PR 0.0213
LIQUIC ****** DEG F -40.00 -30.00	DENSITY COM DENS IN CALCD 0.899E+02 0.888E+02	PARISON ****** LB/FT3 PUBLD 0.880E+02 0.870E+02	ABS ERROR 0.187E+01 0.185E+01	PERCENT ERROR 2.12 2.13	N 1 2	TABLE E5	CONTINUED PR 0.0213 0.0273
LIQUIC ****** DEG F -40.00 -30.00 -20.00	DENSITY COM DENSIN CALCD 0.899E+02 0.888E+02 0.878E+02 0.878E+02	PARISON ****** LB/FT3 PUBLD 0.880E+02 0.870E+02 0.860E+02	ABS ERROR 0.187E+01 0.185E+01 0.181E+01	PERCENT ERROR 2.12 2.13 2.11	N 1 2 3	TABLE E5 TR 0.6316 0.6466 0.6617	CONTINUED PR 0.0213 0.0273 0.0346
LIQUIC ****** DEG F -40.00 -30.00 -20.00 -10.00	DENSITY COM DENSIN CALCD 0.899E+02 0.888E+02 0.878E+02 0.867E+02 0.855E+02	PARISON ****** LB/FT3 PUBLD 0.880E+02 0.870E+02 0.860E+02 0.860E+02 0.849E+02 0.849E+02	ABS ERROR 0.187E+01 0.185E+01 0.181E+01 0.175E+01	PERCENT ERROR 2.12 2.13 2.11 2.06 4 98	N 1 2 3 4 5	TABLE E5 TR 0.6316 0.6466 0.6467 0.6767	CONTINUED PR 0.0213 0.0273 0.0346 0.0433 0.0574
LIQUIC ****** TEMP DEG F -40.00 -30.00 -20.00 -10.00 0.00	DENSITY COM DENSIN CALCD 0.899E+02 0.888E+02 0.878E+02 0.867E+02 0.855E+02 0.855E+02 0.843F+02	PARISON ****** LB/FT3 PUBLD 0.880E+02 0.870E+02 0.860E+02 0.860E+02 0.849E+02 0.838E+02 0.838E+02	ABS ERROR 0.187E+01 0.185E+01 0.181E+01 0.175E+01 0.166E+01	PERCENT ERROR 2.12 2.13 2.11 2.06 1.99 1.88	N 1 2 3 4 5 4	TABLE E5 TR 0.6316 0.6466 0.6466 0.6617 0.6767 0.6718 0.7048	CONTINUED PR 0.0213 0.0273 0.0346 0.0433 0.0536
LIQUIC ****** TEMP DEG F -40.00 -30.00 -20.00 -10.00 0.00 10.00 20.00	D DENSITY COM DENSIN CALCD 0.899E+02 0.888E+02 0.878E+02 0.867E+02 0.855E+02 0.843E+02 0.830E+02	PARISON ****** LB/FT3 PUBLD 0.880E+02 0.870E+02 0.860E+02 0.849E+02 0.838E+02 0.827E+02 0.816F+02	ABS ERROR 0.187E+01 0.185E+01 0.181E+01 0.175E+01 0.166E+01 0.155E+01 0.155E+01	PERCENT ERROR 2.12 2.13 2.11 2.06 1.99 1.88 1.73	N 1 2 3 4 5 6 7	TABLE E5 TR 0.6316 0.6466 0.6466 0.6617 0.6767 0.6718 0.7068 0.7068	CONTINUED PR 0.0213 0.0273 0.0346 0.0433 0.0536 0.0657 0.0799
LIQUIC ****** TEMP DEG F -40.00 -30.00 -20.00 -10.00 0.00 10.00 20.00 30.00	D DENSITY COM DENSITY COM DENS IN CALCD 0.899E+02 0.888E+02 0.878E+02 0.867E+02 0.855E+02 0.843E+02 0.830E+02 0.817E+02	PARISON ****** LB/FT3 PUBLD 0.880E+02 0.870E+02 0.860E+02 0.849E+02 0.827E+02 0.816E+02 0.804E+02	ABS ERROR 0.187E+01 0.185E+01 0.181E+01 0.181E+01 0.166E+01 0.155E+01 0.141E+01 0.125E+01	PERCENT ERROR 2.12 2.13 2.11 2.06 1.99 1.88 1.73 1.55	N 1 2 3 4 5 6 7 8	TABLE E5 TR 0.6316 0.6466 0.6466 0.6617 0.6767 0.6718 0.7068 0.7219 0.7369	CONTINUED PR 0.0213 0.0273 0.0346 0.0433 0.0536 0.0657 0.0799 0.0962
LIQUIC ****** TEMP DEG F -40.00 -30.00 -20.00 -10.00 10.00 10.00 20.00 30.00 40.00	D DENSITY COM DENSIN CALCD 0.899E+02 0.888E+02 0.878E+02 0.867E+02 0.855E+02 0.843E+02 0.830E+02 0.817E+02 0.817E+02 0.803E+02	PARISON ****** LB/FT3 PUBLD 0.880E+02 0.870E+02 0.860E+02 0.849E+02 0.838E+02 0.827E+02 0.816E+02 0.804E+02 0.804E+02 0.804E+02 0.804E+02	ABS ERROR 0.187E+01 0.185E+01 0.181E+01 0.175E+01 0.166E+01 0.125E+01 0.125E+01 0.105E+01	PERCENT ERROR 2.12 2.13 2.11 2.06 1.99 1.88 1.73 1.55 1.32	N 1 2 3 4 5 6 7 8 9	TABLE E5 TR 0.6316 0.6466 0.6466 0.6617 0.6767 0.6718 0.7068 0.7219 0.7369 0.7369	CONTINUED PR 0.0213 0.0273 0.0346 0.0433 0.0536 0.0457 0.0799 0.0792 0.0792 0.1150
LIQUIC ****** TEMP DEG F -40.00 -30.00 -20.00 -10.00 10.00 10.00 20.00 30.00 40.00 50.00	DENSITY COM DENSITY COM DENS IN CALCD 0.899E+02 0.888E+02 0.878E+02 0.867E+02 0.855E+02 0.843E+02 0.830E+02 0.817E+02 0.803E+02 0.788E+02	PARISON ****** LB/FT3 PUBLD 0.880E+02 0.870E+02 0.847E+02 0.847E+02 0.838E+02 0.816E+02 0.816E+02 0.816E+02 0.804E+02 0.773E+02 0.780E+02	ABS ERROR 0.187E+01 0.185E+01 0.181E+01 0.175E+01 0.166E+01 0.125E+01 0.125E+01 0.105E+01 0.816E+00	PERCENT ERROR 2.12 2.13 2.11 2.06 1.99 1.88 1.73 1.55 1.32 1.05	N 1 2 3 4 5 6 7 8 9 10	TABLE E5 TR 0.6316 0.6466 0.6466 0.6617 0.6767 0.6918 0.7068 0.7219 0.7369 0.7520 0.7520 0.7520	CONTINUED PR 0.0213 0.0273 0.0346 0.0433 0.0536 0.0457 0.0799 0.0792 0.1150 0.1364
LIQUIC ****** TEMP DEG F -40.00 -30.00 -20.00 -10.00 10.00 10.00 20.00 30.00 40.00 50.00 60.00	DENSITY COM DENSITY COM DENS IN CALCD 0.899E+02 0.888E+02 0.878E+02 0.867E+02 0.855E+02 0.843E+02 0.843E+02 0.830E+02 0.817E+02 0.803E+02 0.773E+02	PARISON ****** LB/FT3 PUBLD 0.880E+02 0.870E+02 0.860E+02 0.849E+02 0.838E+02 0.827E+02 0.816E+02 0.816E+02 0.804E+02 0.793E+02 0.768E+02	ABS ERROR 0.187E+01 0.185E+01 0.181E+01 0.166E+01 0.155E+01 0.141E+01 0.125E+01 0.125E+01 0.105E+01 0.816E+00 0.551E+00	PERCENT ERROR 2.12 2.13 2.11 2.06 1.99 1.88 1.73 1.55 1.32 1.05 0.72	N 1 2 3 4 5 6 7 8 9 10 11	TABLE E5 TR 0.6316 0.6466 0.6466 0.6617 0.6767 0.6718 0.7068 0.7219 0.7369 0.7520 0.7520 0.7670 0.7821	CONTINUED PR 0.0213 0.0273 0.0346 0.0433 0.0536 0.0457 0.0799 0.0792 0.1150 0.1364 0.1607
LIQUIC ****** TEMP DEG F -40.00 -30.00 -20.00 -10.00 0.00 10.00 20.00 30.00 40.00 50.00 40.00 70.00	DENSITY COM DENSITY COM DENS IN CALCD 0.899E+02 0.888E+02 0.878E+02 0.867E+02 0.855E+02 0.855E+02 0.843E+02 0.830E+02 0.817E+02 0.803E+02 0.738E+02 0.773E+02 0.757E+02	PARISON ****** LB/FT3 PUBLD 0.880E+02 0.870E+02 0.847E+02 0.847E+02 0.838E+02 0.816E+02 0.816E+02 0.816E+02 0.773E+02 0.773E+02 0.768E+02 0.755E+02	ABS ERROR 0.187E+01 0.185E+01 0.181E+01 0.175E+01 0.166E+01 0.155E+01 0.125E+01 0.125E+01 0.105E+01 0.816E+00 0.551E+00 0.250E+00	PERCENT ERROR 2.12 2.13 2.11 2.06 1.99 1.88 1.73 1.55 1.32 1.05 0.72 0.33	N 1 2 3 4 5 6 7 8 9 10 11 12	TABLE E5 TR 0.6316 0.6466 0.6466 0.6417 0.6767 0.6918 0.7068 0.7219 0.7369 0.7520 0.7520 0.7520 0.7670 0.7821 0.7821	CONTINUED PR 0.0213 0.0273 0.0273 0.0346 0.0433 0.0536 0.0457 0.0799 0.0962 0.1150 0.1364 0.1607 0.1882
LIQUIC ****** TEMP DEG F -40.00 -30.00 -20.00 -10.00 0.00 10.00 20.00 30.00 40.00 50.00 40.00 70.00 80.00	DENSITY COM DENSITY COM DENS IN CALCD 0.899E+02 0.888E+02 0.878E+02 0.867E+02 0.855E+02 0.843E+02 0.830E+02 0.837E+02 0.837E+02 0.773E+02 0.757E+02 0.757E+02 0.740E+02	PARISON ****** LB/FT3 PUBLD 0.880E+02 0.870E+02 0.847E+02 0.847E+02 0.838E+02 0.827E+02 0.816E+02 0.816E+02 0.816E+02 0.773E+02 0.775E+02 0.768E+02 0.755E+02 0.741E+02	ABS ERROR 0.187E+01 0.185E+01 0.181E+01 0.166E+01 0.155E+01 0.141E+01 0.125E+01 0.105E+01 0.105E+01 0.551E+00 0.551E+00 0.250E+00 -0.898E-01	PERCENT ERROR 2.12 2.13 2.11 2.06 1.99 1.88 1.73 1.55 1.32 1.05 0.72 0.33 -0.12	N 1 2 3 4 5 6 7 8 9 10 11 12 12 13	TABLE E5 TR 0.6316 0.6466 0.6466 0.6617 0.6767 0.6718 0.7068 0.7219 0.7369 0.7520 0.7520 0.7670 0.7671 0.7821 0.7971 0.8122	CONTINUED PR 0.0213 0.0273 0.0273 0.0346 0.0433 0.0536 0.0457 0.0799 0.0962 0.1150 0.1364 0.1607 0.1882 0.2190
LIQUIC ****** TEMP DEG F -40.00 -30.00 -20.00 -10.00 0.00 10.00 20.00 30.00 40.00 50.00 40.00 70.00 80.00 90.00	DENSITY COM DENSITY COM DENS IN CALCD 0.899E+02 0.888E+02 0.878E+02 0.867E+02 0.855E+02 0.855E+02 0.843E+02 0.830E+02 0.817E+02 0.817E+02 0.738E+02 0.773E+02 0.757E+02 0.740E+02 0.722E+02	PARISON ****** LB/FT3 PUBLD 0.880E+02 0.870E+02 0.847E+02 0.847E+02 0.838E+02 0.816E+02 0.816E+02 0.816E+02 0.773E+02 0.773E+02 0.768E+02 0.768E+02 0.768E+02 0.755E+02 0.741E+02 0.727E+02	ABS ERROR 0.187E+01 0.185E+01 0.181E+01 0.175E+01 0.146E+01 0.155E+01 0.125E+01 0.105E+01 0.105E+01 0.816E+00 0.551E+00 0.250E+00 -0.898E-01 -0.471E+00	PERCENT ERROR 2.12 2.13 2.11 2.06 1.99 1.88 1.73 1.55 1.32 1.05 0.72 0.33 -0.12 -0.65	N 1 2 3 4 5 6 7 8 9 10 11 12 12 13 14	TABLE E5 TR 0.6316 0.6466 0.6466 0.6617 0.6767 0.6717 0.7068 0.7219 0.7369 0.7520 0.7520 0.7520 0.7670 0.7821 0.7821 0.7971 0.8122 0.8272	CONTINUED PR 0.0213 0.0273 0.0273 0.0346 0.0433 0.0536 0.0457 0.0799 0.0792 0.1150 0.1364 0.1607 0.1882 0.2190 0.2534
LIQUIC ****** TEMP DEG F -40.00 -30.00 -20.00 -10.00 0.00 10.00 20.00 10.00 20.00 30.00 40.00 50.00 40.00 70.00 80.00 90.00	DENSITY COM DENSITY COM DENS IN CALCD 0.899E+02 0.888E+02 0.878E+02 0.867E+02 0.855E+02 0.843E+02 0.843E+02 0.830E+02 0.817E+02 0.817E+02 0.738E+02 0.757E+02 0.757E+02 0.722E+02 0.703E+02	PARISON ****** LB/FT3 PUBLD 0.880E+02 0.870E+02 0.860E+02 0.849E+02 0.849E+02 0.838E+02 0.816E+02 0.816E+02 0.816E+02 0.793E+02 0.793E+02 0.768E+02 0.768E+02 0.755E+02 0.768E+02 0.727E+02 0.712E+02	ABS ERROR 0.187E+01 0.185E+01 0.181E+01 0.175E+01 0.146E+01 0.155E+01 0.125E+01 0.125E+01 0.105E+01 0.816E+00 0.551E+00 0.250E+00 -0.898E-01 -0.471E+00 -0.895E+00	PERCENT ERROR 2.12 2.13 2.11 2.06 1.99 1.88 1.73 1.55 1.32 1.05 0.72 0.33 -0.12 -0.65 -1.26	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15	TABLE E5 TR 0.6316 0.6466 0.6466 0.6617 0.6767 0.6717 0.7068 0.7219 0.7369 0.7520 0.7520 0.7520 0.7670 0.7821 0.7821 0.7971 0.8122 0.8272 0.8423	CONTINUED PR 0.0213 0.0273 0.0273 0.0346 0.0433 0.0536 0.0457 0.0792 0.0792 0.1150 0.1364 0.1607 0.1882 0.2190 0.2534 0.2917
LIQUIC ****** TEMP DEG F -40.00 -30.00 -20.00 -10.00 0.00 10.00 20.00 30.00 40.00 50.00 40.00 70.00 80.00 90.00 10.00	DENSITY COM DENSITY COM DENS IN CALCD 0.899E+02 0.888E+02 0.878E+02 0.867E+02 0.855E+02 0.843E+02 0.843E+02 0.830E+02 0.837E+02 0.738E+02 0.773E+02 0.757E+02 0.757E+02 0.740E+02 0.703E+02 0.683E+02	PARISON ****** LB/FT3 PUBLD 0.880E+02 0.870E+02 0.870E+02 0.849E+02 0.849E+02 0.827E+02 0.816E+02 0.816E+02 0.816E+02 0.816E+02 0.793E+02 0.768E+02 0.768E+02 0.768E+02 0.755E+02 0.741E+02 0.712E+02 0.697E+02	ABS ERROR 0.187E+01 0.185E+01 0.181E+01 0.175E+01 0.146E+01 0.155E+01 0.141E+01 0.125E+01 0.105E+01 0.105E+01 0.816E+00 0.551E+00 0.250E+00 -0.878E-01 -0.471E+00 -0.875E+00 -0.137E+01	PERCENT ERROR 2.12 2.13 2.11 2.06 1.99 1.88 1.73 1.55 1.32 1.05 0.72 0.33 -0.12 -0.65 -1.26 -1.96	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16	TABLE E5 TR 0.6316 0.6466 0.6466 0.6617 0.6767 0.6767 0.7068 0.7219 0.7369 0.7520 0.7520 0.7520 0.7670 0.7821 0.7821 0.7871 0.7971 0.8122 0.8122 0.8272 0.8423 0.8573	CONTINUED PR 0.0213 0.0273 0.0273 0.0346 0.0433 0.0536 0.0457 0.0799 0.0962 0.1150 0.1364 0.1607 0.1882 0.2190 0.2534 0.2917 0.3342
LIQUIC ****** TEMP DEG F -40.00 -30.00 -20.00 -10.00 10.00 10.00 20.00 30.00 40.00 50.00 40.00 50.00 40.00 70.00 80.00 90.00 10.00	DENSITY COM DENSITY COM DENS IN CALCD 0.899E+02 0.888E+02 0.878E+02 0.867E+02 0.855E+02 0.843E+02 0.843E+02 0.830E+02 0.817E+02 0.817E+02 0.738E+02 0.773E+02 0.757E+02 0.757E+02 0.703E+02 0.703E+02 0.683E+02 0.663E+02	PARISON ****** LB/FT3 PUBLD 0.880E+02 0.870E+02 0.870E+02 0.849E+02 0.849E+02 0.827E+02 0.816E+02 0.816E+02 0.793E+02 0.793E+02 0.768E+02 0.768E+02 0.755E+02 0.741E+02 0.712E+02 0.697E+02 0.681E+02	ABS ERROR 0.187E+01 0.185E+01 0.185E+01 0.181E+01 0.155E+01 0.141E+01 0.125E+01 0.125E+01 0.105E+01 0.551E+00 0.551E+00 0.551E+00 0.250E+00 -0.898E-01 -0.471E+00 -0.137E+01 -0.189E+01	PERCENT ERROR 2.12 2.13 2.11 2.06 1.99 1.88 1.73 1.55 1.32 1.05 0.72 0.33 -0.12 -0.65 -1.26 -1.96 -2.77	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17	TABLE E5 TR 0.6316 0.6466 0.6466 0.6417 0.6767 0.6717 0.7068 0.7219 0.7369 0.7520 0.7520 0.7520 0.7670 0.7521 0.7521 0.7521 0.7671 0.7771 0.8122 0.8272 0.8423 0.8573 0.8724	CONTINUED PR 0.0213 0.0273 0.0273 0.0346 0.0433 0.0536 0.0457 0.0792 0.0792 0.1150 0.1364 0.1607 0.1882 0.2190 0.2534 0.2917 0.3342 0.3811
LIQUIC ****** TEMP DEG F -40.00 -30.00 -20.00 -10.00 10.00 20.00 10.00 20.00 40.00 50.00 40.00 70.00 80.00 90.00 10.00 10.00 10.00	DENSITY COM DENSITY COM DENS IN CALCD 0.899E+02 0.888E+02 0.878E+02 0.867E+02 0.855E+02 0.843E+02 0.843E+02 0.843E+02 0.817E+02 0.817E+02 0.73E+02 0.757E+02 0.757E+02 0.722E+02 0.703E+02 0.683E+02 0.663E+02 0.639E+02 0.639E+02	PARISON ****** LB/FT3 PUBLD 0.880E+02 0.870E+02 0.870E+02 0.849E+02 0.849E+02 0.827E+02 0.816E+02 0.816E+02 0.793E+02 0.793E+02 0.768E+02 0.768E+02 0.755E+02 0.741E+02 0.727E+02 0.712E+02 0.697E+02 0.663E+02	ABS ERROR 0.187E+01 0.185E+01 0.185E+01 0.181E+01 0.155E+01 0.141E+01 0.125E+01 0.125E+01 0.105E+01 0.105E+01 0.551E+00 0.551E+00 0.551E+00 0.250E+00 -0.878E-01 -0.471E+00 -0.137E+01 -0.189E+01 -0.246E+01	PERCENT ERROR 2.12 2.13 2.11 2.06 1.99 1.88 1.73 1.55 1.32 1.05 0.72 0.33 -0.12 -0.65 -1.26 -1.96 -2.77 -3.71	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18	TABLE E5 TR 0.6316 0.6466 0.6466 0.6417 0.6767 0.6717 0.7068 0.7219 0.7369 0.7520 0.7520 0.7520 0.7670 0.7521 0.82722 0.8272 0.8573 0.8573	CONTINUED PR 0.0213 0.0273 0.0273 0.0346 0.0433 0.0536 0.0657 0.0792 0.0792 0.1150 0.1364 0.1607 0.1882 0.2190 0.2534 0.2917 0.3342 0.3811 0.4327
LIQUIC ****** TEMP DEG F -40.00 -30.00 -20.00 -10.00 0.00 10.00 20.00 30.00 40.00 50.00 40.00 50.00 40.00 70.00 80.00 90.00 10.00 10.00 10.00	DENSITY COM DENSITY COM DENS IN CALCD 0.899E+02 0.888E+02 0.878E+02 0.867E+02 0.855E+02 0.843E+02 0.843E+02 0.830E+02 0.817E+02 0.738E+02 0.738E+02 0.773E+02 0.757E+02 0.757E+02 0.703E+02 0.703E+02 0.663E+02 0.639E+02 0.614E+02 0.6414E+02	PARISON ****** LB/FT3 PUBLD 0.880E+02 0.870E+02 0.870E+02 0.849E+02 0.849E+02 0.827E+02 0.816E+02 0.816E+02 0.793E+02 0.768E+02 0.768E+02 0.768E+02 0.755E+02 0.741E+02 0.727E+02 0.712E+02 0.697E+02 0.663E+02 0.663E+02 0.663E+02	ABS ERROR 0.187E+01 0.185E+01 0.185E+01 0.181E+01 0.175E+01 0.146E+01 0.155E+01 0.141E+01 0.125E+01 0.105E+01 0.551E+00 0.551E+00 0.551E+00 0.250E+00 -0.878E-01 -0.471E+00 -0.137E+01 -0.189E+01 -0.246E+01 -0.309E+01	PERCENT ERROR 2.12 2.13 2.11 2.06 1.99 1.88 1.73 1.55 1.32 1.05 0.72 0.33 -0.12 -0.65 -1.26 -1.96 -2.77 -3.71 -4.79	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19	TABLE E5 TR 0.6316 0.6466 0.6466 0.6617 0.6767 0.6717 0.7068 0.7219 0.7369 0.7520 0.7520 0.7520 0.7670 0.7521 0.7522 0.8573 0.8573 0.8724	CONTINUED PR 0.0213 0.0273 0.0273 0.0346 0.0433 0.0536 0.0457 0.0792 0.0792 0.1150 0.1364 0.1607 0.1882 0.2190 0.2534 0.2917 0.3342 0.3811 0.4327 0.4894
LIQUIC ****** TEMP DEG F -40.00 -30.00 -20.00 -10.00 0.00 10.00 20.00 30.00 40.00 50.00 40.00 50.00 40.00 50.00 10.000 10.000 10.00000000	DENSITY COM DENSITY COM DENS IN CALCD 0.899E+02 0.888E+02 0.878E+02 0.867E+02 0.855E+02 0.843E+02 0.843E+02 0.830E+02 0.817E+02 0.738E+02 0.738E+02 0.773E+02 0.773E+02 0.773E+02 0.773E+02 0.773E+02 0.703E+02 0.663E+02 0.663E+02 0.614E+02 0.586E+02 0.586E+02	PARISON ****** LB/FT3 PUBLD 0.880E+02 0.870E+02 0.870E+02 0.849E+02 0.849E+02 0.827E+02 0.816E+02 0.816E+02 0.793E+02 0.768E+02 0.768E+02 0.755E+02 0.768E+02 0.712E+02 0.712E+02 0.697E+02 0.641E+02 0.644E+02 0.624E+02 0	ABS ERROR 0.187E+01 0.185E+01 0.185E+01 0.181E+01 0.175E+01 0.146E+01 0.155E+01 0.141E+01 0.155E+01 0.105E+01 0.105E+01 0.551E+00 0.551E+00 0.551E+00 0.551E+00 0.250E+00 -0.875E+00 -0.137E+01 -0.246E+01 -0.378E+01 -0.378E+01	PERCENT ERROR 2.12 2.13 2.11 2.06 1.99 1.88 1.73 1.55 1.32 1.05 0.72 0.33 -0.12 -0.65 -1.26 -1.96 -2.77 -3.71 -4.79 -6.05	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20	TABLE E5 TR 0.6316 0.6466 0.6466 0.6617 0.6767 0.6767 0.7068 0.7219 0.7369 0.7520 0.7520 0.7520 0.7670 0.7821 0.7521 0.7871 0.7821 0.7971 0.8122 0.8272 0.8423 0.8573 0.8724 0.8874 0.9025 0.9175	CONTINUED PR 0.0213 0.0273 0.0273 0.0346 0.0433 0.0536 0.0457 0.0792 0.0792 0.1150 0.1364 0.1607 0.1882 0.2190 0.2534 0.2917 0.3342 0.3811 0.4327 0.4894 0.5513
LIQUIC ****** TEMP DEG F -40.00 -30.00 -20.00 -10.00 0.00 10.00 20.00 30.00 40.00 50.00 40.00 70.00 80.00 90.00 10.00 10.00 10.00 20.00 10.00 20.00 10.00 20.00 10.00	DENSITY COM DENSITY COM DENS IN CALCD 0.899E+02 0.888E+02 0.878E+02 0.867E+02 0.855E+02 0.843E+02 0.843E+02 0.843E+02 0.843E+02 0.843E+02 0.738E+02 0.773E+02 0.773E+02 0.773E+02 0.773E+02 0.773E+02 0.703E+02 0.703E+02 0.6639E+02 0.6639E+02 0.614E+02 0.556E+02 0.556E+02 0.556E+02	PARISON ****** LB/FT3 PUBLD 0.880E+02 0.870E+02 0.870E+02 0.849E+02 0.849E+02 0.827E+02 0.816E+02 0.816E+02 0.793E+02 0.768E+02 0.768E+02 0.768E+02 0.768E+02 0.7712E+02 0.712E+02 0.697E+02 0.643E+02 0.643E+02 0.644E+02 0.624E+02 0.601E+02	ABS ERROR 0.187E+01 0.185E+01 0.185E+01 0.185E+01 0.166E+01 0.155E+01 0.141E+01 0.125E+01 0.125E+01 0.105E+01 0.551E+00 0.551E+00 0.551E+00 0.551E+00 0.250E+00 -0.875E+00 -0.137E+01 -0.246E+01 -0.378E+01 -0.453E+01 -0.453E+01	PERCENT ERROR 2.12 2.13 2.11 2.06 1.99 1.88 1.73 1.55 1.32 1.05 0.72 0.33 -0.12 -0.65 -1.26 -1.96 -2.77 -3.71 -4.79 -6.05 -7.53	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21	TABLE E5 TR 0.6316 0.6466 0.6466 0.6417 0.6767 0.6767 0.7068 0.7219 0.7369 0.7520 0.7520 0.7670 0.7821 0.7520 0.7671 0.7821 0.7971 0.8122 0.8272 0.8423 0.8573 0.8724 0.8573 0.8724 0.8874 0.9025 0.9175 0.9326	CONTINUED PR 0.0213 0.0273 0.0273 0.0346 0.0433 0.0536 0.0657 0.0792 0.1150 0.1364 0.1607 0.1882 0.2190 0.2534 0.2917 0.3342 0.3811 0.4327 0.4894 0.5513 0.6190
LIQUIC ****** TEMP DEG F -40.00 -30.00 -20.00 -10.00 0.00 10.00 20.00 30.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 10.00 10.00 10.00 20.00 10.00 20.00 10.00 20.00 10.00 20.00 10.00 20.000 20.000 20.000 20.000 20.0000000	DENSITY COM DENSITY COM DENS IN CALCD 0.899E+02 0.888E+02 0.878E+02 0.867E+02 0.855E+02 0.843E+02 0.843E+02 0.830E+02 0.837E+02 0.837E+02 0.738E+02 0.738E+02 0.773E+02 0.757E+02 0.740E+02 0.703E+02 0.643E+02 0.642E+02 0.642E+02 0.556E+02 0.556E+02 0.522E+02 0.683E+02 0.522E+02 0.683E+02 0.522E+02 0.683E+02 0.522E+02 0.683E+02 0.522E+02 0.683E+02 0.522E+02 0.683E+02	PARISON ****** LB/FT3 PUBLD 0.880E+02 0.870E+02 0.870E+02 0.849E+02 0.849E+02 0.827E+02 0.816E+02 0.816E+02 0.793E+02 0.768E+02 0.768E+02 0.768E+02 0.755E+02 0.741E+02 0.727E+02 0.741E+02 0.727E+02 0.697E+02 0.643E+02 0.663E+02 0.663E+02 0.624E+02 0.624E+02 0.624E+02 0.624E+02 0.624E+02 0.6576E+02 0.6576E+02 0.6576E+02 0.6576E+02 0.6576E+02 0.6576E+02 0.6576E+02 0.6576E+02 0.6576E+02 0.6576E+02 0.6576E+02 0.6576E+02 0.6576E+02 0.6576E+02 0.6576E+02 0.5576E+02	ABS ERROR 0.187E+01 0.185E+01 0.185E+01 0.181E+01 0.175E+01 0.146E+01 0.155E+01 0.141E+01 0.155E+01 0.155E+01 0.105E+01 0.551E+00 0.551E+00 0.551E+00 0.551E+00 0.551E+00 -0.875E+01 -0.471E+01 -0.378E+01 -0.378E+01 -0.53E+01 -0.53E+01 -0.53E+01 -0.53E+01 -0.53E+01	PERCENT ERROR 2.12 2.13 2.11 2.06 1.99 1.88 1.73 1.55 1.32 1.05 0.72 0.33 -0.12 -0.65 -1.26 -1.96 -2.77 -3.71 -4.79 -6.05 -7.53 -9.27	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 21 22	TABLE E5 TR 0.6316 0.6466 0.6466 0.6417 0.6767 0.6918 0.7068 0.7219 0.7369 0.7520 0.7520 0.7670 0.7821 0.7520 0.7671 0.7821 0.7971 0.8122 0.8272 0.8423 0.8573 0.8724 0.8573 0.8724 0.8874 0.9025 0.9175 0.9326 0.9476	CONTINUED PR 0.0213 0.0273 0.0273 0.0346 0.0433 0.0536 0.0657 0.0792 0.1150 0.1364 0.1607 0.1882 0.2190 0.2534 0.2917 0.3342 0.3811 0.4327 0.4894 0.5513 0.6190 0.6926
LIQUIC ****** TEMP DEG F -40.00 -30.00 -20.00 -10.00 0.00 10.00 20.00 30.00 40.00 50.00 40.00 50.00 40.00 50.00 10.00 20.00 10.00 20.00 10.00 20.00 10.00 20.00 10.00 20.000 20.000 20.000 20.000 20.0000000	DENSITY COM DENSITY COM DENS IN CALCD 0.899E+02 0.888E+02 0.878E+02 0.878E+02 0.855E+02 0.843E+02 0.843E+02 0.843E+02 0.830E+02 0.817E+02 0.738E+02 0.738E+02 0.757E+02 0.773E+02 0.773E+02 0.773E+02 0.703E+02 0.643E+02 0.642E+02 0.642E+02 0.556E+02 0.556E+02 0.522E+02 0.483E+02 0.485E+02 0.485E+02	PARISON ****** LB/FT3 PUBLD 0.880E+02 0.870E+02 0.870E+02 0.849E+02 0.849E+02 0.827E+02 0.827E+02 0.816E+02 0.793E+02 0.793E+02 0.768E+02 0.755E+02 0.741E+02 0.727E+02 0.741E+02 0.727E+02 0.741E+02 0.727E+02 0.643E+02 0.663E+02 0.655E+0200E+020E+020E+020E+020E+020E+	ABS ERROR 0.187E+01 0.185E+01 0.185E+01 0.185E+01 0.166E+01 0.155E+01 0.141E+01 0.155E+01 0.155E+01 0.105E+01 0.105E+01 0.551E+00 0.551E+00 0.551E+00 0.551E+00 0.250E+00 -0.895E+00 -0.137E+01 -0.246E+01 -0.309E+01 -0.378E+01 -0.453E+01 -0.453E+01 -0.453E+01 -0.534E+01 -0.620E+01	PERCENT ERROR 2.12 2.13 2.11 2.06 1.99 1.88 1.73 1.55 1.32 1.05 0.72 0.33 -0.12 -0.65 -1.26 -1.96 -2.77 -3.71 -4.79 -6.05 -7.53 -9.27 -11.37	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23	TABLE E5 TR 0.6316 0.6466 0.6466 0.6417 0.6767 0.6918 0.7068 0.7219 0.7369 0.7520 0.7520 0.7670 0.7520 0.7670 0.76721 0.76721 0.77520 0.76721 0.7771 0.8122 0.8272 0.8272 0.8423 0.8573 0.8724 0.8573 0.8724 0.8272 0.8272 0.8423 0.8573 0.8724 0.9025 0.9175 0.9326 0.9476 0.9627	CONTINUED PR 0.0213 0.0273 0.0273 0.0346 0.0433 0.0536 0.0457 0.0799 0.0792 0.1150 0.1364 0.1607 0.1882 0.2190 0.2534 0.2917 0.3342 0.3811 0.4327 0.4894 0.5513 0.6190 0.6926 0.7726
LIQUIC ****** TEMP DEG F -40.00 -30.00 -20.00 -10.00 0.00 10.00 20.00 30.00 40.00 50.00 40.00 50.00 40.00 70.00 80.00 10.000 10.000 10.00000000	DENSITY COM DENSITY COM DENS IN CALCD 0.899E+02 0.888E+02 0.878E+02 0.878E+02 0.855E+02 0.855E+02 0.843E+02 0.843E+02 0.817E+02 0.830E+02 0.738E+02 0.738E+02 0.73E+02 0.757E+02 0.757E+02 0.740E+02 0.703E+02 0.643E+02 0.643E+02 0.645E+02 0.556E+02 0.522E+02 0.436E+02 0.436E+02 0.436E+02 0.465E+02	PARISON ****** LB/FT3 PUBLD 0.880E+02 0.870E+02 0.870E+02 0.849E+02 0.849E+02 0.827E+02 0.827E+02 0.827E+02 0.816E+02 0.793E+02 0.793E+02 0.768E+02 0.755E+02 0.741E+02 0.727E+02 0.741E+02 0.727E+02 0.643E+02 0.645E+02 0.645E+02 0.645E+02 0.645E+02 0.545E+02 0.576E+02 0.507E+02 0.507E+02	ABS ERROR 0.187E+01 0.185E+01 0.185E+01 0.181E+01 0.175E+01 0.146E+01 0.155E+01 0.141E+01 0.125E+01 0.105E+01 0.105E+01 0.551E+00 0.551E+00 0.551E+00 0.551E+00 0.250E+00 -0.898E-01 -0.471E+00 -0.137E+01 -0.246E+01 -0.309E+01 -0.534E+01 -0.534E+01 -0.534E+01 -0.534E+01 -0.5708E+01 -0.708E+01	PERCENT ERROR 2.12 2.13 2.11 2.06 1.99 1.88 1.73 1.55 1.32 1.05 0.72 0.33 -0.12 -0.65 -1.26 -1.96 -2.77 -3.71 -4.79 -6.05 -7.53 -9.27 -11.37 -13.97	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24	TABLE E5 TR 0.6316 0.6466 0.6466 0.6417 0.6767 0.6918 0.7068 0.7219 0.7369 0.7520 0.7670 0.7520 0.7670 0.7520 0.7671 0.77520 0.76721 0.76721 0.76721 0.76721 0.76721 0.76721 0.76721 0.76721 0.76721 0.76721 0.76721 0.81222 0.8272 0.8272 0.8272 0.8272 0.9255 0.9175 0.9326 0.9476 0.9627 0.9777	CONTINUED PR 0.0213 0.0273 0.0273 0.0346 0.0433 0.0536 0.0457 0.0799 0.0962 0.1150 0.1364 0.1607 0.1882 0.2190 0.2534 0.2917 0.3342 0.3811 0.4327 0.4894 0.5513 0.6190 0.6926 0.7726 0.8592
LIQUIC ****** TEMP DEG F -40.00 -30.00 -20.00 -10.00 0.00 10.00 20.00 40.00 50.00 40.00 50.00 40.00 50.00 10.00 20.00 10.00 20.000 20.000 20.000 20.000 20.0000000	DENSITY COM DENSITY COM DENS IN CAL CD 0.899E+02 0.888E+02 0.878E+02 0.878E+02 0.855E+02 0.843E+02 0.843E+02 0.843E+02 0.817E+02 0.817E+02 0.738E+02 0.757E+02 0.757E+02 0.740E+02 0.757E+02 0.745E+02 0.745E+02 0.639E+02 0.643E+02 0.643E+02 0.556E+02 0.556E+02 0.556E+02 0.556E+02 0.556E+02 0.436E+02 0.369E+02	PARISON ****** LB/FT3 PUBLD 0.880E+02 0.870E+02 0.870E+02 0.849E+02 0.849E+02 0.827E+02 0.827E+02 0.816E+02 0.793E+02 0.793E+02 0.768E+02 0.768E+02 0.768E+02 0.727E+02 0.712E+02 0.643E+02 0.643E+02 0.643E+02 0.643E+02 0.643E+02 0.644E+02 0.644E+02 0.576E+02 0.507E+02 0.507E+02 0.446E+02	ABS ERROR 0.187E+01 0.185E+01 0.185E+01 0.181E+01 0.175E+01 0.146E+01 0.155E+01 0.141E+01 0.125E+01 0.105E+01 0.816E+00 0.551E+00 0.250E+00 -0.878E+01 -0.471E+00 -0.875E+00 -0.137E+01 -0.246E+01 -0.378E+01 -0.453E+01 -0.534E+01 -0.708E+01 -0.766E+01	PERCENT ERROR 2.12 2.13 2.11 2.06 1.99 1.88 1.73 1.55 1.32 1.05 0.72 0.33 -0.12 -0.65 -1.26 -1.96 -2.77 -3.71 -4.79 -6.05 -7.53 -9.27 -11.37 -13.97 -17.19	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24 25	TABLE E5 TR 0.6316 0.6466 0.6466 0.6467 0.6767 0.6918 0.7068 0.7219 0.7369 0.7520 0.7670 0.7821 0.7971 0.8122 0.8272 0.8423 0.8573 0.8724 0.8573 0.8724 0.8573 0.8724 0.8874 0.9025 0.9175 0.9326 0.9476 0.9476 0.9777 0.9728	CONTINUED PR 0.0213 0.0273 0.0273 0.0346 0.0433 0.0536 0.0657 0.0799 0.0962 0.1150 0.1364 0.1407 0.1882 0.2190 0.2534 0.2917 0.3342 0.3811 0.4327 0.4894 0.5513 0.6190 0.6926 0.7726 0.8592 0.9529

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D1= 3.270701 6.0678329E-02

****** LIQUID ENTROPY COMPARISON

TABLE E5 CONTINUED

TEMP DEG F	BTU/ Calcd	'LB F PUBLD	ABS Error	PERCENT Error	N	TR	PR
-40.00	0.000E+00	0.000E+00	0.000E+00	0.00	1	0.6316	0.0213
-30.00	0.637E-02	0.598E-02	0.389E-03	6.50	2	0.6466	0.0273
-20.00	0.127E-01 0.189E-01	0.119E-01	0.781E-03	6.57	3	0.6617	0.0346
0.00	0.251E-01	0.1782-01	0.1545-02	6.51	5	U:0/0/ N 4918	0.0433
10.00	0.313E-01	0.293E-01	0.1948-02	6.62	6	0.7068	0.0657
20.00	0.374E-01	0.350E-01	0.235E-02	6.70	7	0.7219	0.0799
30.00	0.435E-01	0.407E-01	0.277E-02	6.80	8	0.7369	0.0962
40.00	0.495E-01	0.463E-01	0.322E-02	6.95	9	0.7520	0.1150
50.00	0.556E-01	0.519E-01	0.370E-02	7.13	10	0.7670	0.1364
60.00	0.617E-01	0.575E-01	0.421E-02	7.34	11	0.7821	0.1607
20.00	0.6//E-01 0.7795-04	U.63UE-01	0.478E-02	7.59	12	0.7971	0.1882
90.00	0.7382-01	0.0055-01	0.5366-02	7.00 8.16	13	0.8272	0.2190
100.00	0.862E-01	0.794E-01	0.674E-02	8.49	15	0.8423	0.2917
110.00	0.924E-01	0.849E-D1	0.752E-02	8.86	16	0.8573	0.3342
120.00	0.988E-01	0.904E-01	0.837E-02	9.26	17	0.8724	0.3811
130.00	0.105E+00	0.960E-01	0.931E-02	9.70	18	0.8874	0.4327
140.00	0.112E+00	0.102E+00	0.103E-01	10.15	19	0.9025	0.4894
150.00	0.119E+00	0.107E+00	0.115E-01	10.66	20	0.9175	0.5513
160.00	0.126E+00	0.113E+00	0.127E-01	11.21	21	0.9326	0.6190
120.00	0.1342+00	0.120E+00	0.141E-01	11.79	22	0.9476	0.6926
100.00	0.1426+00	0.1262+00	U.157E-01	12.41	23	0.9627	0.7726
200.00	0.164E+00	0.145E+00	0.1752-01	13.04	24	0.9777	0.0572
AAD= SD1= SD2=	8.412169 8.9903465E-0 9.0413079E-0)3 32			it.		
						TADLE DE CON	T.T.1111CD
*****	**********	######################################	÷			TABLE ED COM	IINDED
TEMP DEG F	BTU/ CALCD	PARISON ******** LB F PUBLD	ABS Error	PERCENT Error	N	TABLE E 5 CON	PR
TEMP DEG F	BTU/ CALCD	LB F PUBLD 0.239E+00	ABS Error -0.511E-03	PERCENT Error -0.21	N 1	TABLE E 5 CON TR 0.6316	PR 0.0213
TEMP DEG F -40.00 -30.00	BTU/ CALCD 0.238E+00 0.236E+00	LB F PUBLD 0.239E+00 0.236E+00	ABS ERROR -0.511E-03 0.235E-03	PERCENT ERROR -0.21 0.10	N 1 2	TABLE E 5 CON TR 0.6316 0.6466	PR 0.0213 0.0273
TEMP DEG F 	BTU/ CALCD 0.238E+00 0.236E+00 0.234E+00	PARISON ******* B F PUBLD 0.239E+00 0.236E+00 0.233E+00	ABS ERROR -0.511E-03 0.235E-03 0.973E-03	PERCENT ERROR -0.21 0.10 0.42	N 1 2 3	TABLE E CON TR 0.6316 0.6466 0.6617	PR 0.0213 0.0273 0.0346
TEMP DEG F -40.00 -30.00 -20.00 -10.00	BTU/ CALCD 0.238E+00 0.236E+00 0.234E+00 0.234E+00 0.232E+00	PARISON ******* DLB F PUBLD 0.239E+00 0.236E+00 0.231E+00 0.231E+00	ABS ERROR -0.511E-03 0.235E-03 0.973E-03 0.170E-02	PERCENT ERROR 0.21 0.10 0.42 0.74	N 1 2 3 4	TABLE E CON TR 0.6316 0.6466 0.6617 0.6767 0.6767	PR 0.0213 0.0273 0.0346 0.0433 0.0534
TEMP DEG F -40.00 -30.00 -20.00 -10.00 0.00	BTU/ CALCD 0.238E+00 0.236E+00 0.236E+00 0.234E+00 0.232E+00 0.231E+00 0.231E+00	PARISON ******* DEBF PUBLD 0.239E+00 0.234E+00 0.231E+00 0.231E+00 0.228E+00	ABS ERROR -0.511E-03 0.235E-03 0.973E-03 0.170E-02 0.242E-02 0.342E-02	PERCENT ERROR 0.21 0.10 0.42 0.74 1.06 1 38	N 1 2 3 4 5 4	TABLE CON TR 0.6316 0.6466 0.6617 0.6767 0.6918 0.7068	PR 0.0213 0.0273 0.0346 0.0433 0.0536 0.0556
TEMP DE6 F 	BTU/ CALCD 0.238E+00 0.236E+00 0.236E+00 0.234E+00 0.232E+00 0.231E+00 0.229E+00 0.229E+00	PARISON ******* DEB F PUBLD 0.239E+00 0.233E+00 0.231E+00 0.228E+00 0.226E+00 0.226E+00	ABS ERROR -0.511E-03 0.235E-03 0.773E-03 0.170E-02 0.242E-02 0.312E-02 0.381E-02	PERCENT ERROR -0.21 0.10 0.42 0.74 1.06 1.38 1.70	N 1 2 3 4 5 6 7	TABLE CON TR 0.6316 0.6466 0.6617 0.6767 0.6718 0.7068 0.7068 0.7219	PR 0.0213 0.0273 0.0346 0.0433 0.0536 0.0557 0.0599
TEMP DEG F 	BTU/ CALCD 0.238E+00 0.236E+00 0.236E+00 0.232E+00 0.231E+00 0.229E+00 0.229E+00 0.228E+00 0.228E+00 0.226E+00	PARISON ******* DEBF PUBLD 0.239E+00 0.233E+00 0.231E+00 0.228E+00 0.226E+00 0.224E+00 0.224E+00 0.222E+00	ABS ERROR -0.511E-03 0.235E-03 0.773E-03 0.170E-02 0.242E-02 0.312E-02 0.381E-02 0.447E-02	PERCENT ERROR -0.21 0.10 0.42 0.74 1.06 1.38 1.70 2.02	N 1 2 3 4 5 6 7 8	TABLE CON TR 0.6316 0.6466 0.6617 0.6767 0.6718 0.7068 0.7219 0.7369	PR 0.0213 0.0273 0.0346 0.0433 0.0536 0.0536 0.0657 0.0799 0.0962
TEMP DEG F 	BTU/ CALCD 0.238E+00 0.236E+00 0.234E+00 0.232E+00 0.231E+00 0.231E+00 0.229E+00 0.228E+00 0.226E+00 0.225E+00	PARISON ******* LB F PUBLD 0.239E+00 0.234E+00 0.231E+00 0.224E+00 0.224E+00 0.224E+00 0.222E+00 0.220E+00	ABS ERROR -0.511E-03 0.235E-03 0.973E-03 0.170E-02 0.242E-02 0.312E-02 0.312E-02 0.381E-02 0.447E-02 0.513E-02	PERCENT ERROR -0.21 0.10 0.42 0.74 1.06 1.38 1.70 2.02 2.33	N 1 2 3 4 5 6 7 8 9	TABLE CON TR 0.6316 0.6466 0.6466 0.6417 0.6767 0.6718 0.7068 0.7068 0.7219 0.7369 0.7520	PR 0.0213 0.0273 0.0346 0.0433 0.0536 0.0457 0.0799 0.0962 0.1150
TEMP DEG F 	BTU/ CALCD 0.238E+00 0.236E+00 0.236E+00 0.232E+00 0.231E+00 0.229E+00 0.228E+00 0.228E+00 0.226E+00 0.225E+00 0.224E+00	PARISON ******* LB F PUBLD 0.239E+00 0.234E+00 0.231E+00 0.224E+00 0.224E+00 0.222E+00 0.220E+00 0.220E+00 0.218E+00	ABS ERROR -0.511E-03 0.235E-03 0.773E-03 0.170E-02 0.242E-02 0.312E-02 0.381E-02 0.447E-02 0.513E-02 0.576E-02	PERCENT ERROR -0.21 0.10 0.42 0.74 1.06 1.38 1.70 2.02 2.33 2.64	N 1 2 3 4 5 6 7 8 9 10	TR 0.6316 0.6466 0.6466 0.6467 0.6767 0.6718 0.7068 0.7219 0.7369 0.7520 0.7670	PR 0.0213 0.0273 0.0273 0.0346 0.0433 0.0536 0.0657 0.0799 0.0962 0.1150 0.1364
TEMP DEG F 	BTU/ CALCD 0.238E+00 0.236E+00 0.236E+00 0.234E+00 0.232E+00 0.231E+00 0.229E+00 0.228E+00 0.226E+00 0.225E+00 0.225E+00 0.224E+00 0.223E+00	PARISON ******* LB F PUBLD 0.237E+00 0.234E+00 0.231E+00 0.228E+00 0.226E+00 0.224E+00 0.222E+00 0.220E+00 0.220E+00 0.216E+00	ABS ERROR -0.511E-03 0.235E-03 0.773E-03 0.170E-02 0.242E-02 0.312E-02 0.381E-02 0.447E-02 0.513E-02 0.576E-02 0.636E-02	PERCENT ERROR -0.21 0.10 0.42 0.74 1.06 1.38 1.70 2.02 2.33 2.64 2.94	N 1 2 3 4 5 6 7 8 9 10 11	TR 0.6316 0.6466 0.6466 0.6477 0.6767 0.6918 0.7068 0.7219 0.7369 0.7520 0.7670 0.7821	PR 0.0213 0.0273 0.0346 0.0433 0.0536 0.0657 0.0799 0.0962 0.1150 0.1364 0.1607
TEMP DEG F 	BTU/ CALCD 0.238E+00 0.236E+00 0.236E+00 0.234E+00 0.232E+00 0.231E+00 0.229E+00 0.228E+00 0.226E+00 0.226E+00 0.225E+00 0.224E+00 0.223E+00 0.221E+00	PARISON ******* LB F PUBLD 0.237E+00 0.234E+00 0.231E+00 0.228E+00 0.226E+00 0.222E+00 0.222E+00 0.220E+00 0.218E+00 0.216E+00 0.215E+00	ABS ERROR -0.511E-03 0.235E-03 0.773E-03 0.170E-02 0.242E-02 0.312E-02 0.381E-02 0.447E-02 0.513E-02 0.576E-02 0.636E-02 0.693E-02	PERCENT ERROR -0.21 0.10 0.42 0.74 1.06 1.38 1.70 2.02 2.33 2.64 2.94 3.23	N 1 2 3 4 5 6 7 8 9 10 11 12	TR TR 0.6316 0.6466 0.6466 0.6477 0.6767 0.6718 0.7068 0.7219 0.7369 0.7520 0.7670 0.7821 0.7971	PR 0.0213 0.0273 0.0273 0.0346 0.0433 0.0536 0.0657 0.0799 0.0962 0.1150 0.1364 0.1607 0.1882
TEMP DEG F 	BTU/ CALCD 0.238E+00 0.236E+00 0.236E+00 0.234E+00 0.232E+00 0.231E+00 0.229E+00 0.228E+00 0.226E+00 0.225E+00 0.224E+00 0.223E+00 0.221E+00 0.220E+00	PARISON ******* LB F PUBLD 0.239E+00 0.236E+00 0.231E+00 0.223E+00 0.226E+00 0.224E+00 0.222E+00 0.220E+00 0.220E+00 0.218E+00 0.215E+00 0.213E+00	ABS ERROR -0.511E-03 0.235E-03 0.773E-03 0.170E-02 0.242E-02 0.312E-02 0.381E-02 0.447E-02 0.513E-02 0.576E-02 0.636E-02 0.693E-02 0.749E-02	PERCENT ERROR -0.21 0.10 0.42 0.74 1.06 1.38 1.70 2.02 2.33 2.64 2.94 3.23 3.52	N 1 2 3 4 5 6 7 8 9 10 11 12 13	TR 0.6316 0.6466 0.6466 0.6477 0.6767 0.6718 0.7068 0.7219 0.7369 0.7520 0.7670 0.7821 0.7971 0.8122	PR 0.0213 0.0273 0.0346 0.0433 0.0536 0.0657 0.0799 0.0962 0.1150 0.1364 0.1607 0.1882 0.2190
TEMP DEG F 	BTU/ CALCD 0.238E+00 0.236E+00 0.236E+00 0.234E+00 0.232E+00 0.231E+00 0.229E+00 0.229E+00 0.228E+00 0.226E+00 0.225E+00 0.225E+00 0.223E+00 0.221E+00 0.221E+00 0.219E+00	PARISON ******* LB F PUBLD 0.237E+00 0.236E+00 0.231E+00 0.223E+00 0.224E+00 0.224E+00 0.222E+00 0.220E+00 0.218E+00 0.215E+00 0.213E+00 0.211E+00	ABS ERROR -0.511E-03 0.235E-03 0.773E-03 0.170E-02 0.242E-02 0.312E-02 0.381E-02 0.447E-02 0.513E-02 0.576E-02 0.636E-02 0.693E-02 0.749E-02 0.801E-02	PERCENT ERROR -0.21 0.10 0.42 0.74 1.06 1.38 1.70 2.02 2.33 2.64 2.94 3.23 3.52 3.79	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14	TR 0.6316 0.6466 0.6466 0.6467 0.6767 0.6718 0.7068 0.7219 0.7369 0.7520 0.7670 0.7821 0.7971 0.8122 0.8272 0.8272	PR 0.0213 0.0273 0.0346 0.0433 0.0536 0.0657 0.0799 0.0962 0.1150 0.1364 0.1607 0.1882 0.2190 0.2534
XAPOUR ****** TEMP DEG F -40.00 -30.00 -20.00 -10.00 20.00 10.00 20.00 30.00 40.00 50.00 60.00 70.00 80.00 90.00 00.00	BTU/ CALCD 0.238E+00 0.236E+00 0.236E+00 0.234E+00 0.232E+00 0.231E+00 0.229E+00 0.228E+00 0.226E+00 0.225E+00 0.225E+00 0.221E+00 0.221E+00 0.219E+00 0.219E+00 0.218E+00 0.218E+00	PARISON ******* LB F PUBLD 0.237E+00 0.236E+00 0.231E+00 0.223E+00 0.226E+00 0.226E+00 0.222E+00 0.220E+00 0.218E+00 0.215E+00 0.213E+00 0.211E+00 0.210E+00 0.210E+00	ABS ERROR -0.511E-03 0.235E-03 0.773E-03 0.170E-02 0.242E-02 0.312E-02 0.381E-02 0.447E-02 0.513E-02 0.576E-02 0.576E-02 0.693E-02 0.801E-02 0.801E-02 0.850E-02	PERCENT ERROR -0.21 0.10 0.42 0.74 1.06 1.38 1.70 2.02 2.33 2.64 2.94 3.23 3.52 3.79 4.06 6 74	N 1 2 3 4 5 6 7 8 9 10 11 11 12 13 14 15	TABLE CON TR 0.6316 0.6466 0.6466 0.6467 0.6767 0.6918 0.7068 0.7219 0.7369 0.7520 0.7670 0.7821 0.7971 0.8122 0.8272 0.8423 0.8423	PR 0.0213 0.0273 0.0346 0.0433 0.0536 0.0657 0.0799 0.0962 0.1150 0.1364 0.1607 0.1882 0.2190 0.2534 0.2917 0.7742
XAPOUR ****** TEMP DEG F -40.00 -30.00 -20.00 10.00 20.00 10.00 20.00 30.00 40.00 50.00 60.00 70.00 80.00 90.00 10.00 20.00	BTU/ CALCD 0.238E+00 0.236E+00 0.236E+00 0.234E+00 0.232E+00 0.231E+00 0.229E+00 0.228E+00 0.228E+00 0.226E+00 0.224E+00 0.223E+00 0.221E+00 0.221E+00 0.219E+00 0.217E+00 0.217E+00 0.217E+00	PARISON ******* LB F PUBLD 0.237E+00 0.236E+00 0.231E+00 0.223E+00 0.224E+00 0.224E+00 0.222E+00 0.220E+00 0.218E+00 0.215E+00 0.213E+00 0.211E+00 0.210E+00 0.208E+00 0.206E+00	ABS ERROR -0.511E-03 0.235E-03 0.773E-03 0.170E-02 0.242E-02 0.312E-02 0.381E-02 0.447E-02 0.513E-02 0.576E-02 0.693E-02 0.801E-02 0.896E-02 0.876E-02	PERCENT ERROR -0.21 0.10 0.42 0.74 1.06 1.38 1.70 2.02 2.33 2.64 2.94 3.23 3.52 3.79 4.06 4.31 6.54	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17	TABLE CON TR 0.6316 0.6466 0.6466 0.6467 0.6767 0.6918 0.7068 0.7219 0.7369 0.7520 0.7670 0.7520 0.7670 0.7821 0.7971 0.8122 0.8272 0.8423 0.8573 0.8726	PR 0.0213 0.0273 0.0346 0.0433 0.0536 0.0657 0.0799 0.0962 0.1150 0.1364 0.1607 0.1882 0.2190 0.2534 0.2917 0.3342 0.344
XAPOOR ****** TEMP DEG F -40.00 -30.00 -20.00 -10.00 20.00 10.00 20.00 30.00 40.00 50.00 60.00 70.00 80.00 90.00 10.00 20.00 30.00	BTU/ CALCD 0.238E+00 0.236E+00 0.236E+00 0.234E+00 0.232E+00 0.231E+00 0.229E+00 0.229E+00 0.226E+00 0.224E+00 0.224E+00 0.221E+00 0.221E+00 0.219E+00 0.216E+00 0.216E+00 0.216E+00	PARISON ******* LB F PUBLD 0.237E+00 0.236E+00 0.233E+00 0.231E+00 0.228E+00 0.226E+00 0.226E+00 0.220E+00 0.218E+00 0.215E+00 0.213E+00 0.211E+00 0.210E+00 0.206E+00 0.206E+00	ABS ERROR -0.511E-03 0.235E-03 0.973E-03 0.170E-02 0.242E-02 0.312E-02 0.381E-02 0.447E-02 0.513E-02 0.576E-02 0.693E-02 0.801E-02 0.896E-02 0.939E-02 0.976E-02	PERCENT ERROR -0.21 0.10 0.42 0.74 1.06 1.38 1.70 2.02 2.33 2.64 2.94 3.23 3.52 3.79 4.06 4.31 4.56 4.79	N 1 2 3 4 5 6 7 8 9 10 11 11 12 13 14 15 16 17 18	TABLE CON TR 0.6316 0.6466 0.6466 0.6467 0.6767 0.6918 0.7068 0.7219 0.7369 0.7520 0.7670 0.7821 0.7971 0.8122 0.8272 0.8423 0.8573 0.8724 0.8724 0.8874	PR 0.0213 0.0273 0.0346 0.0433 0.0536 0.0657 0.0799 0.0962 0.1150 0.1364 0.1607 0.1882 0.2190 0.2534 0.2917 0.3342 0.3811 0.4327
XAPOUR ****** TEMP DEG F -40.00 -30.00 -20.00 10.00 20.00 10.00 20.00 30.00 40.00 50.00 60.00 70.00 80.00 90.00 10.00 20.00 30.00 40.00	BTU/ CALCD 0.238E+00 0.236E+00 0.236E+00 0.234E+00 0.232E+00 0.231E+00 0.229E+00 0.229E+00 0.226E+00 0.225E+00 0.225E+00 0.224E+00 0.221E+00 0.219E+00 0.217E+00 0.216E+00 0.214E+00 0.212E+00	PARISON ******* LB F PUBLD 0.237E+00 0.236E+00 0.233E+00 0.231E+00 0.224E+00 0.224E+00 0.224E+00 0.224E+00 0.218E+00 0.215E+00 0.213E+00 0.213E+00 0.211E+00 0.210E+00 0.208E+00 0.204E+00 0.202E+00	ABS ERROR -0.511E-03 0.235E-03 0.973E-03 0.170E-02 0.242E-02 0.312E-02 0.381E-02 0.447E-02 0.513E-02 0.513E-02 0.576E-02 0.693E-02 0.801E-02 0.896E-02 0.896E-02 0.939E-02 0.978E-02 0.978E-02 0.101E-01	PERCENT ERROR -0.21 0.10 0.42 0.74 1.06 1.38 1.70 2.02 2.33 2.64 2.94 3.23 3.52 3.79 4.06 4.31 4.56 4.79 5.01	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19	TABLE CON TR 0.6316 0.6466 0.6466 0.6617 0.6767 0.6718 0.7068 0.7219 0.7369 0.7520 0.7670 0.7821 0.7971 0.8122 0.8272 0.8423 0.8573 0.8573 0.8724 0.8874 0.9025	PR 0.0213 0.0273 0.0273 0.0346 0.0433 0.0536 0.0457 0.0799 0.0762 0.1150 0.1364 0.1607 0.1364 0.1607 0.1364 0.2534 0.2917 0.3342 0.3811 0.4327 0.4894
XAPOUR ****** TEMP DE6 F -40.00 -30.00 -20.00 10.00 20.00 10.00 20.00 30.00 40.00 50.00 80.00 90.00 10.00 20.00 30.00 40.00 50.00 10.00 50.00	BTU/ CALCD 0.238E+00 0.236E+00 0.236E+00 0.234E+00 0.232E+00 0.231E+00 0.229E+00 0.229E+00 0.226E+00 0.225E+00 0.225E+00 0.225E+00 0.221E+00 0.219E+00 0.217E+00 0.217E+00 0.212E+00 0.212E+00 0.212E+00 0.212E+00 0.212E+00 0.212E+00 0.212E+00 0.211E+00	PARISON ******* LB F PUBLD 0.237E+00 0.236E+00 0.233E+00 0.231E+00 0.224E+00 0.224E+00 0.224E+00 0.224E+00 0.214E+00 0.215E+00 0.213E+00 0.213E+00 0.214E+00 0.216E+00 0.206E+00 0.202E+00 0.202E+00 0.202E+00 0.202E+00 0.202E+00 0.202E+00 0.202E+00	ABS ERROR -0.511E-03 0.235E-03 0.973E-03 0.170E-02 0.242E-02 0.312E-02 0.381E-02 0.447E-02 0.513E-02 0.576E-02 0.636E-02 0.693E-02 0.801E-02 0.896E-02 0.896E-02 0.939E-02 0.978E-02 0.978E-02 0.101E-01 0.105E-01	PERCENT ERROR -0.21 0.10 0.42 0.74 1.06 1.38 1.70 2.02 2.33 2.64 2.94 3.23 3.52 3.79 4.06 4.31 4.56 4.31 4.56 4.79 5.01 5.23	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20	TABLE CON TR 0.6316 0.6466 0.6466 0.6617 0.6767 0.6718 0.7068 0.7219 0.7369 0.7520 0.7670 0.7821 0.7971 0.8122 0.8272 0.8423 0.8573 0.8573 0.8724 0.8874 0.9025 0.9175	PR 0.0213 0.0273 0.0273 0.0346 0.0433 0.0536 0.0433 0.0536 0.0457 0.0799 0.0762 0.1150 0.1364 0.1607 0.1882 0.2190 0.2534 0.2917 0.3342 0.3811 0.4327 0.4894 0.5513
XAPOUR ****** TEMP DE6 F -40.00 -30.00 -20.00 10.00 20.00 10.00 20.00 30.00 40.00 50.00 40.00 50.00 80.00 90.00 10.00 20.00 30.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 50.00 40.00 50.00 50.00 40.00 50.00	BTU/ CALCD 0.238E+00 0.236E+00 0.236E+00 0.234E+00 0.232E+00 0.231E+00 0.229E+00 0.229E+00 0.226E+00 0.225E+00 0.225E+00 0.225E+00 0.221E+00 0.219E+00 0.218E+00 0.218E+00 0.216E+00 0.214E+00 0.214E+00 0.212E+00 0.212E+00 0.212E+00 0.212E+00 0.212E+00 0.212E+00 0.212E+00 0.212E+00 0.212E+00 0.212E+00 0.212E+00 0.212E+00 0.212E+00	PARISON ******* UB F PUBLD 0.239E+00 0.236E+00 0.231E+00 0.223E+00 0.224E+00 0.224E+00 0.224E+00 0.214E+00 0.215E+00 0.213E+00 0.215E+00 0.215E+00 0.214E+00 0.208E+00 0.208E+00 0.202E+00 0.216E+00 0.206E+00 00	ABS ERROR -0.511E-03 0.235E-03 0.973E-03 0.170E-02 0.242E-02 0.312E-02 0.381E-02 0.447E-02 0.513E-02 0.576E-02 0.576E-02 0.693E-02 0.801E-02 0.896E-02 0.896E-02 0.939E-02 0.939E-02 0.978E-02 0.978E-02 0.101E-01 0.105E-01 0.108E-01	PERCENT ERROR -0.21 0.10 0.42 0.74 1.06 1.38 1.70 2.02 2.33 2.64 2.94 3.23 3.52 3.79 4.06 4.31 4.56 4.31 4.56 4.79 5.01 5.23 5.44	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21	TABLE CON TR 0.6316 0.6466 0.6466 0.6617 0.6767 0.6718 0.7068 0.7219 0.7369 0.7520 0.7670 0.7821 0.7971 0.8122 0.8272 0.8423 0.8272 0.8423 0.8573 0.8724 0.8573 0.8724 0.8874 0.9025 0.9175 0.9326	PR 0.0213 0.0273 0.0273 0.0346 0.0433 0.0536 0.0433 0.0536 0.0433 0.0536 0.0433 0.0546 0.0799 0.0799 0.07962 0.1150 0.1364 0.1407 0.1882 0.2190 0.2534 0.2917 0.3342 0.3811 0.4327 0.4894 0.5513 0.6190
XAPOUR ****** TEMP DE6 F -40.00 -30.00 -20.00 10.00 20.00 10.00 20.00 30.00 40.00 50.00 50.00 40.00 50.00	BTU/ CALCD 0.238E+00 0.236E+00 0.236E+00 0.234E+00 0.232E+00 0.231E+00 0.229E+00 0.228E+00 0.226E+00 0.225E+00 0.225E+00 0.225E+00 0.225E+00 0.221E+00 0.219E+00 0.218E+00 0.218E+00 0.218E+00 0.218E+00 0.218E+00 0.214E+00 0.226E+00 0.2276E+00 0.2276E+00 0.2176E+00 0.2176E+00 0.216E+00 0.206E+00 0.206E+00 0.206E+00 0.206E+00 0.206E+00 0.206E+00 0.206E+00 0.206E+00 0.206E+00 0.206E+00	PARISON ******* UB F PUBLD 0.239E+00 0.236E+00 0.231E+00 0.223E+00 0.224E+00 0.224E+00 0.224E+00 0.214E+00 0.215E+00 0.215E+00 0.215E+00 0.215E+00 0.216E+00 0.208E+00 0.208E+00 0.208E+00 0.202E+00 0.215E+00 0.205E+00 00	ABS ERROR -0.511E-03 0.235E-03 0.973E-03 0.170E-02 0.242E-02 0.312E-02 0.381E-02 0.447E-02 0.513E-02 0.576E-02 0.576E-02 0.693E-02 0.749E-02 0.896E-02 0.896E-02 0.939	PERCENT ERROR -0.21 0.10 0.42 0.74 1.06 1.38 1.70 2.02 2.33 2.64 2.94 3.23 3.52 3.79 4.06 4.31 4.56 4.31 4.56 4.56 5.23 5.44 5.66	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22	TABLE CON TR 0.6316 0.6466 0.6466 0.6477 0.6767 0.6718 0.7068 0.7219 0.7369 0.7520 0.7670 0.7821 0.7971 0.8122 0.8272 0.8423 0.8272 0.8423 0.8573 0.8724 0.8573 0.8724 0.8874 0.9025 0.9175 0.9326 0.9476	PR 0.0213 0.0273 0.0273 0.0346 0.0433 0.0536 0.0433 0.0536 0.0657 0.0799 0.0962 0.1150 0.1364 0.1407 0.1882 0.2190 0.2534 0.2917 0.3342 0.3811 0.4327 0.4894 0.5513 0.6190 0.6926
XAPOUR ****** TEMP DE6 F -40.00 -30.00 -20.00 10.00 20.00 10.00 20.00 30.00 40.00 50.00 50.00 40.00 50.00 50.00 40.00 50.00	BTU/ CALCD 0.238E+00 0.236E+00 0.236E+00 0.234E+00 0.232E+00 0.231E+00 0.229E+00 0.229E+00 0.225E+00 0.225E+00 0.225E+00 0.225E+00 0.225E+00 0.225E+00 0.225E+00 0.225E+00 0.219E+00 0.218E+00 0.218E+00 0.218E+00 0.218E+00 0.214E+00 0.225E+00 0.217E+00 0.217E+00 0.216E+00 0.206E+00 0.206E+00 0.206E+00	PARISON ******* LB F PUBLD 0.239E+00 0.236E+00 0.231E+00 0.223E+00 0.224E+00 0.224E+00 0.224E+00 0.214E+00 0.215E+00 0.215E+00 0.215E+00 0.214E+00 0.216E+00 0.208E+00 0.208E+00 0.208E+00 0.202E+00 0.215E+00 00	ABS ERROR -0.511E-03 0.235E-03 0.773E-03 0.170E-02 0.242E-02 0.312E-02 0.381E-02 0.447E-02 0.513E-02 0.576E-02 0.673E-02 0.673E-02 0.749E-02 0.896E-02 0.896E-02 0.939E-02 0.939E-02 0.939E-02 0.939E-02 0.939E-02 0.939E-02 0.939E-02 0.939E-02 0.939E-02 0.939E-02 0.939E-02 0.939E-02 0.939E-02 0.101E-01 0.108E-01 0.113E-01	PERCENT ERROR -0.21 0.10 0.42 0.74 1.06 1.38 1.70 2.02 2.33 2.64 2.94 3.23 3.52 3.79 4.06 4.31 4.56 4.31 4.56 4.79 5.01 5.23 5.44 5.66 5.91	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23	TABLE CON TR 0.6316 0.6466 0.6466 0.6477 0.6767 0.6718 0.7068 0.7219 0.7369 0.7520 0.7670 0.7821 0.7971 0.8122 0.8272 0.8423 0.8272 0.8423 0.8573 0.8724 0.8573 0.8724 0.8874 0.9025 0.9175 0.9326 0.9476 0.9627	PR 0.0213 0.0273 0.0273 0.0346 0.0433 0.0536 0.0657 0.0799 0.0962 0.1150 0.1364 0.1407 0.1882 0.2190 0.2534 0.2917 0.3342 0.3811 0.4327 0.4894 0.5513 0.6190 0.6926 0.7726
XAPOUR ****** TEMP DEG F -40.00 -30.00 -20.00 10.00 20.00 10.00 20.00 30.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 50.00 40.00 50.00 50.00 40.00 50.00 50.00 40.00 50.00	BTU/ CALCD 0.238E+00 0.236E+00 0.236E+00 0.234E+00 0.232E+00 0.232E+00 0.227E+00 0.228E+00 0.228E+00 0.225E+00 0.225E+00 0.225E+00 0.2219E+00 0.217E+00 0.217E+00 0.217E+00 0.217E+00 0.214E+00 0.214E+00 0.214E+00 0.214E+00 0.214E+00 0.214E+00 0.215E+00 0.225E+00 0.217E+00 0.215E+000000000000000000000000000	PARISON ******* 	ABS ERROR -0.511E-03 0.235E-03 0.773E-03 0.170E-02 0.242E-02 0.312E-02 0.381E-02 0.447E-02 0.513E-02 0.576E-02 0.673E-02 0.673E-02 0.876E-02 0.876E-02 0.876E-02 0.876E-02 0.978E-02 0.978E-02 0.978E-02 0.978E-02 0.101E-01 0.108E-01 0.113E-01 0.117E-01	PERCENT ERROR -0.21 0.10 0.42 0.74 1.06 1.38 1.70 2.02 2.33 2.64 2.94 3.23 3.52 3.79 4.06 4.31 4.56 4.31 4.56 4.79 5.01 5.23 5.44 5.66 5.91 6.25	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24	TABLE CON TR 0.6316 0.6466 0.6466 0.6477 0.6767 0.6718 0.7068 0.7219 0.7369 0.7520 0.7670 0.7821 0.7971 0.8122 0.8272 0.8423 0.8272 0.8423 0.8573 0.8724 0.8874 0.9025 0.9175 0.9326 0.9476 0.9627 0.9777	PR 0.0213 0.0273 0.0273 0.0346 0.0433 0.0536 0.0536 0.0536 0.057 0.0799 0.0962 0.1150 0.1364 0.1407 0.1382 0.2190 0.2534 0.2917 0.3342 0.3811 0.4327 0.4894 0.5513 0.6190 0.6926 0.7726 0.8592
XAPOUR ****** TEMP DEG F -40.00 -30.00 -20.00 10.00 20.00 10.00 20.00 30.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 40.00 50.00 50.00 40.00 50.00	BTU/ CALCD 0.238E+00 0.236E+00 0.236E+00 0.234E+00 0.232E+00 0.232E+00 0.227E+00 0.228E+00 0.228E+00 0.226E+00 0.225E+00 0.221E+00 0.217E+00 0.217E+00 0.217E+00 0.217E+00 0.217E+00 0.214E+0000000000000000000000000000	PARISON ******* 	ABS ERROR -0.511E-03 0.235E-03 0.773E-03 0.170E-02 0.242E-02 0.312E-02 0.381E-02 0.447E-02 0.513E-02 0.576E-02 0.673E-02 0.673E-02 0.876E-02 0.876E-02 0.876E-02 0.876E-02 0.978E-02 0.978E-02 0.978E-02 0.978E-02 0.101E-01 0.108E-01 0.113E-01 0.117E-01 0.124E-01	PERCENT ERROR -0.21 0.10 0.42 0.74 1.06 1.38 1.70 2.02 2.33 2.64 2.94 3.23 3.52 3.79 4.06 4.31 4.56 4.31 4.56 4.79 5.01 5.23 5.44 5.66 5.91 6.25 6.99	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24 25	TABLE CON TR 0.6316 0.6466 0.6466 0.6477 0.6767 0.6718 0.7068 0.7219 0.7369 0.7520 0.7670 0.7821 0.7971 0.8122 0.8423 0.8272 0.8423 0.8573 0.8724 0.8573 0.8724 0.8874 0.9025 0.9175 0.9326 0.9476 0.9476 0.9777 0.9928	PR 0.0213 0.0273 0.0273 0.0346 0.0433 0.0536 0.0536 0.0536 0.0536 0.0597 0.0799 0.0962 0.1150 0.1364 0.1407 0.1882 0.2190 0.2534 0.2534 0.2917 0.3342 0.3811 0.4327 0.4894 0.5513 0.6190 0.6926 0.7726 0.8592 0.9529

6

3.371461 7.9773273E-03 4.0038630E-02 AD= ;D1=

;02=

TABLE E6

				PERCENT	 N	 TR	PR
DEGF	CALCD	PUBLD	ERROR	ERROR			
			0.2505-01	12 14		0.4785	0.0005
-40.00	0.2316+00	0.2005+00	0.2506-01	11.70	2	0.4899	0.0007
-30.00	0.3342400	0.2772+00	0.3502-01	10 44	3	0.5013	0.0009
-20.00	0.4/46+00	0.4276700	0.4472-01	9,19	· 4	0.5127	0.0013
-10.00	0.0012700	0.8795+00	0.5582-01	8.09	5	0.5241	0.0018
0.00	0.4005+00	0.44/5+04	0.8782-01	7.10	-	0.5355	0.0025
10.00	0.1222701	0.1146701	0.8112-01	6.13	7	0.5469	0.0033
20.00	0.24/5+04	0.1936+01	0.4075+00	5.27	å	0.5583	0.0043
30.00	0.2146+01	0.2032+01	0.1072.00	4.43	9	0.5697	0.0056
40.00 %	0.7555+01	0.2050701	0.1182+00	3.72	10	0.5811	0.0071
50.00	0.3552401	0.3432401	0.1335+00	3.05	11	0.5925	0.0090
20.00	0.4512701	0.4376401	0.1346+00	2.42	12	0.6039	0.0113
/0.00	0.3000+01	0.3326+01	0.1295+00	1.88	13	0.6153	0.0141
	0.7032+01	0.8702+01	0.1175+00	1.37	14	0.6267	0.0174
70.00	0.0002+01	0.05552401	0.1015+00	n.97	15	0.6381	0.0212
100.00	0.4285+02	0.4295+02	0.4305-01	0.49	16	0.6495	0.0257
110.00	0.1200102	0.1202+02	0.0302 01	0.16	× 17	0.6609	0.0309
120.00	0 1845+02	0.1342402	-0.2758-01	-0.15	18	0.6723	0.0369
140.00	0.1072702	D 219F+02	-0.711E-01	-0.32	19	0.6837	0.0438
150.00	0.2172102	0.2595+02	-0.155E+00	-0.60	20	0.6951	0.0517
140 00	0.1002+02	0.3045+02	-0.225E+00	-0.74	21	0.7065	0.0606
170.00	0.3526+02	0.355E+02	-0.307E+00	-0.86	22	0.7179	0.0706
180.00	0.408E+02	0.412E+02	-0.373E+00	-0.91	23	0.7293	0.0819
190.00	0.471E+02	0.476E+02	-0.465E+00	-0.98	24	0.7407	0.0945
200.00	0.541E+02	0.547E+02	-0.523E+00	-0.96	25	0.7521	0.1085
210.00	0.619E+02	0,625E+02	-0.596E+00	-0.95	26	0.7635	0.1241
220.00	0.705E+02	0.711E+02	-0.580E+00	-0.82	27	0.7749	0.1413
230.00	0.799E+02	0.813E+02	-0.132E+01	-1.63	28	0.7863	0.1602
240.00	0.903E+02	0.915E+02	-0.113E+01	-1.24	29	0.7977	0.1811
250.00	0.102E+03	0.102E+03	-0.918E-01	-0.09	30	0.8091	0.2037
260.00	0.114E+03	0.114E+03	0.566E+00	0.50	31	0.8205	0.2288
270.00	0.128E+03	0.127E+03	0.328E+DD	0.26	32	0.8319	0.2559
280.00	0.142E+03	0.142E+03	0.257E+00	0.18	33	0.8433	0.2853
290.00	0,158E+03	0.158E+03	0.416E+00	0.26	34	0.8547	0.3172
300.00	0.175E+03	0.176E+03	-0.131E+00	-0.07	35	0.8661	0.3517
310.00	0.194E+03	0.193E+03	0.905E+00	0.47	36	0.8776	0.3889
320.00	0.214E+03	0.214E+03	0.371E+00	0.17	37	0.8890	0.4290
330.00	0.236E+03	0.235E+03	0.339E+00	0.14	38	0.9004	0.4722
340.00	0.259E+03	0.258E+03	0.879E+00	0.34	39	0.9118	0.5184
350.00	0.283E+03	0.283E+03	D.681E-01	0.02	40	0.9232	0.5680
360.00	0.310E+03	0.310E+03	0.227E-01	0.01	41	0.9346	0.6210
370.00	0.338E+03	0.338E+03	0.819E-01	0.02	42	0.9460	0.6777
380.00	0.368E+03	0.368E+03	0.129E+00	0.04	43	0.9574	0.7381
390.00	0.400E+03	0.400E+03	0.348E+00	0.09	44	0.9688	0.8025
400.00	0.435E+03	0.434E+03	0.228E+00	0.05	45	0.9802	0.8710
410.00	0.471E+03	0.471E+03	0.261E+00	0.06	46	0.9916	0.9433

AAD= 2.206517 SD1= 0.4079235 SD2= 4.0156055E-02

R152a

TABLE E7

******* PRESSURE COMPARISON ***************

TEMD				DEDCENT	N	TP	PR
	PRESSURE	PSIA	A85 F0000	FERGENI	n		
DEG F	CALCD	PUBLD	ERROR	ERNON			
	0 7005.04	0 7//5+04	-0 3425+00	-4 86	1	0.6030	0.0109
-40.00	0.7082+01	0.7445701	-0.3622+00	-3.59	2	0.6174	0.0143
~30.00	0.4345+02	0.1245+02	-0.3472.00	-2.54	3	0.6317	0.0186
-20.00	0.1212702	0.1246402	-0.263E+00	-1.67	4	0.6461	0.0238
-10.00	0.1552+02	0.1985+02	-D.192E+00	-0.97	5	0.6605	0.0301
40.00	0.1702+02	0.1702+04	-0.104E+00	-0.42	6	0.6749	0.0377
10.00	0.2462402	0.2472.02	-0.1268-02	0.00	7	0.6892	0.0467
20.00	0.3042+02 N.374F+N2	0.372E+02	0.113E+00	0.30	8	0.7036	0.0573
40.00	0.4545+02	0.452E+02	0.230E+00	0.51	9	0.7180	0.0696
50.00	0.547E+02	0.544E+02	0.348E+00	0.64	10	0.7323	0.0840
60.00	0.4555+02	0.650E+02	0.456E+00	0.70	11	0.7467	0.1004
70.00	0.778E+02	0.772E+02	0.547E+00	0.71	12	0.7611	0.1193
80.00	0.917E+02	0.911E+02	0.613E+00	0.67	13	0.7754	0.1407
90.00	0.107E+03	0.107E+03	0.645E+00	0.60	14	0.7898	0.1649
100.00	0.125E+03	0.125E+03	0.638E+00	0.51	15	0.8042	0.1921
110.00	0.145E+03	0.145E+03	0.585E+00	0.40	16	0.8185	D.2226
120.00	0.167E+03	0.167E+03	0.483E+00	0.29	17	D.8329	0.2565
130.00	0,192E+03	0.192E+03	0.333E+00	0.17	18	0.8473	0.2942
140.00	0.219E+03	0.219E+03	0.136E+00	0.06	19	0.8616	0.3360
150.00	0.249E+03	0.249E+03	-0.968E-01	-0.04	20	0.8760	0.3820
160.00	0.282E+03	0.282E+03	-0.355E+00	-0.13	21	0.8904	0.4325
170.00	0.318E+03	0.319E+03	-D.616E+0D	-0.19	22	0.9047	D.4879
180.00	0.358E+03	0.358E+03	-0.854E+00	-0.24	23	0.9191	0.5484
190.00	0.401E+03	0.402E+03	-0.103E+01	-0.26	24	D.9335	0.6144
200.00	0.447E+03	0.448E+03	-0.110E+01	-0.25	25	0.9478	0.6862
210.00	0.498E+03	0.499E+03	-0.101E+01	-0.20	26	0.9622	0.7640
******* LIQUID ******	DENSITY COM	******* PARISON *******				TABLE E7	ONTINUED
ТЕМР	DENS IN	LB/FT3	 ABS	PERCENT	N	TR	PR
TEMP DEG F	DENS IN Calcd	LB/FT3 PUBLD	ABS Error	PERCENT Error	N	TR	PR
TEMP DEG F -40.00	DENS IN CALCD 0.608E+02	LB/FT3 PUBLD 0.649E+02	ABS Error -0.416E+01	PERCENT Error -6.40	N 1	TR 0.6030	PR 0.0109
TEMP DEG F -40.00 -30.00	DENS IN CALCD 0.608E+02 0.602E+02	LB/FT3 PUBLD 0.649E+02 0.643E+02	ABS ERROR -0.416E+01 -0.408E+01	PERCENT ERROR -6.40 -6.35	N 1 2	TR 0.6030 0.6174	PR 0.0109 0.0143
TEMP DEG F -40.00 -30.00 -20.00	DENS IN CALCD 0.608E+02 0.602E+02 0.596E+02	LB/FT3 PUBLD 0.649E+02 0.643E+02 0.636E+02	ABS ERROR -0.416E+01 -0.408E+01 -0.402E+01	PERCENT ERROR -6.40 -6.35 -6.33	N 1 2 3	TR 0.6030 0.6174 0.6317	PR 0.0109 0.0143 0.0186
TEMP DEG F -40.00 -30.00 -20.00 -10.00	DENS IN CALCD 0.608E+02 0.602E+02 0.596E+02 0.589E+02 0.589E+02	LB/FT3 PUBLD 0.649E+02 0.643E+02 0.636E+02 0.636E+02 0.629E+02	ABS ERROR -0.416E+01 -0.408E+01 -0.402E+01 -0.397E+01	PERCENT ERROR -6.40 -6.35 -6.33 -6.32	N 1 2 3 4	TR 0.6030 0.6174 0.6317 0.6461	PR 0.0109 0.0143 0.0186 0.0238
TEMP DEG F -40.00 -30.00 -20.00 -10.00 0.00	DENS IN CALCD 0.608E+02 0.602E+02 0.596E+02 0.589E+02 0.583E+02 0.583E+02	LB/FT3 PUBLD 0.649E+02 0.643E+02 0.636E+02 0.636E+02 0.629E+02 0.622E+02	ABS ERROR -0.416E+01 -0.408E+01 -0.402E+01 -0.397E+01 -0.393E+01	PERCENT ERROR -6.40 -6.35 -6.33 -6.32 -6.33	N 1 2 3 4 5	TR 0.6030 0.6174 0.6317 0.6461 0.6605	PR 0.0109 0.0143 0.0144 0.0238 0.0301
TEMP DEG F -40.00 -30.00 -20.00 -10.00 0.00 10.00	DENS IN CALCD 0.608E+02 0.602E+02 0.596E+02 0.589E+02 0.583E+02 0.583E+02 0.576E+02	LB/FT3 PUBLD 0.649E+02 0.643E+02 0.636E+02 0.636E+02 0.629E+02 0.622E+02 0.615E+02	ABS ERROR -0.416E+01 -0.408E+01 -0.402E+01 -0.397E+01 +0.393E+01 -0.391E+01	PERCENT ERROR -6.40 -6.35 -6.33 -6.32 -6.33 -6.36	N 1 2 3 4 5 6	TR 0.6030 0.6174 0.6317 0.6461 0.6605 0.6749	PR 0.0109 0.0143 0.0186 0.0238 0.0301 0.0377
TEMP DEG F -40.00 -30.00 -20.00 -10.00 0.00 10.00 20.00	DENS IN CALCD 0.608E+02 0.602E+02 0.596E+02 0.589E+02 0.583E+02 0.583E+02 0.568E+02	LB/FT3 PUBLD 0.649E+02 0.643E+02 0.636E+02 0.636E+02 0.629E+02 0.622E+02 0.615E+02 0.607E+02	ABS ERROR -0.416E+01 -0.408E+01 -0.402E+01 -0.377E+01 -0.373E+01 -0.371E+01 -0.389E+01	PERCENT ERROR -6.40 -6.35 -6.33 -6.32 -6.33 -6.36 -6.41	N 1 2 3 4 5 6 7	TR 0.6030 0.6174 0.6317 0.6461 0.6605 0.6749 0.6892	PR 0.0109 0.0143 0.0186 0.0238 0.0301 0.0377 0.0467
TEMP DEG F -40.00 -30.00 -20.00 -10.00 0.00 10.00 20.00 30.00	DENS IN CALCD 0.608E+02 0.602E+02 0.596E+02 0.589E+02 0.583E+02 0.583E+02 0.568E+02 0.568E+02 0.561E+02	LB/FT3 PUBLD 0.649E+02 0.643E+02 0.636E+02 0.629E+02 0.622E+02 0.622E+02 0.615E+02 0.607E+02 0.600E+02	ABS ERROR -0.416E+01 -0.408E+01 -0.402E+01 -0.377E+01 -0.373E+01 -0.371E+01 -0.389E+01 -0.389E+01	PERCENT ERROR -6.40 -6.35 -6.33 -6.32 -6.33 -6.36 -6.41 -6.48	N 1 2 3 4 5 6 7 8	TR 0.6030 0.6174 0.6317 0.6461 0.6605 0.6749 0.6892 0.7036	PR 0.0109 0.0143 0.0143 0.0238 0.0238 0.0301 0.0377 0.0467 0.0467
TEMP DEG F -40.00 -30.00 -20.00 -10.00 0.00 10.00 20.00 30.00 40.00	DENS IN CALCD 0.608E+02 0.602E+02 0.596E+02 0.589E+02 0.583E+02 0.568E+02 0.568E+02 0.564E+02 0.564E+02 0.553E+02	LB/FT3 PUBLD 0.649E+02 0.643E+02 0.636E+02 0.629E+02 0.622E+02 0.615E+02 0.607E+02 0.600E+02 0.592E+02	ABS ERROR -0.416E+01 -0.408E+01 -0.402E+01 -0.397E+01 -0.393E+01 -0.389E+01 -0.389E+01 -0.390E+01	PERCENT ERROR -6.40 -6.35 -6.33 -6.32 -6.33 -6.36 -6.41 -6.48 -6.58	N 1 2 3 4 5 6 7 8 9	TR 0.6030 0.6174 0.6317 0.6461 0.6605 0.6749 0.6892 0.7036 0.7180	PR 0.0109 0.0143 0.0186 0.0238 0.0301 0.0377 0.0467 0.0573 0.0696
TEMP DEG F -40.00 -30.00 -20.00 -10.00 0.00 10.00 20.00 30.00 40.00 50.00	DENS IN CALCD 0.608E+02 0.602E+02 0.596E+02 0.589E+02 0.583E+02 0.568E+02 0.568E+02 0.564E+02 0.553E+02 0.553E+02	LB/FT3 PUBLD 0.649E+02 0.643E+02 0.636E+02 0.629E+02 0.622E+02 0.615E+02 0.607E+02 0.607E+02 0.592E+02 0.584E+02	ABS ERROR -0.416E+01 -0.408E+01 -0.397E+01 -0.393E+01 -0.391E+01 -0.389E+01 -0.389E+01 -0.390E+01 -0.392E+01	PERCENT ERROR -6.40 -6.35 -6.33 -6.32 -6.33 -6.33 -6.36 -6.41 -6.48 -6.58 -6.71	N 1 2 3 4 5 6 7 8 9 10	TR 0.6030 0.6174 0.6317 0.6461 0.6605 0.6749 0.6892 0.7036 0.7180 0.7323	PR 0.0109 0.0143 0.0186 0.0238 0.0301 0.0377 0.0467 0.0573 0.0496 0.0840
TEMP DEG F -40.00 -30.00 -20.00 -10.00 10.00 10.00 20.00 30.00 40.00 50.00 60.00	DENS IN CALCD 0.608E+02 0.602E+02 0.596E+02 0.589E+02 0.583E+02 0.568E+02 0.568E+02 0.564E+02 0.554E+02 0.553E+02 0.554E+02	LB/FT3 PUBLD 0.649E+02 0.643E+02 0.636E+02 0.629E+02 0.622E+02 0.615E+02 0.607E+02 0.600E+02 0.592E+02 0.584E+02 0.576E+02	ABS ERROR -0.416E+01 -0.408E+01 -0.402E+01 -0.377E+01 -0.373E+01 -0.371E+01 -0.387E+01 -0.387E+01 -0.370E+01 -0.372E+01 -0.376E+01	PERCENT ERROR -6.40 -6.35 -6.33 -6.32 -6.33 -6.32 -6.33 -6.41 -6.48 -6.58 -6.71 -6.87	N 1 2 3 4 5 6 7 8 9 10 11	TR 0.6030 0.6174 0.6317 0.6461 0.6605 0.6749 0.6892 0.7036 0.7180 0.7323 0.7467	PR 0.0109 0.0143 0.0186 0.0238 0.0301 0.0377 0.0467 0.0573 0.0696 0.0840 0.1004
TEMP DEG F -40.00 -30.00 -20.00 -10.00 10.00 10.00 20.00 30.00 40.00 50.00 60.00	DENS IN CALCD 0.608E+02 0.602E+02 0.596E+02 0.587E+02 0.583E+02 0.568E+02 0.564E+02 0.564E+02 0.554E+02 0.554E+02 0.554E+02 0.554E+02 0.554E+02	LB/FT3 PUBLD 0.649E+02 0.643E+02 0.636E+02 0.629E+02 0.622E+02 0.615E+02 0.607E+02 0.600E+02 0.592E+02 0.584E+02 0.584E+02 0.567E+02	ABS ERROR -0.416E+01 -0.408E+01 -0.402E+01 -0.377E+01 -0.373E+01 -0.371E+01 -0.387E+01 -0.387E+01 -0.370E+01 -0.370E+01 -0.376E+01 -0.401E+01	PERCENT ERROR -6.40 -6.35 -6.33 -6.32 -6.33 -6.32 -6.33 -6.41 -6.48 -6.58 -6.71 -6.87 -7.07	N 1 2 3 4 5 6 7 8 9 10 11 11 12	TR 0.6030 0.6174 0.6317 0.6461 0.6605 0.6749 0.6892 0.7036 0.7180 0.7323 0.7467 0.7611	PR 0.0109 0.0143 0.0186 0.0238 0.0301 0.0377 0.0467 0.0573 0.0696 0.0840 0.1004 0.1193
TEMP DEG F -40.00 -30.00 -20.00 -10.00 10.00 10.00 20.00 30.00 40.00 50.00 60.00 70.00 80.00	DENS IN CALCD 0.608E+02 0.602E+02 0.596E+02 0.589E+02 0.583E+02 0.568E+02 0.564E+02 0.564E+02 0.554E+02 0.554E+02 0.554E+02 0.536E+02 0.518E+02 0.518E+02	LB/FT3 PUBLD 0.649E+02 0.643E+02 0.636E+02 0.629E+02 0.622E+02 0.615E+02 0.607E+02 0.600E+02 0.592E+02 0.584E+02 0.584E+02 0.567E+02 0.567E+02	ABS ERROR -0.416E+01 -0.408E+01 -0.402E+01 -0.397E+01 -0.397E+01 -0.389E+01 -0.389E+01 -0.389E+01 -0.390E+01 -0.392E+01 -0.396E+01 -0.401E+01 -0.408E+01	PERCENT ERROR -6.40 -6.35 -6.33 -6.32 -6.33 -6.32 -6.33 -6.41 -6.48 -6.58 -6.71 -6.87 -7.07 -7.30	N 1 2 3 4 5 6 7 8 9 10 11 11 12 13	TR 0.6030 0.6174 0.6317 0.6461 0.6605 0.6749 0.6892 0.7036 0.7180 0.7323 0.7467 0.7611 0.7754	PR 0.0109 0.0143 0.0186 0.0238 0.0301 0.0377 0.0467 0.0573 0.0467 0.0573 0.0696 0.0840 0.1004 0.1193 0.1407
TEMP DEG F -40.00 -30.00 -20.00 -10.00 10.00 10.00 20.00 30.00 40.00 50.00 40.00 70.00 80.00	DENS IN CALCD 0.608E+02 0.602E+02 0.596E+02 0.589E+02 0.583E+02 0.568E+02 0.564E+02 0.564E+02 0.553E+02 0.553E+02 0.553E+02 0.518E+02 0.518E+02 0.508E+02 0.508E+02	LB/FT3 PUBLD 0.649E+02 0.643E+02 0.636E+02 0.629E+02 0.622E+02 0.615E+02 0.607E+02 0.607E+02 0.592E+02 0.592E+02 0.584E+02 0.557E+02 0.559E+02 0.550E+02	ABS ERROR -0.416E+01 -0.408E+01 -0.402E+01 -0.377E+01 -0.373E+01 -0.379E+01 -0.387E+01 -0.387E+01 -0.387E+01 -0.370E+01 -0.376E+01 -0.401E+01 -0.416E+01 -0.426E+01	PERCENT ERROR -6.40 -6.35 -6.33 -6.32 -6.33 -6.32 -6.33 -6.41 -6.48 -6.58 -6.71 -6.87 -7.07 -7.30 -7.56 -7.88	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15	TR 0.6030 0.6174 0.6317 0.6461 0.6605 0.6749 0.6892 0.7036 0.7180 0.7323 0.7467 0.7611 0.7754 0.7898 0.8922	PR 0.0109 0.0143 0.0186 0.0238 0.0301 0.0377 0.0467 0.0573 0.0467 0.0573 0.0696 0.0840 0.1004 0.1193 0.1407 0.1649
TEMP DEG F -40.00 -30.00 -20.00 -10.00 10.00 10.00 20.00 30.00 40.00 50.00 40.00 70.00 80.00 90.00	DENS IN CALCD 0.608E+02 0.602E+02 0.596E+02 0.589E+02 0.589E+02 0.568E+02 0.564E+02 0.564E+02 0.564E+02 0.553E+02 0.553E+02 0.554E+02 0.518E+02 0.518E+02 0.518E+02 0.498E+02 0.497E+02	LB/FT3 PUBLD 0.649E+02 0.643E+02 0.636E+02 0.629E+02 0.622E+02 0.615E+02 0.607E+02 0.607E+02 0.592E+02 0.592E+02 0.584E+02 0.559E+02 0.559E+02 0.540E+02 0.540E+02	ABS ERROR -0.416E+01 -0.408E+01 -0.402E+01 -0.377E+01 -0.373E+01 -0.379E+01 -0.387E+01 -0.387E+01 -0.370E+01 -0.370E+01 -0.401E+01 -0.408E+01 -0.416E+01 -0.426E+01	PERCENT ERROR -6.40 -6.35 -6.33 -6.32 -6.33 -6.32 -6.33 -6.41 -6.48 -6.58 -6.71 -6.87 -7.07 -7.30 -7.56 -7.88 -8.24	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 14	TR 0.6030 0.6174 0.6317 0.6461 0.6605 0.6749 0.6892 0.7036 0.7180 0.7323 0.7467 0.7611 0.7754 0.7898 0.8042 0.8042	PR 0.0109 0.0143 0.0186 0.0238 0.0301 0.0377 0.0467 0.0573 0.0696 0.0840 0.1004 0.1193 0.1407 0.1649 0.1921
TEMP DEG F -40.00 -30.00 -20.00 -10.00 10.00 20.00 30.00 40.00 50.00 40.00 70.00 80.00 90.00 100.00	DENS IN CALCD 0.608E+02 0.602E+02 0.596E+02 0.589E+02 0.583E+02 0.568E+02 0.564E+02 0.564E+02 0.564E+02 0.553E+02 0.553E+02 0.536E+02 0.518E+02 0.518E+02 0.518E+02 0.498E+02 0.487E+02 0.487E+02	LB/FT3 PUBLD 0.649E+02 0.643E+02 0.636E+02 0.629E+02 0.622E+02 0.615E+02 0.607E+02 0.607E+02 0.592E+02 0.592E+02 0.576E+02 0.559E+02 0.559E+02 0.540E+02 0.530E+02 0.530E+02	ABS ERROR -0.416E+01 -0.408E+01 -0.402E+01 -0.397E+01 -0.397E+01 -0.397E+01 -0.389E+01 -0.389E+01 -0.390E+01 -0.392E+01 -0.396E+01 -0.401E+01 -0.408E+01 -0.416E+01 -0.426E+01 -0.437E+01	PERCENT ERROR -6.40 -6.35 -6.33 -6.32 -6.33 -6.32 -6.33 -6.41 -6.48 -6.58 -6.71 -6.87 -7.07 -7.30 -7.56 -7.88 -8.24 -8.45	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17	TR 0.6030 0.6174 0.6317 0.6461 0.6605 0.6749 0.6892 0.7036 0.7180 0.7323 0.7467 0.7611 0.7754 0.7898 0.8042 0.8185 0.8328	PR 0.0109 0.0143 0.0186 0.0238 0.0301 0.0377 0.0467 0.0573 0.0696 0.0840 0.1004 0.1193 0.1407 0.1649 0.1921 0.2226 0.2545
TEMP DEG F -40.00 -30.00 -20.00 -10.00 10.00 20.00 30.00 40.00 50.00 40.00 70.00 80.00 70.00 100.00 110.00	DENS IN CALCD 0.608E+02 0.602E+02 0.596E+02 0.589E+02 0.583E+02 0.568E+02 0.564E+02 0.564E+02 0.564E+02 0.553E+02 0.553E+02 0.536E+02 0.518E+02 0.518E+02 0.518E+02 0.498E+02 0.498E+02 0.475E+02 0.475E+02	LB/FT3 PUBLD 0.649E+02 0.643E+02 0.636E+02 0.629E+02 0.622E+02 0.615E+02 0.607E+02 0.607E+02 0.592E+02 0.592E+02 0.576E+02 0.559E+02 0.559E+02 0.540E+02 0.520E+02 0.520E+02	ABS ERROR -0.416E+01 -0.408E+01 -0.402E+01 -0.397E+01 -0.397E+01 -0.397E+01 -0.389E+01 -0.389E+01 -0.390E+01 -0.392E+01 -0.401E+01 -0.401E+01 -0.416E+01 -0.426E+01 -0.455E+01	PERCENT ERROR -6.40 -6.35 -6.33 -6.32 -6.33 -6.32 -6.33 -6.41 -6.48 -6.58 -6.71 -6.87 -7.07 -7.56 -7.88 -8.24 -8.65 -9.13	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18	TR 0.6030 0.6174 0.6317 0.6461 0.6605 0.6749 0.6892 0.7036 0.7180 0.7323 0.7467 0.7611 0.7754 0.7898 0.8042 0.8185 0.8329 0.8477	PR 0.0109 0.0143 0.0186 0.0238 0.0301 0.0377 0.0467 0.0573 0.0467 0.0573 0.0696 0.0840 0.1004 0.1094 0.1193 0.1407 0.1449 0.1921 0.2226 0.2565 0.2565
TEMP DEG F -40.00 -30.00 -20.00 -10.00 10.00 20.00 30.00 40.00 50.00 40.00 70.00 80.00 70.00 100.00 110.00 110.00 130.00	DENS IN CALCD 0.608E+02 0.602E+02 0.596E+02 0.589E+02 0.589E+02 0.568E+02 0.564E+02 0.564E+02 0.564E+02 0.553E+02 0.553E+02 0.536E+02 0.518E+02 0.518E+02 0.518E+02 0.498E+02 0.475E+02 0.475E+02 0.463E+02 0.463E+02	LB/FT3 PUBLD 0.649E+02 0.643E+02 0.636E+02 0.629E+02 0.629E+02 0.622E+02 0.615E+02 0.607E+02 0.607E+02 0.592E+02 0.592E+02 0.550E+02 0.550E+02 0.520E+02 0.509E+02 0.509E+02 0.509E+02	ABS ERROR -0.416E+01 -0.408E+01 -0.402E+01 -0.397E+01 -0.397E+01 -0.397E+01 -0.389E+01 -0.389E+01 -0.390E+01 -0.392E+01 -0.401E+01 -0.401E+01 -0.426E+01 -0.450E+01 -0.465E+01 -0.465E+01	PERCENT ERROR -6.40 -6.35 -6.33 -6.32 -6.33 -6.32 -6.33 -6.41 -6.48 -6.58 -6.71 -6.87 -7.07 -7.30 -7.56 -7.88 -8.24 -8.65 -9.13 -9.44	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19	TR 0.6030 0.6174 0.6317 0.6461 0.6605 0.6749 0.6892 0.7036 0.7180 0.7323 0.7467 0.7611 0.7754 0.7898 0.8042 0.8185 0.8329 0.8473 0.8473	PR 0.0109 0.0143 0.0186 0.0238 0.0301 0.0377 0.0467 0.0573 0.0696 0.0840 0.1004 0.1094 0.1094 0.1193 0.1407 0.1649 0.1921 0.2226 0.2565 0.2942 0.3340
TEMP DEG F -40.00 -30.00 -20.00 -10.00 10.00 20.00 30.00 40.00 50.00 40.00 70.00 80.00 70.00 100.00 110.00 110.00 120.00 140.00 150.00	DENS IN CALCD 0.608E+02 0.602E+02 0.596E+02 0.589E+02 0.589E+02 0.568E+02 0.564E+02 0.564E+02 0.553E+02 0.553E+02 0.554E+02 0.518E+02 0.518E+02 0.518E+02 0.498E+02 0.498E+02 0.475E+02 0.450E+02 0.450E+02 0.450E+02 0.450E+02	LB/FT3 PUBLD 0.649E+02 0.643E+02 0.636E+02 0.629E+02 0.629E+02 0.622E+02 0.615E+02 0.607E+02 0.607E+02 0.592E+02 0.592E+02 0.550E+02 0.550E+02 0.520E+02 0.509E+02 0.509E+02 0.509E+02 0.498E+02 0.486F+02	ABS ERROR -0.416E+01 -0.408E+01 -0.402E+01 -0.397E+01 -0.397E+01 -0.397E+01 -0.389E+01 -0.389E+01 -0.390E+01 -0.392E+01 -0.401E+01 -0.408E+01 -0.426E+01 -0.450E+01 -0.450E+01 -0.481E+01 -0.481E+01	PERCENT ERROR -6.40 -6.35 -6.33 -6.32 -6.33 -6.32 -6.33 -6.41 -6.48 -6.58 -6.71 -6.87 -7.07 -7.58 -7.88 -8.24 -8.65 -9.13 -9.66 -10.28	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20	TR 0.6030 0.6174 0.6317 0.6461 0.6605 0.6749 0.6892 0.7036 0.7180 0.7323 0.7467 0.7611 0.7754 0.7898 0.8042 0.8185 0.8329 0.8473 0.8616 0.8740	PR 0.0109 0.0143 0.0186 0.0238 0.0301 0.0377 0.0467 0.0573 0.0696 0.0840 0.1004 0.1094 0.1094 0.1193 0.1407 0.1649 0.1921 0.2226 0.2565 0.2942 0.3360 0.3820
TEMP DEG F -40.00 -30.00 -20.00 -10.00 10.00 20.00 30.00 40.00 50.00 50.00 40.00 70.00 80.00 70.00 100.00 110.00 110.00 120.00 130.00 140.00 160.00	DENS IN CALCD 0.608E+02 0.602E+02 0.596E+02 0.589E+02 0.589E+02 0.568E+02 0.568E+02 0.564E+02 0.553E+02 0.553E+02 0.554E+02 0.518E+02 0.518E+02 0.508E+02 0.498E+02 0.498E+02 0.475E+02 0.450E+02 0.450E+02 0.436E+02 0.436E+02 0.421E+02	LB/FT3 PUBLD 0.649E+02 0.643E+02 0.636E+02 0.629E+02 0.629E+02 0.622E+02 0.615E+02 0.607E+02 0.607E+02 0.592E+02 0.592E+02 0.550E+02 0.550E+02 0.520E+02 0.509E+02 0.509E+02 0.509E+02 0.498E+02 0.473E+02	ABS ERROR -0.416E+01 -0.408E+01 -0.402E+01 -0.397E+01 -0.397E+01 -0.397E+01 -0.389E+01 -0.389E+01 -0.390E+01 -0.390E+01 -0.401E+01 -0.408E+01 -0.426E+01 -0.450E+01 -0.450E+01 -0.450E+01 -0.481E+01 -0.497E+01 -0.519E+01	PERCENT ERROR -6.40 -6.35 -6.33 -6.32 -6.33 -6.32 -6.33 -6.41 -6.48 -6.58 -6.71 -6.87 -7.07 -7.58 -7.88 -8.24 -8.65 -9.13 -9.66 -10.28 -10.98	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21	TR 0.6030 0.6174 0.6317 0.6461 0.6605 0.6749 0.6892 0.7036 0.7180 0.7323 0.7467 0.7611 0.7754 0.7898 0.8042 0.8185 0.8329 0.8473 0.8616 0.8760 0.8904	PR 0.0109 0.0143 0.0186 0.0238 0.0301 0.0377 0.0467 0.0573 0.0696 0.0840 0.1004 0.1004 0.1193 0.1407 0.1649 0.1921 0.2226 0.2565 0.2942 0.3360 0.3820 0.4325
TEMP DEG F -40.00 -30.00 -20.00 -10.00 10.00 20.00 30.00 40.00 50.00 40.00 70.00 80.00 70.00 100.00 110.00 120.00 130.00 140.00 150.00 160.00	DENS IN CALCD 0.608E+02 0.602E+02 0.596E+02 0.589E+02 0.589E+02 0.568E+02 0.564E+02 0.564E+02 0.553E+02 0.553E+02 0.554E+02 0.518E+02 0.518E+02 0.508E+02 0.498E+02 0.475E+02 0.436E+02 0.436E+02 0.421E+02 0.405E+02	LB/FT3 PUBLD 0.649E+02 0.643E+02 0.636E+02 0.629E+02 0.629E+02 0.622E+02 0.615E+02 0.607E+02 0.607E+02 0.592E+02 0.592E+02 0.550E+02 0.550E+02 0.550E+02 0.520E+02 0.509E+02 0.509E+02 0.478E+02 0.473E+02 0.459E+02	ABS ERROR -0.416E+01 -0.408E+01 -0.402E+01 -0.397E+01 -0.397E+01 -0.397E+01 -0.389E+01 -0.389E+01 -0.390E+01 -0.390E+01 -0.401E+01 -0.408E+01 -0.416E+01 -0.426E+01 -0.450E+01 -0.450E+01 -0.450E+01 -0.499E+01 -0.519E+01 -0.519E+01 -0.540E+01	PERCENT ERROR -6.40 -6.35 -6.33 -6.32 -6.33 -6.32 -6.33 -6.41 -6.48 -6.58 -6.71 -6.87 -7.07 -7.58 -7.88 -8.24 -8.65 -9.13 -9.66 -10.28 -10.98 -11.77	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22	TR 0.6030 0.6174 0.6317 0.6461 0.6605 0.6749 0.6892 0.7036 0.7180 0.7323 0.7467 0.7611 0.7754 0.7898 0.8042 0.8185 0.8329 0.8473 0.8616 0.8760 0.8904 0.9047	PR 0.0109 0.0143 0.0186 0.0238 0.0301 0.0377 0.0467 0.0573 0.0467 0.0573 0.0696 0.0840 0.1004 0.1004 0.1193 0.1407 0.1449 0.1921 0.2226 0.2565 0.2942 0.3360 0.3820 0.4325 0.4879
TEMP DEG F -40.00 -30.00 -20.00 -10.00 10.00 20.00 30.00 40.00 50.00 40.00 50.00 40.00 70.00 100.00 100.00 110.00 120.00 130.00 140.00 150.00 160.00 170.00	DENS IN CALCD 0.608E+02 0.602E+02 0.596E+02 0.589E+02 0.589E+02 0.568E+02 0.564E+02 0.564E+02 0.554E+02 0.554E+02 0.554E+02 0.558E+02 0.518E+02 0.518E+02 0.508E+02 0.487E+02 0.487E+02 0.463E+02 0.450E+02 0.436E+02 0.436E+02 0.436E+02 0.436E+02 0.436E+02 0.436E+02 0.436E+02 0.436E+02 0.436E+02 0.436E+02 0.436E+02 0.436E+02 0.436E+02 0.436E+02 0.436E+02 0.436E+02	LB/FT3 PUBLD 0.649E+02 0.643E+02 0.636E+02 0.629E+02 0.629E+02 0.615E+02 0.607E+02 0.607E+02 0.592E+02 0.592E+02 0.550E+02 0.550E+02 0.550E+02 0.550E+02 0.520E+02 0.509E+02 0.509E+02 0.486E+02 0.486E+02 0.473E+02 0.459E+02 0.459E+02	ABS ERROR -0.416E+01 -0.408E+01 -0.402E+01 -0.377E+01 -0.373E+01 -0.379E+01 -0.387E+01 -0.387E+01 -0.370E+01 -0.370E+01 -0.401E+01 -0.408E+01 -0.416E+01 -0.426E+01 -0.450E+01 -0.450E+01 -0.457E+01 -0.457E+01 -0.457E+01 -0.457E+01 -0.517E+01 -0.517E+01 -0.540E+01 -0.563E+01	PERCENT ERROR -6.40 -6.35 -6.33 -6.32 -6.33 -6.32 -6.33 -6.41 -6.48 -6.58 -6.71 -6.87 -7.07 -7.56 -7.88 -8.24 -8.65 -9.13 -9.66 -10.28 -10.98 -11.77 -12.69	N 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23	TR 0.6030 0.6174 0.6317 0.6461 0.6605 0.6749 0.6892 0.7036 0.7180 0.7323 0.7467 0.7611 0.7754 0.7611 0.7754 0.7898 0.8042 0.8185 0.8329 0.8473 0.8473 0.8616 0.8760 0.8904 0.9047 0.9191	PR 0.0109 0.0143 0.0186 0.0238 0.0301 0.0377 0.0467 0.0573 0.0696 0.0840 0.1004 0.1004 0.1097 0.1407 0.1649 0.1921 0.2226 0.2565 0.2942 0.3360 0.3820 0.4325 0.4879 0.5484

190.00

200.00

210.00

AAD=

SD2= 9.3973741E-02

0.368E+02

0.346E+02

0.427E+02 -0.586E+01 0.407E+02 -0.609E+01

0.322E+02 0.384E+02 -0.629E+01

-13.73

-14.95

-16.36

24

25

26

0.9335

0.9478

0.9622

0.6144

0.6862

0.7640

************* PRESSURE COMPARISON ****************

TABLE E8

A CONTRACTOR OF A CONTRACTOR O							
TEMP DEG F	PRESSUR Calcd	E PSIA PUBLD	ABS Error	PERCENT Error	N	TR	PR
0.00	0.571E+03	0.611E+03	-0.404E+02	-6.61	1	0.4220	0.0000
10.00	0.116E+04	0.123E+04	-0.631E+02	-5.14	2	0.4374	0.0001
20.00	D-225E+04	0.234E+04	-0.926E+02	-3,96	3	0.4529	0.0001
30.00	0.412E+04	0.425E+04	-0.121E+D3	-2.86	4	0.4683	0.0602
40.00	D.725E+04	0.738E+04	-0.136E+03	-1.84	5	0.4838	0.0003
50.00	0.122E+05	0.123E+05	-0.116E+03	-0.94	6	0.4992	0.0006
60.00	0.199E+05	0.199E+05	-0.300E+02	-0.15	7	0.5147	0.0009
70.00	0.314E+05	0.312E+05	D.166E+D3	0.53	8	0.5301	0.0014
80.00	0.479E+05	0.474E+05	0.538E+03	1.14	9	0.5456	0.0022
90.00	0.713E+05	0.701E+05	0.116E+04	1.65	10	D.5610	0.0032
100.00	0.103E+06	0.101E+06	0.211E+04	2.08	11	0.5765	0.0047
110.00	0.147E+06	0.143E+06	0.350E+04	2.44	12	0.5919	0.0067
120.00	0.204E+06	0.199E+06	0.542E+04	2.73	13	0.6074	0.0093
130.00	0.278E+06	0.270E+06	0.798E+D4	2.95	14	0.6228	0.0126
140.00	0.373E+06	0.361E+06	0.113E+05	3.13	15	0.6383	0.0169
150.00	0.491E+06	0.476E+06	0.156E+05	3.27	16	0.6537	0.0223
160.00	0.637E+06	0.618E+D6	0.207E+05	3.36	17	0.6692	0.0290
170.00	0.819E+D6	0.792E+06	0.269E+05	3.40	18	D.6846	0.0371
180.00	0.104E+07	0.100E+07	0.344E+05	3.43	19	0.7001	0.0470
190.00	0.130E+07	0.125E+07	0.429E+05	3.42	20	0.7155	0.0588
200.00	0.161E+07	0.155E+07	0.529E+05	3.40	21	0.7310	0.0729
210.00	0.197E+07	0.191E+07	0.640E+05	3.36	22	0.7464	0.0894
220.00	0.239E+07	0.232E+07	0.762E+05	3.29	23	0.7619	0.1086
230.00	D.288E+07	0.280E+07	0.899E+05	3.21	24	0.7773	0.1308
240.00	0.345E+07	0.334E+07	0.105E+06	3.14	25	0.7928	0.1564
250.00	0.409E±07	0.397E+07	0.120E+06	3.03	26	0.8082	0.1857
260.00	D.483E+07	0.469E+07	0.1372+06	2.93	27	0.8237	0.2189
270.00	0.565E+07	0.550E+07	0.154E+06	2.80	28	0.8391	0.2564
280.00	0.658E+07	C.641E+C7	0.171E+06	2.67	29	0.8546	0.2986
290.00	0.762E+07	0.744E+07	0.188E+06	2.53	30	0.8700	0.3458
300.00	0.879E+07	0.858E+07	0.204E+C6	2.38	31	0.8855	0.3985
310.00	0.101E+C8	0,985E+07	0.217E+06	2.22	32	0.9009	0.4569
320.00	0.115E+08	0.113E+08	0.2282+06	2.02	33	0.9164	0.5217
330.00	0.131E+08	0.128E+08	0.232E+06	1.80	34	0.9318	0.5931
340.00	0.148E+08	0.146E+D8	0.2232+06	1.53	35	0.9473	0.6717
350.00	D.167E+D8	0.165E+08	0.197E+06	1.19	36	0.9627	0.7579
360.00	0.188E+08	0.187E+08	0.139E+06	0.74	37	0.9782	0.8522
370.00	0.211E+D8	0.210E+08	0.312E+05	0.15	38	0.9936	0.9552

AAD= SD1= SD2= 2.564522 111741.2 2.9003220E-02

TABLE E9

TEMP DEG F	PRESSUR Calcd	E PSIA PUBLD	ABS Error	PERCENT Error	N	TR	PR
-40.00	0.107E+02	0.109E+02	-0.242E+00	-2.21	1	0.6158	0.0167
-30.00	0.138E+02	0.141E+02	-0.287E+00	-2.05	2	0.6304	0.0215
-20.00	0.176E+02	0.179E+02	-0.334E+00	-1.87	3	0.6451	0.0274
-10.00	0.221E+02	0.225E+02	-0.395E+00	-1.75	4	0.6598	0.0345
0.00	0.275E+02	0.280E+02	-0.449E+00	-1.60	5	0.6744	0.0429
10.00	0.339E+D2	0.344E+02	-0.521E+00	-1.51	6	0.6891	0.0528
20.00	0.414E+02	0.420E+02	-0.587E+00	-1.40	7	0.7038	0.0645
30.00	0.500E+02	0.507E+02	-0.666E+00	-1.31	8	D.7185	0.0779
40.00	0.600E+02	0.608E+02	-0.733E+00	-1.21	9	0.7331	0.0935
50.00	0.714E+02	0.723E+02	-0.812E+00	-1.12	10	0.7478	0.1113
60.00	0.845E+02	0.853E+02	-0.875E+00	-1.03	11	0.7625	0.1316
70.00	0.992E+02	0.100E+03	-0.927E+00	-0.93	12	0.7771	0.1545
80.00	0.116E+03	0.117E+03	-0.960E+00	-0.82	13	0.7918	0.1803
90.00	0.134E+03	0.135E+03	-0.100E+01	-0.74	14	0.8065	0.2092
100.00	0.155E+03	0.156E+03	-0.902E+00	-0.58	15	0.8212	0.2415
110.00	0.178E+03	0.179E+03	-0.813E+00	-0.45	16	0.8358	0.2773
120.00	0,203E+03	0.204E+03	-0.679E+00	-0.33	17	0.8505	0.3169
130.00	D.231E+03	0.232E+03	-0.438E+00	-0.19	18	0.8652	0.3606
140.00	0.262E+03	0.262E+03	-0.125E+00	-0.05	19	0.8798	0.4086
150.00	0.296E+03	0.296E+03	0.231E+00	0.08	20	0.8945	0.4612
160.00	0.333E+03	0.332E+03	0.804E+00	0.24	21	0.9092	0.5186
170.00	0.373E+03	0.372E+03	0.138E+01	0.37	22	0.9239	0.5812
180.00	0.417E+03	0.415E+03	0.215E+01	0.52	23	0.9385	0.6492
190.00	0.464E+03	0.461E+03	0.311E+01	0.67	24	0.9532	0.7230
200.00	0.515E+03	0.511E+03	0.426E+01	0.83	25	0.9679	0.8029
210.00	0.571E+03	0.567E+03	0.342E+01	0.60	26	0.9825	0.8891
220.00	0.630E+03	0.630E+03	0.710E+00	0.11	27	0.9972	0.9821
AAD=	0.9106809						
SD1=	1.465343						82

SD2= 1.1217823E-D2

LIQUID DENSITY COMPARISON

______ TEMP DENS IN LB/FT3 ABS DEG F CALCD PUBLD ERROR ABS PERCENT N PR TR DEG F ERROR -40.000.904E+020.843E+020.609E+017.2310.6158-30.000.894E+020.833E+020.606E+017.2720.6304-20.000.884E+020.824E+020.601E+017.2930.6451-10.000.874E+020.814E+020.593E+017.2840.65980.000.863E+020.805E+020.583E+017.2550.674410.000.852E+020.795E+020.571E+017.1960.689120.000.840E+020.784E+020.556E+017.0970.703830.000.828E+020.774E+020.537E+016.9480.718540.000.815E+020.774E+020.517E+016.7790.733150.000.802E+020.733E+020.473E+016.55100.747860.000.773E+020.743E+020.435E+016.27110.762570.000.778E+020.730E+020.435E+015.97120.777180.000.758E+020.718E+020.401E+015.97130.791890.000.742E+020.706E+020.364E+015.15140.80650.0167 0.0215 0.0274 0.0345 0.0429 0.0528 0.0645 0.0779 0.0935 0.1113 0.1316 0.773E+02 0.730E+02 0.435E+01 0.758E+02 0.718E+02 0.401E+01 0.742E+02 0.706E+02 0.364E+01 0.725E+02 0.693E+02 0.322E+01 0.707E+02 0.679E+02 0.276E+01 0.1545 13 14 0.1803 0.8065 5.15 0.2092 90.00 100.00 15 0.8212 0.2415 4.65 15 16 17 18 19 20 21 22 0.8358 110.00 4.05 0.2773 0.688E+02 0.666E+02 0.225E+01 0.668E+02 0.651E+02 0.170E+01 0.646E+02 0.635E+02 0.111E+01 0.623E+02 0.618E+02 0.454E+00 0.8505 120.00 3.39 0.3169 0.8652 2.61 130.00 0.3606 1.74 0.8798 140.00 0.4086 0.73 150.00 0.8945 0.4612 0.598E+02 0.600E+02 -0.252E+00 0.570E+02 0.580E+02 -0.101E+01 -0.42 0.9092 160.00 0.5186 170.00 0.9239 0.5812 0.540E+02 0.558E+02 -D.182E+01 180.00 23 0.9385 0.6492 -3.26 24 25 26 27 190.00 0.506E+02 0.533E+02 -0.270E+01 -5.06 0.9532 0.7230 0.465E+02 0.502E+02 -0.362E+01 0.415E+02 0.460E+02 -0.451E+01 0.336E+02 0.384E+02 -0.476E+01 -7.21 200.00 0.9679 0.8029 0.9825 210.00 ~9.80 0.8891 0.9972 220.00 -12.39 0.9821

AAD= 5.587523 SD1= 4.330630 SD2= 6.3137342E-02 TABLE E9 CONTINUED

TABLE E9 CONTINUED

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TEMP Deg F	H IN B Calcd	TU/LB Publd	ABS Error	PERCENT Error	N	TR	PR
-40.00	0.000E+00	0.000£+00	0.000E+00	0.00	1	0.6158	0.0167
-30.00	0.259E+01	0.238E+01	0.211E+00	8.85	2	0.6304	0.0215
20.00	0.520E+01	0.479E+01	0.411E+00	8.59	3	0.6451	0.0274
-10.00	0.784E+01	0.722E+01	D.615E+00	8.52	4	0.6598	0.0345
0.00	0.105E+02	0.971E+01	0.785E+00	8.09	5	0.6744	0.0429
10.00	0.132E+02	0.122E+02	0.954E+00	7.80	6	0.6891	0.0528
20.00	0.159E+02	0.148E+02	0.112E+01	7.55	. 7	0.7038	0.0645
30.00	0.187E+02	0.174E+D2	0.126E+01	7.26	8	0.7185	0.0779
40.00	0.215E+02	0.200E+02	0.141E+01	7.04	9	0.7331	0.0935
50.00	0.243E+02	0.228E+02	0.155E+01	6.82	10	0.7478	0.1113
60.00	0.272E+02	0.255E+02	0.171E+01	6.72	11	0.7625	0.1316
70.00	0.301E+02	0.283E+02	0.185E+01	6.55	12	0.7771	0.1545
80.00	0.331E+02	0.311E+02	0.201E+01	6.47	13 :	0.7918	0.1803
90.00	D.362E+02	0.340E+02	0.219E+01	6.44	14	0.8065	0.2092
100.00	0.393E+02	0.370E+02	0.237E+01	6.40	15	0.8212	0.2415
110.00	0.426E+02	0.400E+02	0.255E+01	6.38	16	0.8358	0.2773
120.00	0.459E+02	0.431E+02	0.276E+01	6.41	17	0.8505	0.3169
130.00	0.493E+02	0.463E+02	0.298E+01	6.45	18	0.8652	0.3606
140.00	0.528E+02	0.496E+02	0.322E+01	6.50	19	0.8798	0.4086
150.00	0.565E+D2	0.530E+02	0.349E+01	6.59	20	0.8945	0.4613
160.00	0.603E+02	0.565E+02	0.380E+01	6.73	21	0.9092	0.5186
170.00	0.644E+02	0.602E+02	0.414E+01	6.88	22	D.9239	0.5812
180.00	0.687E+02	0.641E+02	0.454E+01	7.08	23	0.9385	0.6493
190.00	0.733E+02	0.683E+02	0.503E+01	7.36	24	0.9532	0.7230
200.00	0.785E+02	0.729E+02	0.566E+01	7.77	25	0.9679	0.8029
210.00	D.847E+02	0.768E+02	0,793E+D1	10.33	26	0.9825	0.8891
220.00	D.937E+D2	0.840E+02	0.767E+01	8.92	27	0.9972	0.9821
AAD=	7.055314						
SD1=	3.436757						
SD2=	7.3944241E-0	2					
*****	*******						TINUES
VAPOUR	ENIHALPY					INGLE FA COM	THOED

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TEMP BTU/LB DEG F CALCD PUBLD		PUBLD	ABS Error	PERCENT Error	N	TR	PR
-40,00	0.895E+02	0.877E+02	0.178E+01	2.03	1	0.6158	0.0167
-30.00	0.910E+02	0.890E+02	0.199E+01	2.24	2	0.6304	0.0215
-20.00	0.925E+02	0.903E+02	0.222E+01	2.45	3	0.6451	0.0274
-10.00	0.940E+02	0.916E+02	0.243E+01	2.65	4	0.6598	0.0345
0.00	0.954E+02	0.928E+02	0.264E+01	2.85	5	0.6744	0.0429
10.00	0.969E+02	0.940E+02	0.283E+01	3.01	6	0.6891	0.0528
20.00	0.983E+02	0.952E+02	0.303E+01	3.18	7	0.7038	0.0645
30.00	0.996E+02	0.964E+D2	0.322E+01	3.34	8	0.7185	0.0779
40.00	0.101E+03	0.975E+02	0.340£+01	3.49	9	0.7331	0.0935
50.00	0.102E+03	0.986E+D2	0.357E+01	3.62	10	0.7478	0.1113
60.00	0.103E+03	0.997E+02	0.374E+01	3.76	11	0.7625	0.1316
70.00	0.105E+03	0.101E+03	0.390E+01	3.87	12	0.7771	0.1545
80.00	0.106E+03	0.102E+03	0.404E+01	3.97	13	D.7918	0.1803
90.00	0.107E+03	0.103E+03	0.417E+01	4.06	14	0.8065	0.2092
100.00	0.108E+03	0.104E+03	0.429E+01	4.14	15	0.8212	0.2415
110.00	0.109E+03	0.104E+03	0.439E+01	4.20	16	0.8358	0.2773
120.00	0.110E+03	0.105E+03	0.446E+01	4.24	17	0.8505	D.3169
130.00	D.110E+03	0.106E+D3	0.453E+01	4.28	18	0.8652	0.3606
140.00	0.111E+03	0.107E+03	0.457E+01	4.29	19	0.8798	0.4086
150.00	0.112E+03	0.107E+03	0.459E+01	4.29	20	0.8945	0.4612
160.00	0.112E+03	0.107E+03	0.457E+01	4.26	21	0.9092	0.5186
170.00	0.112E+03	0.108E+03	0.453E+01	4.21	22	0.9239	0.5812
180.00	0.112E+03	0.107E+03	0.445E+01	4.14	23	0.9385	0.6492
190.00	0.111E+03	0.107E+03	0.431E+01	4.02	24	0.9532	0.7230
200.00	0.110E+03	0.106E+D3	0.408E+01	3.83	25	0.9679	0.8029
210.00	0.108E±03	0.104E+03	0.400E+01	3.83	26	0.9825	0.8891
220.00	0.103E+03	0.987E+02	0.444E+01	4.50	27	0.9972	0.9821
AAD=	3.657464	8					
004-	7 880480						

SD1= 3.882189 SD2= 3.7921775E-D2

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PERCHLOROETHYLENE

TABLE E10

TEMP	PRESSUR	E PSIA	ABS	PERCENT	N	TR	PR
DEG F	CALCD	PUBLD	ERROR	ERROR			
-40.00	0.229E+02	0.211E+02	0.170E+01	8.05	1	0.3761	0.0000
-30.00	0.598E+02	0.543E+02	0.553E+01	10.17	2	0.3922	0.0000
-20.00	0.134E+03	0.127E+03	0.756E+01	5.96	3	0.4083	0.0000
-10.00	0.281E+03	0.274E+03	0.740E+01	2.70	4	0.4245	0.0001
0.00	0.552E+03	0.550E+03	0.184E+01	0.33	5	0.4406	0.0001
10.00	0.103E+04	0.104E+04	-0.120E+02	-1.15	6	0.4567	0.0002
20.00	0.183E+04	0.187E+04	-0.459E+02	-2.45	7	0.4728	0.0004
30.00	0.310E+04	0.321E+04	-0.106E+03	-3.30	8	0.4890	0.0007
40.00	0.507E+04	0.527E+04	-0.203E+03	-3.86	9	0.5051	0.0011
50.00	0.801E+04	0.836E+04	-0.352E+03	-4.21	10	0.5212	0.0017
60.00	0.123E+05	0.128E+05	-0.567E+03	-4.42	11	0.5374	0.0026
70.00	0.182E+05	0.191E+05	-0.868E+03	-4.54	12	0.5535	0.0038
80.00	0.264E+05	0.277E+05	-0.127E+04	-4.60	13	0.5696	0.0056
90.00	0.375E+05	0.393E+05	-0.180E+04	-4.59	14	0.5857	0.0079
100.00	0.5192+05	D.544E+05	-0.248E+04	-4.55	15	0.6019	0.0109
110.00	0.706E+05	0.740E+05	-D.332E+D4	-4.49	° 16	0.6180	0.0148
120.00	0.944E+05	0.987E+05	-0.435E+04	-4.41	17	0.6341	0.0198
130.00	0.124E+06	0.130E+06	-0.558E+04	-4.31	18	0.6503	0.0260
140.00	0.161E+06	0.168E+06	-0.701E+04	-4.19	19	0.6664	0.0337
150.00	0.205E+06	0,214E+06	-0.865E+04	-4.04	20	0.6825	0.0431
160.00	0.259E+06	0.269E+06	-0.105E+05	-3.88	21	0.6786	0.0543
170.00	0.323E+06	0.335E+06	-0.124E+05	-3.70	22	0.7148	0.0677
180.00	0.3985+04	0.412E+06	-D.144E+05	-3.49	23	0.7309	0.0836
190.00	0.4865+06	0.503E+06	-0.164E+05	-3.25	24	0.7470	0.1021
200 00	0 5892+06	D. 607E+06	-0.181E+05	-2.99	25	0.7632	0.1236
210.00	0.707E+06	0.726E+06	-0.195E+05	-2.68	26	0.7793	0.1484
220 00	D 847E+04	0 8675+06	-0.2025+05	-2.35	27	0.7954	0.1767
230.00	0.9945+04	n.102E+07	-0.200E+05	-1.97	28	0.8115	0.2090
240.00	0 1176+07	0-1195+07	-0.184E+05	-1.55	29	0.8277	0.2455
250.00	0.1375+07	0 1385+07	-0.151E+05	-1.09	30	0.8438	0.2856
250.00	0.1572+07	0.1595+07	-R 931E+64	-0.58	31	0.8577	0.3326
280.00	0.1975+07	0.1975+07	-0 5255+0%	-0.03	32	0.8761	0.3840
270.00	0.1032+07	0.1032+07	-0.525E+05	0.58	33	0.8922	0.4410
200.00	0.2100407	0.2072+07	0.1216.03	1.24	34	0.9083	0.5042
270,00	0.2776107	0.2425107	0.5325405	1.95	35	0.9245	0.5739
300.00	0.2/36+07	0.2002707	n 8245+05	2.72	36	0.9406	0,6505
310.00	0.3100707	D 3725+07	0.0212+03	3.55	37	0.9567	0.7346
320.00	0.3306707	0.3306407	D.1205+00	6.44	38	0.9728	0.8265
330.00	0.3745707	0.3//240/	0.1002-00	5 40	20	0.9890	0.9268
340.00	0.4422+07	0.4176+07	0.20/5+04	3.7U 1 77	40	1.0051	1.0352
350.00	U.493E+U/	U.464E+U/	U.274E+U0	0.33	70	1.0031	

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AAD= 3.503570 SD1= 70456.25 SD2= 4.0931337E-02

BROMOTRIFLUOROMETHANE

I VAP MOL CONC VAP MASS CONC I LIQ MASS CONC I LIQ MOL CONC E PRESSURE (PA) TEMP I PUB I CALC I PUB I CALC I PUB I CALC I PUB I CALC I 1 CALC PUB DEG C I _____ · · · I. 1 1 0.6296 | 0.6321 | 1 1 0.2491 1 -17.80 | 0.30310E+06 | 0.30871E+06 | 1 I 0.7593 I 0.7625 I 1 0.4985 1 -17.80 | 0.39543E+06 | 0.38924E+06 | 1 1 0.8570 1 0.8598 1 1 0.7483 1 -17.80 | 0.45125E+06 | 0.44387E+06 | 1 0.6817 | 0.6845 | -17.80 | 0.32655E+06 | 0.33954E+06 | 1 0.3277 1 | 0.7162 | 0.7192 | 1 0.3956 1 -17.80 | 0.36172E+06 | 0.36145E+06 | 1 0.7612 1 0.7644 1 1 0.5034 1 -17.80 | 0.39410E+06 | 0.39045E+06 | I 0.8119 | 0.8151 | -17.80 1 0.42714E+06 1 0.42087E+06 1 1 0.6370 1 I 0.8792 | 0.8817 | -17.80 | 0.46032E+06 | 0.45354E+06 | I 0.7974 I 1 0.9541 1 0.9552 1 0.9343 -17.80 | 0.48443E+06 | 0.47855E+06 | 1 0,9833 1 0.9837 1 1 0.9774 1 0.48989E+06 | 0.48708E+06 | -17.70 1 1.0000 1 1.0000 1 1 1.0000 1 -17.60 | 0.49055E+06 | 0.49204E+06 | I 0.5041 I 0.4977 I I 0.2475 I 25.00 | 0.10349E+07 | 0.10381E+07 | 1 0.6811 1 0.6771 1 1 0.4959 1 25.00 | 0.13070E+07 | 0.12968E+07 | 1 0.8229 | 0.8210 | 1 0.7465 .1 0.15055E+07 | 0.14843E+07 | 25.00 | 1 1.0000 | 1.0000 | 1 1.0000 l 25.00 | 0.16174E+07 | 0.16192E+07 | | 0.4187 | 0.4121 | 1 0.2440 1 54.40 | 0.20091E+07 | 0.20099E+07 | I 0.6185 | 0.6135 | 1 0.4932 1 54.40 | 0.25061E+07 | 0.24850E+07 | 1 0.7920 | 0.7896 | 1 0.7448 1 54.40 | 0.28711E+07 | 0.28490E+07 | 1 1.0000 | 1.0000 | 1 1.0000 1 0.30923E+07 | 0.30969E+07 | 54.40 1

AND DIFLUOROETHANE

CHLORODIFLUOROMETHANE

AND TRICHLOROFLUOROMETHANE

TEMP	1	PRES	sui	RE (PA)	 	LIQ	MASS	CONC		LIQ	HOL	CONC	 I	VAP	MASS	CONC	Ī	VAP	MOL	CONC	1
DEG C	Ē	PUB	1	CALC	1	PUB	I	CALC	1	PUB	- 1	CALC	1	PUB	l 	CALC	1	PUB	1	CALC	
	1		1		1		1		1		1		I	*	1		1		1		1
25.00	i.	0.31000E+06	1	0.29925E+06	1		1		1	0.156	8		1		1		1	0.678	38 1	0.6847	1
25.00	î.	0.40000E+06	-î	0.40975E+06	1		1		1	0.259	0 1		1		1		1	0.783	31 I	0.7869	I
25.00	i.	0.51000E+06	1	0.53396E+06	1		1		1	0.388	4 1		1	14	1		1	0.852	21 1	0.8543	1
25,00	i.	0.66000E+06	î	0.66486E+06	1		. 1		1	0.542	6 1		1	а. 8	1		1	0.90	10	0.9018	1
25.00	1	0.78000E+06	1	0.78392E+06	1		1		1	0.694	2 1		1		1		1	0.93	55 1	0.9356	1
25.00	i.	0.90000E+06	Ť	0.90821E+06	1		1		1	0.850	5 1		1		1		I	0.96	72	0.9669	1
25.00	i.	0.92000E+06	1	0.92485E+06	1		I		1	0.870	4 1		1		1		1	0.97	13 1	0.9710	1
50.00	î.	0.57000E+06	1	0.55578E+06	1		1		1	0.156	8 1		1		1		1	0.60	73	0.6130	1
50.00	î.	0.74000E+06	E	0.74634E+06			1		1	0.259	0		1		1		1	0.72	54 1	0.7294	1
50.00	i i	0.96000E+06	-î	0.96726E+06	1		1		1	0.388	4 1		1		1		1	0.80	35 I	0.8108	1
50.00	î.	0.12000E+07	1	0.12089E+07	1		1		1	0.542	6 1		1		1		1	0.869	78	0.8708	1
50.00	î -	0.14300E+07	1	0.14366E+07	1		1		1	0.694	2 1		- E .,		1		1	0.914	45 1	0.9146	1
50.00	î.	0.16800E+07	- î	0.16804E+07	1		1		I	0.850	5 1		1		1		1	0.95	52 1	0.9560	1
50.00	i -	0.17100E+07	i	0.17134E+07	1		1		1	0.870	4 1		1		1		1	0.96	17 1	0.9614	1
75.00	î.	0.96000E+06	1	0.94668E+06	1		1		1	0.156	8 1		1		1		1	0.53	53 I	0.5409	1
75.00	î.	0.12300E+07	î.	0.12444E+07	1		1		1	0.259	0 1		1		1		1	0.663	26 1	0.6663	1
75.00	î.	0.15900E+07	È	0.16001E+07	1		1		1	0.388	4 Î		1		1		1	0.75	72 1	0.7594	1
75.00	î.	0.19800E+07	ì	0.20032E+07	1		1		1	0.542	6 1		1		1		1	0.83	33 1	0.8312	1
75.00	î.	0.23700E+07	î	0.23974E+07	1		1		L	0.694	2 1		1		1		1	0.88	55 1	0.8856	1
87.00	î.	0.12500E+07	÷Ì.	0.12364E+07	1		1		I	0.156	8 1		I		1		1	0.49	58	0.5004	I
95.00	î.	0.17700E+07	- î	0.17799E+07	1		1		1	0.259	0 1		1		1		1	0.60	BO 1	0.6114	1
100.00	i	0.30500E+07	i	0.30756E+07	1	÷.	i		1	0.542	6 1		I		1		1	0.77	75 I	0.7781	1

TEMP	 I	PRESSURE (PA)		LIQ MASS	CONC		LIQ MO	L CONC		VAP MA	SS CO	DNC	1 1	AP MOL	CONC
DEG C	् ।	PUB I CALC		PUB I	CALC	!	PUB	I CALC	I	PUB	I C/	ALC		PUB	CALC
		Image: Construct of the construction of the constrelation of the constrelation of the construction of the construct	1 06	I 1.0000 I 0.9499 I 0.8998 I 0.8018 I 0.7477 I 0.7496 I 0.7395 I 0.7395 I 0.7302 I 0.7302 I 0.7200 I 0.6984 I 0.5972 I 0.5001 I 0.3974 I 0.2912 I 0.1997 I	5.					1.0000 0.9224 0.8632 0.8219 0.7849 0.7610 0.7449 0.7410 0.7367 0.7367 0.7366 0.7253 0.7078 0.6455 0.5685 0.5685 0.4918 0.3894 0.3894		.0000 9182 8571 8090 7705 7511 7276 7276 7246 7217 7154 7022 .6465 .5885 .5237 .4422 .3511			
0.00	L	0.28682E+06 0.29282E+	06	0.0993 I		4		1		0.10/1					

AND

DICHLORODIFLUOROMETHANE

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DIFLUOROETHANE

TENP	1	PRESSUR	RE	(PA)	1	LIQ	MASS	CONC	I	LIQ	10L	CONC	I	VAP	MASS	CONC	T	VAP	MOL	CONC	1
DEG C	ī	PUB I		CALC	1	PUB	1	CALC	1	PUB	1	CALC	1	PUB	1	CALC	L	PUB	1	CALC	
					1		1		1		1		1		1		I		I		1
-18 10	÷.	0 214505+04	п.	212448+06	î –		î		í.	0.0360	ı İ.		1		I		1	0.230	11	0.2447	1
-18.10	÷.	0.272105+06	n.	26850E+06	î –		î		î.	0.079	i i		1		1		1	0.398	5 1	0.4178	1
-18 10	2	0 354705+04 1	n.	35988E+06	i		î.		î.	0.1539	Ĩ.		1		1		1	0.565	3 [0.5864	1
-18.10	1	0.338702700 1	n.	45757F+06	î -		i		÷.	0.2392			1		1		1	0.686	2 1	0.6945	1
-18 10	1	0 544205+04 1	n.	57001E+06	î -		i		÷.	0.346	i i		1		1		I.	0.779	3 L	0.7759	1
-18 40	4	0 474705+04 1	n.	48134F+04	i		i		÷.	0.4602	1		1		1		1	0.840	11	0.8331	1
-18.10	1	0.745005+04	о. П	794245+04	i		í		֔.	0.5841	i i		i -		1		1	0.887	7 🖡	0.8789	1
-18.10	1	0.703702400 1	n.	97887F+04	i –		- i		i	0.7817	1		1		1		1	0.946	0 1	0.9375	1
-18.10	1	0.440525+07	n.	440895+07	i i		- i		i	0.9123	ιî.		Î.		1		1	0.979	4 1	0.9741	1
-18.00	i	0.12078E+07	ο.	12086E+07	i –		i		i	1.0000	ī į		i		1		1	1.000	0 1	1.0000	1

AND

DICHLORODIFLUOROMETHANE

E15 R12 - R2:

CHLORODIFLUOROMETHANE

TEMP	I	PRESS	SUR	E (PA)	1	LIQ	MASS	CONC	1	LIQ	HOL	CONC	I	VAP	MASS	CONC	I	VAP	MOL	CONC	1
DEG C	Ē	PUB	T	CALC	1	PUB	1	CALC	1	PUB	1	CALC	I	PUB	I	CALC	1	PUB	1	CALC	
	7		7		1		1		1		1		1		1		1		1		1
-40 80	1	0 404335404	1	0 102045+04	i	1 000	n i	×.			i		- î -		i		i -		- î		1
-40.00	÷.	0.101332+04	i -	0.10254E+06	÷.	0.984	n i		i i		i i		- î -		i		i		1		1
-40.70	÷.	0.101332+00	1	0.102342.00	1	0.700	01		1		i		1		i		i		1		1
-40.80	÷.	0.101332+00	1	0.102076+06	1	0.771			1		- 1		i		i		i -		i		I
-/0.99	4	0.404775+04	4	0.102372700	1	0.740	0				- 2		1		- i		i		- i		1
-40.67	4	0.101332700	1	0.102002700	-	0.731					- 1				- i		÷		- 1		i
-40.77	÷.	0.101332400	÷.,	0.102042400	4	0.710					- 1		- 1		- i -		÷.		- î		-i
-41.31	4	0.101335+06	4	0.101002700	1	0.031							1		i		i		i		i
-41.41	4	0.101332400	4.1	0,101242+00	4	0.701			-				- i -		- i -		- í -		i		î
-41.41	5	U.10133E+06	8.1	U.1U1UUE+U6	÷.	0.724	U 1		1		- 2		1				-		- 1		÷
-41.39	÷.	U.10133E+06	1	U.100/1E+06	4	0.0/5					- 1		1		- 1		1		- 1		- 1
-41.39	5	0.10133E+06	1	0.999172+05	1	0.612					- 5						-		- 1		- 1
-40.93	1	0.10133E+06	1	0.10104E+06	1	0.556			1				1				1		1		1
-40.73	1	0.10133E+06	1	0.10025E+06	<u>.</u>	0.484	01										1		- 1		-1
-40.63	I	0.10133E+06	1	0.10004E+06	1	0.461	0 1		1				1				1		- 1		-1
-40.54	1	0.10133E+06	1	0.99294E+05	1	0.425	0 1		1				1				1		1		- 2
-40.54	1	0.10133E+06	1	D.98980E+05	1	0.416	0 1		1		1				1.0		1		1		1
-39.84	1	0.10133E+06	1	D.99774E+05	1	0.358	0		1		1		1						- !		1
-38.93	1	0.10133E+06	1 1	D.10092E+06	1	0.301	0 1		1		1		1				1		- 1		4
-38.33	1	0.10133E+06	1	0.10161E+06	1	0.269	0 1		1		1		1		1		1		- 1		1
-37.14	1	0.10133E+06	1	D.10294E+06	1	0.216	0 1		1		1		1				1		1		
-35.87	1	0.10133E+06	1.1	D.10330E+06	1	0.161	0 1		1		1		1		1		1		1		1
-34.26	L	0.10133E+06	1	0.10354E+06	1	0.106	0 1		1		1		1		1		1		1		

DICHLORODIFLUOROMETHANE

AND TRICHLOROTRIFLUOROETHANE

			2								CONC			MASS	CONC	 I	 VAP M	OL	CONC	
TEMP	1	PRESSURE	(PA)			MA55		۱ 												
DEG C	Ł	PUB I	CALC	ł	PUB	E.	CALC	I	PUB	1	CALC	ا 	PUB		CALC		PUB		CALC	
				1		1		 I		1		1		1		1		1		1
			400/45+07	1.		- A.		- 1	0.830	in İ		1		Ĩ.		1	0.9700	1	0.9640	1
50.00	1	0.10090E+07 1 U	1.102616+07	:					0.000			- î		1		1	0.9200	1	0.9180	1
60.00	1	0.10090E+07 I L	1.10535E+07	4	1.23	- 2			0.000		20	- i		î î		1	0.8600	1	0.8618	1
70.00	1	0.10090E+07 C	10633E+07	1				- 1-		0 1		1		- î -	- E	1	0.7800	I	0.7887	1
80.00	1	0.10090E+07 C	10492E+07	1		1			0.3/2					i	0	i	0.6800	I.	0.6871	1
90.00	1	0.10090E+07 0	10010E+07	1		1		1	U.265			- 1		- ÷		- î	0.5600	Ť	0.5306	1
100.00	1	0.10090E+07 C	.90058E+06	1		1			0.160			- 1				1	0.9670	i	0.9642	Ĩ
20.00	1	0.42330E+06 0	.44045E+06	1		4		- H.,	0.739	1 Di		1				- 21	0.7070	÷.	0.9270	i
30.00	i.	0.42330E+06 0	.46482E+06	1		1			0.549	50 I						- 1	0.7300		01/270	1
40.00	i.	0.42330E+06 1 0	.47315E+06	1		1		1	0.395	50 I	2	1				1	0.0700		0.0731	
50.00	1	0 42330E+06 0	-46767E+06	L		1		1	0.280	1 00		1				1	0.7800		0.0012	
50.00	τ.	0.423305+06 6	458455+04	÷.		1		- L	0.199	50 I		1		1		ł	0.6800	11	0.7020	
80.00	1		1.4207E+04	i		i i		i	0.130	1 00		1		1		1	0.5700	11	0.5732	
70.00	5	J.42330ETU6 1 4	1.4407UETUO	1		1		÷.	0.04			1		1.		1	0.3800	1	0.3777	1
80.00	1	0.42330E+06 1 L	J.4183UE+U8	1					0.000			- î		1		1	0.1700	1.1	0.1576	, I
90.00	1	0.42330E+06 0	J.40616E+06			1			0.02-			<u>.</u>								

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APPENDIX F

PURE SUBSTANCE PROPERTY TABLES

Table	Fluid
F.1	R11
F.2	R12
F.3	R13
F ₂ .4	R13B1
F.5	R22
F.6	R113
F.7	R152a
F.8	R718
F.9	R500
F.10	Perchloroethylene

Nomenclature

B = Availability D = Density FC = Fugacity coefficient H = Enthalpy P = Pressure PR = Reduced pressure S = Entropy T = Temperature TR = Reduced temperature V = Specific volume Z = Compressibility

(F) = Liquid (G) = Vapour (FG) = Latent value

Note: Availability datum is 24 deg C and 101.325 KPa.

		SATURATED	PROPERTIES	OF TRICHL	OROFLUOROMETHANE	RE	EFRIGERANT 1	1	(CCL3F)	TABLE F1	
т			D(F)	V(G)	 H(F)	H(FG)	н(б)		S(F)	S(G)	т	т
DEG	C F	PA	KG/M3	M3/KG	KJ/KG	KJ/KG	KJ/KG		KJ/KG.K	KJ/KG.K	DEG K	DEG C
			1720.1	2.7132	0.00	199.71	199.71		0.0000	0.8565	233.2	-40.0
-70		3747 7	1701.2	1.5804	7.79	197.10	204.90		0.0353	0.8459	243.2	-30.0
-30	.0 40	770 4	1681.3	0.9669	15.68	194.45	210.13		0.0695	0.8376	253.2	-20.0
-20	.0 29	513.1	1660.6	0.6175	23.67	191.71	215.39		0.1028	0.8313	263.2	-10.0
							200 45		0 4752	0 8267	273.2	0.0
0	.0 39	762.8	1638.8	0.4094	31.77	188.88	220.05		0.1332	0.0207	283 2	10.0
· 10,	.0 59	7827.9	1615.8	0.2804	39.98	185.93	225.92		0.1000	0.0233	203.2	20.0
20	.0 87	7255.8	1591.7	0.1975	48.32	182.84	231.16		0.1978	0.0215	273.2	20.0
30.	.0 123	3777.9	1566.3	0.1427	56.78	179.58	236.36		0.2281	0.0203	303.2	40.0
40	.0 171	313.8	1539.4	0.1053	65.38	176.13	241.51		U.25/9	0.8203	313.4	40.0
	a 374	077 0	4644 4	0.0792	74.13	172.44	246.58		0.2872	0.8208	323.2	50.0
50.	.0 231	1984 3	4684 0	0.0606	83.05	168.50	251.54		0.3161	0.8218	333.2-	60.0
60.	.0 307		4440 4	0.0470	92.14	164.25	256.39		0.3446	0.8232	343.2	70.0
70.	.0 401	512.0	199711	0.0470	101 . 43	159.65	261.09		0.3728	0.8248	353.2	80.0
80	.0 515	5329.5	1413.1	0.0307	110 95	154.66	265.60	2	0.4007	0.8266	363.2	90.0
90,	.0 651	958.3	13/0.0	0.0270	110173	121100						
ក្ម			47/0 0	0 0275	420 70	149.20	269.91		0.4286	0.8284	373.2	100.0
✓ 100.	.0 814	142.0	1340.0	0.0235	430 74	143 22	273.96		0.4563	0.8301	383.2	110.0
110	.0 1004	735.6	1290.2	0.0457	444 40	476 40	277.70		0.4841	0.8316	393.2	120.0
120	.0 1228	5719.0	1253.1	0.0153	454 84	420 27	281.07		0.5121	0.8327	403.2	130.0
130.	.0 1483	5198.1	1204.0	0.0125	151.04	127.25	201.07		0.5404	0.8332	413.2	140.0
140	.0 1777	7404.3	1150.2	0.0102	103.03	120.73	200.77					
		744 5	4090 7	0.0083	174.83	111.51	286.34		0.5693	0.8329	423.2	150.0
150.	.0 2112	(/14.5	4027 8	0.0000	187.36	100.57	287.92		0.5992	0.8314	433.2	160.0
160	.0 2492		1023.0	0.0001	200.94	87.51	288.44		0.6306	0.8280	443.2	170.0
170	.0 2920	19/1.5	740.7	0.0034	214 14	71.12	287.29		0.6647	0.8216	453.2	180.0
180	.0 3401	546.8	034.3	0.0073	210110	48.03	282.72		0.7050	0.8087	463.2	190.0
190	.0 3938	3531.8	/31.7	0.0032	234.07	40100						
191	.0 3999	503.0	716.4	0.0031	236.91	44.98	281.87		0.7097	0.8066	464.2	191.0
192	.0 4052	3082.8	700.0	0.0030	239.24	41.68	280.92		0.7147	0.8043	465.2	192.0
193	0 4111	280-0	682.2	0.0029	241.72	38.08	279.80		0.7199	0.8016	466.2	193.0
194	.0 4170	103.5	662.8	0.0027	244.38	34.08	278.46		0.7256	0.7985	467.2	194.0
195	.0 4229	7551.0	641.2	0.0026	247.30	29.51	276.81		0.7318	0.7948	468.2	195.0
				0 0025	250 40	24.03	276.64		0.7387	0.7900	469.2	196.0
196	.0 . 4289	7630.0	010.2 C 95 7	0.0025	250.00	16.65	271.20		0.7471	0.7825	470.2	197.0
407	-11 4351	14411.11	202.1	0.0023	6 J T B J J							

			SHIC	JANIED FROFER	TIES OF CUES	F FI CONTI	NUEV			nin Sandalaista
T DEG C	TR	Р РА	PR	Z(F)	Z (G)	FC(F)	FC(G)	B(F) KJ/KG	B(G) KJ/KG	T DEG
-40.0	0.4949	5185.8	0.0012	0.0002	0.9970	0.9973	0.9970	0.11E+02	-0.44E+02	-40.0
-30.0	0.5161	9267.7	0.0021	0.0004	0.9952	0.9955	0.9952	0.80E+01	-0.36E+02	-30.0
-20.0	0.5373	15730.4	0.0036	0.0006	0.9926	0,9928	0.9927	0.58E+01	-0.28E+02	-20.0
-10.0	0.5585	25513.1	0.0058	0.0010	0.9891	0.9893	0.9892	0.39E+01	-0.21E+02	-10.0
0.0	0.5798	39762.8	0.0090	.0.0015	0.9845	0.9848	0.9848	0.23E+01	-0.14E+02	0.0
10.0	0.6010	59827.9	0.0136	0.0022	0.9787	0.9791	0.9791	0.11E+01	-0.81E+01	10.0
20.0	0.6222	87255.8	0.0198	0.0031	0.9714	0.9722	0.9721	0.26E+00	-0.22E+01	20.0
30.0	0.6434	123777.9	0.0281	0.0043	0.9626	0.9639	0.9638	-0.29E+00	0.33E+01	30.0
40.0	0.6647	171313.8	0.0389	0.0059	0.9520	0.9541	0.9541	-0.54E+00	- 0.85E+01	40.0
50.0	0.6859	231937.9	0.0526	0.0078	0.9397	0.9429	0.9428	-0.49E+00	0.13E+02	50.0
60.0	0.7071	307881.2	0.0698	0.0103	0.9254	0.9302	0.9301	-0.15E+00	0.18E+02	60.0
70.0	0.7283	401512.0	0.0911	0.0133	0.9090	0.9160	0.9160	0.47E+00	0.22E+02	70.0
80.0	0.7496	515329.5	0.1169	0.0170	0.8904	0.9005	0.9005	0.14E+01	0.27E+02	80.0
90.0	0.7708	651958.3	0.1479	0.0215	0.8695	0.8836	0.8836	0.26E+01	0.31E+02	90°. C
100.0	0.7920	814142.0	0.1846	0.0269	0.8461	0.8655	0.8654	0.41E+01	0.34E+02	100.0
110.0	0.8132	1004735.6	0.2279	0.0334	0.8201	0.8462	0.8461	0.59E+01	0.38E+02	110.0
120.0	0.8345	1226719.0	0.2782	0.0411	0.7913	0.8258	0.8258	0.80E+01	0.41E+02	120.0
130.0	0.8557	1483198.1	0.3364	0.0505	0.7593	0.8044	0.8044	0.10E+02	0.44E+02	130.0
140.0	0.8769	1777404.3	0.4031	0.0618	0.7238	0.7823	0.7822	0.13E+02	0.47E+02	140.0
150.0	0.8781	2112714.5	0.4792	0.0756	0.6842	0.7593	0.7593	0.16E+02	0.50E+02	150.0
160.0	0.9193	2492666.8	0.5653	0.0929	0.6397	0.7358	0.7358	0.20E+02	0.52E+02	160.0
170.0	0.9406	2920971.5	0.6625	0.1150	0.5886	0.7117	0.7117	0.24E+02	0.53E+02	170.0
180.0	0.9618	3401546.8	0.7715	0.1452	0.5278	0.6873	0.6873	0.29E+02	0.54E+02	180.0
190.0	0.9830	3738531.8	0.8933	0.1920	0.4484	0.6625	0.6625	D.36E+02	0.53E+02	190.0
191.0	0.9851	3995503.0	0.9062	0.1985	0.4384	0.6600	0.6600	0.37E+02	0.53E+02	191.0
192.0	0.9873-	4053082.8	0.9192	0.2057	0.4278	0.6575	0.6575	0.38E+02	0.53E+02	192.0
193.0	0.9894	4111280.0	0.9324	0.2136	0.4162	0.6550	0.6550	0.39E+02	0.52E+02	193.0
194.0	0.9915	4170103.5	0,9458	0.2225	0.4035	0.6525	0.6525	0.39E+02	0.52E+02	194.0
195.0	0.9936	4229551.0	0.9593	0.2328	0.3892	0.6500	0.6500	0.41E+02	0.51E+02	195.0
196.0	0.9958	4289630.0	0.9729	0.2451	0.3720	0.6475	0.6475	0.42E+02	0.51E+02	196.0
197.0	0.9979	4350330.0	0.9866	0.2610	0.3478	0.6450	0.6450	0.43E+02	0.49E+02	197.0

SATURATED PROPERTIES OF CCL3F F1 CONTINUED

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	SATD COND		DEGREES OF SUPERHEAT	704	40K
	(DEG C)	- 10K - V(6) 5(6) 5(6)	20K V(6) H(6) S(6)	V(6) H(6) S(6)	V(G) H(G) S(G
	(TR)	M3/KG KJ/KG KJ/KG.K	M3/KG KJ/KG KJ/KG.K	M3/KG KJ/KG KJ/KG.K	M3/KG KJ/KG KJ/K
		B(G) FC(G) Kj/kg	B(G) FC(G) KJ/KG	B(G) FC(G) KJ/KG	"B(6)FC Kj/kg
-		х.			
	(-40.0)	2 870 204 98 0.8812	2.948 210.32 0.9052	3.065 215.74 0.9285	3.182 221.22 0.95
	(0.4949)	-46.1614 0.9973	-47.9420 0.9976	-49.4573 0.9978	-50.7219 0.9980
	(-70 0)	1997 - 1997 - 1998 - 1998 - 1998 - 1998 - 1998 - 1998 - 1998 - 1998 - 1998 - 1998 - 1998 - 1998 - 1998 - 1998 -			2
	(9.268)	1.646 210.25 0.8699	1.712 215.67 0.8932	1.778 221.16 0.9159	
	(0.5161)	-37.5209 0.9957	-39.0368 0.9961	-40.3017 0.9984	-41.3203 0.7700
	(-20.0)	- a b			4 4 9 9 9 7 9 7 1 8 9
	(15.730)	1.006 215.55 0.8609	1.045 221.05 0.8837	1.083 226.61 U.9059 -31.8488 0.9945	-32.6492 0.9950
	(0.5373)	-29.5562 0.9934	-30.8217 0.7740	-31.8466, 667775	7
	(-10.0)		0 /// 00/ // 0 87/7	n 4en 232.10 0.8979	0.714 237.80 0.91
0	(25.513)	0.642 220.89 0.8540 -22.1777 0.9902		-24.0060 0.9918	-24.5905 0.9925
ω	(0.55057		201		
	(0.0)	D 425 224 24 D 8489	R.440 231.88 0.8706	0.456 237.59 0.8918	0.472 243.37 0.91
120	(0.5798)	-15.3037 0.9861	-16.1048 0.9873	-16.6892 0.9884	-17.0671 0.9893
	(10.0)		÷ .		
	(59.828)	· 0.291 231.58 0.8452	0.301 237.31 0.8665	0.312 243.09 0.8872	
	(0.6010)	-8.8697 0.9809	-9.4541 0.9825	-9.8315 0.7637	
	(20.0)				0.226 254.50 0.90
	(87.256)	0.205 236.90 0.8428	0.212 242.71 0.8636 -3 1987 0.9766	-3.3773 0.9784	-3.3662 0.9801
	(0.6222)	-2.8220 0.9745	-3.1787 017100		
	(30.0)		0 453 948 08 0 8418	0.158 254.03 0.8818	0.163 260.03 0.90
	(123.778) (0.6434)	U:145 242.17 U.0414 2.8825 0.9668	2.7055 0.9694	2.7187 0.9718	2.9140 0.9739
	(40.0)	- - - 1.109 247.43 0.8409	0.113 253.41 0.8609	0.117 259.44 0.8805	0.120 265.52 0.89
	(0.6647)	8.2802 0.9577	8.2963 0.9610	8.4951 0.9640	8.8687 U.766/
	(50.0)				
	(231.938)	0.082 252.60 0.8410	0.085 258.66 0.8608	0.088 264.78 0.8801	0.091 270.95 0.85 14.5293 0.9582

	er har		0							
T Deg c	 Р РА	D(F) Kg/M3	V(G) M3/KG	H(F) KJ/KG	H(FG) KJ/KG	H(G) KJ/KG	S(F) KJ/KG.K	S(G) KJ/KG.K	T DEG K	T DEG C
-40.0	45562 A	1613.2	0.2386	0.00		167.91	0.0000	0.7202	233.2	-40.0
-30.0	101847.4	1582.9	0.1586	8.43	164.46	172.90	0.0382	0.7146	243.2	-30.0
-20.0	152257.6	1550-6	0.1090	17.06	160.76	177.82	0.0756	0.7106	253.2	-20.0
-10.0	220172.2	1516.0	0.0772	25.89	156.76	182.65	0.1123	0.7080	263.2	-10.0
0.0	309250.8	1479.0	0.0559	. 34.94	152.40	187.35	0.1484	0.7064	273.2	0.0
10.0	423426.9	1439.2	0.0414	44.25	147.63	191.88	0.1841	0.7054	283.2	10.0
20.0	566877.7	1396.2	0.0311	53.85	142.35	196.20	0.2194	0.7050	293.2	20.0
30.0	744013.8	1349.6	0.0238	63.77	136.49	200.25	0.2546	0.7048	303.2	30.0
40.0	959459.0	1298.8	0.0183	74.06	129.92	203.98	0.2897	0.7046	313.2	40.0
50.0	1218071.5	1243.0	0.0142	84.81	122.49	207.31	0.3250	0.7040	323.2	50.0
60.0	1524937.6	1181.2	0.0111	96.11	114.00	210.11	0.3607	0.7029	333.2	60.0
70.0	1885397.3	1111.7	0.0087	108.11	104.14	212.24	0.3973	0.7008	343.2	70.0
80.0	2305077.8	1032.1	0.0068	121.04	92.40	213.44	0.4353	0.6970	353.2	80.0
90.0	2789927.0	938.1	0.0052	135.37	77.89	213.26	0.4759	0.6903	363.2	90.0
100.0	3346270.5	819.3	0.0039	152.13	58.48	210.60	0.5215	0.6782	373.2	100.0
110.0	3980889.0	629.8	0.0025	176.12	24.13	200.25	0.5844	0.6474	383.2	110.0
111.0	4048918.3	595.4	0.0023	180.14	16.85	196.99	0.5948	0.6386	384.2	111.0
			SATURAT	ED PROPERTIES	OF CCL2F2	CONTINU	IED			
	TR	P PA	PR	Z(F) 2	2(6)	FC(F)	FC(G)	B(F) Kj/kg	B(G) KJ/KG	T Deg C
-40.0	0.6054	65562.8	0.0159	0.0025 0.	.9757	0.9763	0.9762	0.47E+02	0.77E+00	-40.0
-30.0	0.6313	101843.4	0.0247	0.0038 0.	9658	0.9669	0.9669	0.44E+02	0.74E+01	-30.0
-20.0	0.6573	152257.6	0.0370	0.0056 0.	.9536	0.9555	0.9555	0.41E+02	0.14E+02	-20.0
-10.0	0.6832	220172.2	0.0535	0.0080 0.	9386	0.9419	0.9419	0.39E+02	0.19E+02	-10.0
0.0	0.7092	309250.8	0.0751	0.0111 0.	9208	0.9262	0.9262	0.38E+02	0.24E+02	u.u
10.0	0.7352	423426.9	0.1029	0.0151 0.	.8999	0.9084	0.9084	0.36E+02	0.29E+02	10.0
20.0	0.7611	566877.7	0.1377	0.0201 0.	.8757	0.8886	0.8886	0.36E+02	0.34E+02	20.0
30.0	0.7871	744013.8	0.1808	0.0264 0.	8481	0.8669	0.8669	0.35E+02	0.38E+02	30.0
40.0	0.8131	959459.0	0.2331	0.0343 0.	.8166	0.8436	0.8435	0.35E+02	0.41E+02	40.0
50.0	0.8390	1218071.5	0.2960	0.0441 0.	7809	0.8187	0.8187	0.35E+02	0.45E+02	50.0
60.0	0.8650	1524937.6	0.3705	0.0563 0.	7405	0.7925	0.7925	0.36E+02	0.48E+02	60.0
70.0	0.8710	1885397.3	0.4581	0.0719 0.	6947	0.7652	0.7652	0.37E+02	0.51E+02	70.0
80.0	0.9169	2305077.8	0.5601	0.0919 0.	.6419	0.7369	0.7369	0.39E+02	0.53E+02	80.0
90.0	0.9429	2789927.0	0.6779	0.1191 0.	5799	0.7080	0.7080	0.41E+02	0.55E+02	90.0
100.0	0.9688	3346270.5	0.8131	0.1591 0.	5025	0.6784	0.6784	0.44E+02	0.56E+02	100.0
110.0	0.9948	3980889.0	0.9673	0.2399 0.	3793	0.6485	0.6485	0.49E+02	0.55E+02	110.0
111.0	0.9974	4048918.3	0.9838	0.2574 0.	.3538	0.6455	0.6455	0.50E+02	0.54E+02	111.0

SATD COND		DE	SREES OF SUPERHEAT	20K		40	к
(DEG C) (KPA) (TR)	10K V(G) H(G) S M3/KG KJ/KG KJ	(5) V(6) /KG.K M3/KG	20K H(G) S(G) KJ/KG KJ/KG.K	V(G) H(G) M3/KG KJ/KG	2(G) 8(G)	V(G) H(M3/KG KJ	G) S(G) /Kg KJ/Kg.K
	8(G) FC(KJ/KG	G) B(KJ/K	G) FC(G) G	B(G) KJ/KG	FC(6)	B(G) Kj/kg	FC(G)
(-40.0) (65.563) (0.6054)	0.250 173.43 0. -1.4573 0.978	7462 0.260 7 -3.3	179.04 0.7716 893 0.9808	0.271 184.74 -5.0422	0.7964 0.9826	0.282 190 -6.4303	.52 0.8205 0.9842
(-30.0) (101.843) (0.6313)	0.166 178.54 0. 5.4774 0.970	7401 0.173 2 3.8	184.26 0.7649 201 _0.9730	0.180 190.07 2.4292	0.7892 0.9756	0.187 195 1.2917	.96 0.8128 0.9778
(-20.0) (152.258) (0.6573)	0.114 183.59 0. 11.8450 0.959	7356 0.119 7 10.4	187.44 0.7600 500 0.9635	0.124 195.36 9.3100	0.7838 0.9668	0.128 201 8.4129	.36 0.8070 0.9698
(-10.0) (220.172) N (0.6832)	0.081 188.55 0. 17.7194 0.947	7326 0.084 3 16.5	194.53 0.7565 760 0.9521	0.087 200.58 15.6774	0.7799 0.9564	0.091 206 15.0120	.70 0.8027 0.9602
(0.0) (307.251) (0.7072)	0.058 193.40 0. 23.1585 0.932	7306 0.061 9 22.2	199.51 0.7542 577 0.9389	0.063 205.69 21.5926	0.7772 0.9442	0.066 211 21.1519	.94 0.7997 0.9489
(10.0) (423.427) - (0.7352)	.0.043 198.09 0. 28.2067 0.916	7294 0.045 5 27.5	204.36 0.7527 418 0.9238	0.047 210.68 27.1039	0.7755 0.9303	0.049 217 26.8824	.07 0.7977 0.9361
(20.0) (566.878) (0.7611)	0.033 202.60 0. 32.8974 0.898	7288 0.03 4 3 32.4	209.04 0.7519 636 0.9070	0.036 215.52 32.2486	0.7744 0.9147	0.037 222 32.2424	.05 0.7964 0.9217
(30.0) (744.014) (0.7871)	0.025 206.87 0. 37.2537 0.878	7285 0.026 3 37.0	213.50 0.7515 486 0.8885	0.027 220.16 37.0547	0.7739 0.8976	0.028 226 37.2618	.86 0.7957 0.9058
(40.0) (959.459) (0.8131)	0.019 210.85 0. 41.2891 0.856	7283 0.020 7 41.3	217.71 0.7513 139 0.8685	0.021 224.58 41.5416	0.7737 0.8791	0.022 231 41.9624	.46 0.7954 0.8886
(50.0) (1218.072) (0.8390)	0.015 214.50 0. 45.0087 0.833	7281 0.016 57 45.2	221.63 0.7512 685 0.8472	0.017 228.74 45.7219	0.7736 0.8593	0.018 235 46.3597	.83 0.7953 0.8703

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	201	UNRIEV PROIE								
т	 P	D(F)	V(6)	HC	F) H(FG	H(G)	S(F)	S(6) KI/KG K	T DEG K	T DEG C
DEG C	PA	KG/M	3 M3/KG	KJ.	/KG KJ/K	5 KJ/NG				
-40.0	LOU268'3	1425.7	7 0.0267	0	.00 126.6	126.68	0.0000	0.5433	233.2	-40.0
-70.0	878495 7	1361.5	0.0191	10	.19 119.4	129.66	0.0454	0.5367	243.2	-30.0
-30.0	4470827 7	1288.4	6 0.0138	21	.11 111.0	132.18	0.0917	0.5304	253.2	-20.0
-10.0	1514586.9	1205.9	9 0.0101	32	.91 101.1	134.01	0.1393	0.5235	263.2	-10.0
					a. 99 Gi	. 174 81	0.1891	0.5148	273.2	0.0
0.0	1973200.3	1109.4	6 0.0074	45 (0		433.97	0.2429	0.5021	283.2	10.0
10.0	2526816.3	992.4	6 0.0053	60	.55 /3.4	400.85	0 3054	0.4805	293.2	20.0
20.0	3187448.8	837.9	5 0.0036	78	.52 51.3		0 3427	0.4774	294.2	21.0
21.0	3259890.5	818.2	2 0.0035	80	.66 48.4	129.09	0.3127	0 4738	295.2	22.0
22.0	3333546.0	797.	7 0.0033	82	.90 45.3	128.22	0.3203	0.4/00	*	
			7 0 0032	85	.27 41.9	127.22	0.3282	0.4699	296.2	23.0
23.0	3408428.0		0.0000	· 87	.80 38.2	126.05	0.3367	0.4654	297.2	24.0
24.0	3484554.3	(52.)		01	54 34.1	124.66	0.3458	0.4603	298.2	25.0
25.0	3561933.3	/26.0		07	67 29.3	122.93	0.3559	0.4540	299.2	26.0
26.0	3640586.8	696.		73	n4 23.6	120.66	·· 0.3674	0.4460	300.2	27.0
27.0	3720532.3	662.4	• U.UU25	77	.08 23.0		105		- C	
		34		101	79 15.7	117.11	0.3816	0.4338	301.2	28.0
28.0	3801752.3	619.0		404	04 14.6	116.57	0.3833	0.4320	301.3	28.1
28.1	3809946.3	613.0	5 0.0022	101	·71 17·0	115.95	0.3851	0.4299	301.4	28.2
28.2	3818149.5	608.3	3 0.0022	102		115.16	0.3869	0.4273	301.5	28.3
28.3	3826360.5	602.6	5 0.0021	103	.01 12.1	,			20	
			SATURAI	ED PROPERT	IES OF CCLF3	CONTINU	JED			
				7(E)	7(6)	FC(F)	FC(G)	B(F)	B(G)	т
	I K	PA	T K					KJ/KG	KJ/KG	DEG C
								D 845+02	0 528+02	-40.0
-40.0	0.7721	600798.2	0.1553	0.0227	0.8642	0.8794	0.0/74	0.000-02	0.575+02	-30 0
-30.0	0.8052	838495.7	0.2168	0.0318	0.8262	0.8505	0.8505	0.035+02	0.572.02	-20.0
-20.0	0.8383	1139827.3	0.2947	0.0439	0.7816	0.8191	0.8191	0.802402	0.010+02	-10.0
-10.0	0.8714	1514586.9	0.3916	0.0600	0.7294	0.7856	0.7856	U.76ETU2	0.035402	0.0
0.0	0.9045	1973200.3	0.5101	0.0818	0.6677	0.7503	0.7503	U./8E+U2	0.002702	0.0
10 0	0 9376	2526816-3	0.6533	0.1130	0,5931	0.7137	0.7137	0.75E+02	0.71E+02	10.0
20.0	0.0700	2497669 9	D.8241	0.1631	0.4954	0.6762	0.6762	0.74E+02	0.73E+02	20.0
20.0	0.97/0	310/770.0°	0.0271	0.4702	0.4835	0.6724	0.6724	0.74E+02	0.74E+02	21.0
21.0	0.9741	3237070.3	0.0720	0 1779	0.4707	0.6686	0.6686	0.74E+02 *	0.74E+02	22.0
22.0	0.9774	2323340.0	0.0010	0 4844	0 4574	0.6668	0.6648	0.74E+02	0.74E+02	23.0
23.0	0.9807	3408428.8	U.0012	0.1004	014371	0.0040	010010	.		
24.0	0.9840	3484554.3	0.9009	0.1959	0.4424	0.6610	0.6610	0.74E+02	0.74E+02	24.0
25.0	0.9873	3561933.3	0.9209	0.2068	0.4262	0.6571	0.6571	0.74E+02	0./4E+U2	25.0
26.0	0.9906	3640586.8	0.9412	0.2195	0.4078	0.6533	0.6533	0.74E+02	0.74E+02	26.0
27.0	0.9939	3720532.3	0.9619	0.2351	0.3860	0.6495	0.6495	0.74E+02	0.75E+02	27.0
28.0	0.9973	3801752.3	0.9829	0.2562	0.3557	0.6457	0.6457	0.74E+02	0.75E+02	28.0
2010									0.755.00	29.4
28.1	0.9976	3809946.3	0.9850	0.2589	0.3514	0.6453	0.6453	U.74E+02	U.75E+UZ	20.1
28.2	0.9979	3818149.5	0.9871	0.2617	0.3465	0.6449	0.6449	U.74E+U2	U.75E+UZ	20.2
28.3	D.9982	3826360.5	0.9892	0.2647	0.3406	0.6445	0.6445	0.74E+02	0.75E+02	20.3

			SU	PERHEATED P	ROPERTIES	OF CHLOR	OTRIFLUOROME	THANE	REFRIGE	RANT R13	(CCLF3	'	-3
	SATD COND (DEG C) (KPA) (TR)	V(6) M3/KG	10K H(g) KJ/Kg	S(6) Kj/kg.K	DE V(G) M3/KG	GREES OF 20K H(G) KJ/KG	SUPERHEAT S(6) KJ/KG.K	V(G) M3/KG	30K H(G) KJ/KG	S(G). Kj/kg.k	V(G) M3/KG	40K H(g) KJ/Kg	S(G) Kj/kg.k
		B(6 Kj/K) G	FC(6)	В(<u></u> КЈ/К	6) G	FC(6)	KJ/I	B(G) Kg	FC(6)	КЈ/	B(G) Kg 	FC(G)
	(-40.0) (600.798) (0.7721)	0.028	132.68 300 0	0.5719 .8925	0.030 46.9	138.83 628 0	0.5998 .9038	0.032	145.11 0965 0	0.6273 .9137	0.033 43 <i>,</i>	· 151.51 5218 (0.6541].9223
	(-30.0) (838.496) (0.8052)	0.020 54.3	136.10 205 0	0.5659 .8664	0.022 52.4	142.62 178 O	0.5943 .8802	0.023 50.8	149.23 8214 Q	0.6218 .8922	0.024 49,	155.90 5145 (0.6487 3.9028
	(-20.0) (1139.827) (0.8383)	0.015 58.90	139.14 681 0	0.5605 .8380	0.016	146.08 344 D	0.5893 .8544	0.017 56.0	153.04 0084 0	0.6172 .8687	0.018 54.	160.02 9685 C	0.6442).8814
203	(-10.0) (1514.587) (0.8714)	0.011 63.13	141.64 182 O	0.5549 .8077	0.012	149.10 575 0	0.5846 .8268	0.013	156.46 7052 0	0.6129 .8436	0.013	163.79 9355 (0.6402 .8585
	(0.0) (1973.200) (0.9045)	0.008 66.79	143.43 705 0	0.5486 .7759	0.009 65.7	151.55 136 0	0.5796 .7980	0.010	159.43 7452 0	0.6087 .8174	0.010 64.	167.17 4528 (0.6364 .8344
	(10.0) (2526.816) (0.9376)	_ 0.006 69.98	144.27 309 0	0.5407 .7433	0.007	153.34 149 D	0.5738 .7685	0.007 68.3	161.88 7499 0.	0.6041 .7904	0.008 68.	170.12 5495 (0.6325).8098
	(20.0) (3187.449) (0.9708)	0.005 72.68	143.93 544 0	0.5305	0.005	154.39 649 D	0.5671 7388	0.006 72.1	163.77 1383 0.	0.5991 .7634	0.006 72.	172.64 2533 (0.6285).7851
	(30.0) (3764.322) (1.0039)	0.003 74.80	142.45)53 O	0.5183 .6787	0.004 74.8	154.82 693 0.	0.5597 .7103	0.005 75.1	165.23 1295 D	0.5939 7374	0.005 75.	174.81 5890 C	0.6246).7611
	(40.0) (4855.447) (1.0370)	0.003	141.18 761 0	0.5084 .6507	0.003	155.23 915 0.	0.5536 .6847	0.004 77.7	166.65 7723 0	0.5898 7138	0.004 78.	176.95 5998 C	0.6217).7394

	SATURA	TED PROPERTIE	S OF BROMOTR	IFLUOROMETHANE	REF	RIGERANT 13	381 (CBRF3)	TABLE F4	1
T DEG C	 Р РА	D(F) Kg/M3	V(G) M3/KG	H(F) KJ/KG	H(FG) Kj/kg	" H(6) KJ/KG	S(F) Kj/kg.k	S(G) KJ/KG.K	T DEG K	T DEG
-40.0	219762.3	2032.0	0.0555	0.00	111.84	111.84	0.0000	0.4797	233.2	-40.0
-30.0	a 321124.3	1975.1	0.0387	6.85	108.26	115.11	0.0309	0.4761	243.2	-30.0
-20.0	454584.3	1913.2	0.0277	13.95	104.29	118.24	0.0615	0.4734	253.2	-20.0
-10.0	626054.9	1845.6	0.0202	21.33	99.83	121.16	0.0919	0.4712	263.2	-10.0
0.0	841800.0	1771.3	0.0150	29.04	94.78	123.82	0.1223	0.4693	273.2	0.0
10.0	1108436.6	1689.0	0.0113	37.15	89.00	126.15	0.1529	0.4672	283.2	10.0
20.0	1432937.5	1596.8	0.0085	45.74	82.30	128.04	0.1839	0.4647	293.2	20.0
30.0	1822662.9	1491.9	0.0064	54.96	74.38	129.34	0.2159	0.4612	303.2	30.0
40.0	2285418.0	1369.5	0.0048	65.06	64.73	129.79	0.2494	0.4561	313.2	40.0
50.0	2829508.8	1220.2	0.0036	76.54	52.33	128.87	0.2859	0.4479	323.2	50.0
60.0 8	3463855.3	1017.4	0.0025	90.87	34.20	125.07	0.3295	0.4322	333.2	60.0
61.0	3532615.3	991.1	0.0024	92.63	31.71	124.34	0.3348	0.4297	334.2	61.0
62.0	3602386.8	962.7	0.0023	94.51	28.99	123.50	0.3404	0.4269	335.2	62.0
63.0	3673178.8	931.6	0.0022	96.53	25.97	122.50	0.3463	0.4236	336.2	63.0
64.0	3744999.8	896.9	0.0020	98.76	22.51	121.27	0.3529	0.4196	337.2	64.0
65.0	3817859.5	856.6	0.0017	101.29	18.37	119.66	0.3603	0.4146	338.2	65.0
66.0	3891765.8	806.7	0.0017	104.37	12.87	117.24	0.3693	0.4072	337.2	66.0
66.1	3899207.8	800.8	0.0017	104.72	12.17	116.89	0.3703	0.4062	339.3	66.1
66.2	3906669.0	794.8	0.0017	105.09	11.41	116.50	0.3714	0.4050	339.4	66.2
66.3	3914131.3	788.4	0.0017	105.47	10.58	116.06	0.3725	0.4037	339.5	66.3
66 6	3921600.8	781.8	0.0016	105.87	9.64	115.51	0.3737	0.4021	339.6	66 . 4
00.4	7000075 0	774.7	0.0016	106.30	8.45	114.75	0.3749	0.3998	339.7	66.5

T DEG C	TR	 Р РА	PR	Z(F)	Z(G)	FC(F)	FC(G)	B(F) Kj/kg	B(G) KJ/KG	T DEG (
-40.0	0.6854	219762.3	0.0554	0.0083	0.9369	0.9404	0.9404	0.51E+02	0.20E+02	-40.0
-30.0	0.7148	321124.3	0.0810	0.0120	0.9162	0.9222	0.9222	0.49E+02	0.24E+02	-30.0
-20.0	0.7442	454584.3	0.1147	0.0168	0.8915	0.9014	0.9014	0.47E+02	0.28E+02	-20.0
-10.0	0.7736	626054.9	0.1579	0.0231	D.8625	0.8781	0.8780	0.45E+02	0.32E+02	-10.0
0.0	0.8030	841800.0	0.2123	. 0.0312	0.8288	0.8525	0.8525	0.44E+02	0.35E+02	0.0
10.0	0.8324	1108436.6	0.2796	0.0415	0.7900	0.8249	0.8248	0.43E+02	0.38E+02	10.0
20.0	0.8618	1432937.5	0.3614	0.0548	0.7453	0.7955	0.7955	0.42E+02	0.41E+02	20.0
30.0	0.8712	1822662.9	0.4597	0.0722	0.6938	0.7647	0.7647	0.42E+02	0.43E+02	30.0
40.0	0.9206	2285418.0	0.5765	0.0754	0.6334	0.7327	0.7327	0.42E+02	0.45E+02	40.0
50.0	0.9500	2829508.8	0.7137	0.1285	0.5603	0.6778	0.6998	0.42E+02	0.47E+02	50.0
40.0	0,9794	3463855.3	0.8737	0.1830	0.4625	0.6662	0.6662	0.44E+02	0.47E+02	60.0
61.0	0.9824	3532615.3	0.8911	0.1911	0.4498	0.6629	0.6628	0.44E+02	0.47E+02	61.0
62.0	0.9853	3602386.8	0.9087	0.2000	0.4362	0.6595	0.6595	0.44E+02	0.47E+02	62.0
63.0	0.9882	3673178.8	0.9265	0.2101	0.4213	0.6561	0.6561	0.44E+02	0.47E+02	63.0
64.0	0.9912	3744999.8	0.9446	0.2218	0.4045	0.6527	0.6527	0.45E+02	0.47E+02	64.0
65.0	0,9941	3817859.5	0.9630	0.2361	0.3846	0.6493	0.6493	0.45E+02	0.47E+02	65.0
66.0	0.9971	3891765.8	0.9817	0.2548	0.3579	0.6459	0.6459	0.45E+02	0.47E+02	66.0
66.1	0.9974	3899207.8	0.9835	0.2570	0.3544	0.6456	0.6456	0.46E+02	0.47E+02	66.1
66.2	0.9976	3906669.0	0.9854	0.2594	0.3505	0.6452	0.6452	0.46E+02	0.47E+02	66.2
66.3	0.9979	3914131.3	0.9873	0.2619	0.3461	0.6449	0.6449	0.46E+02	0.47E+02	66.3
66.4	0.9982	3921600.8	0.9892	0.2646	0.3408	0.6445	0.6445	0.46E+02	0.47E+02	66.4
66.5	0.9985	3929035.0	0.9911	0.2674	0.3337	0.6442	0.6442	0.46E+02	0.47E+02	66.5

SATURATED PROPERTIES OF CBRF3 F4 CONTINUED

	SATD COND	4.0%	× D	EGREES OF SUPERHEAT	30K			40K	
	(KPA) (TR)	V(G) H(G) S M3/KG KJ/KG KJ	(6) V(6) /KG.K M3/KG	H(G) S(G) KJ/KG KJ/KG.K	V(G) H(G) M3/KG KJ/K	S(G) G KJ/KG.K	V(6) M3/KG	H(G) KJ/KG	S(G) KJ/KG.K
		B(G) FC(Kj/kg	G) B KJ/	(G) FC(G) Kg	B(G) KJ/KG	FC(G)	KJ/K	B(G) G	FC(6)
	(-40.0)				L				
	(219.762) (0.6854)	0.058 116.13 0. 18.3839 0.946	5000 0.061 6 16.	120.49 0.5199 8530 0.9521	0.064 124.93 15.5395	3 0.5392 0.9568	14.4	129.44 324 0.9	610
	(-30.0) (321.124) (0.7148)	0.041 119.55 0. 22.9224 0.930	4963 0.043 1 21.	124.05 0.5159 5994 0.9371	0.045 128.6 20.4863	2 0.5350 0.9431	0.047 19.53	133.25 728 0.9	0.5537 7485
à)	(-20.0) (454.584) (0.7442)	0.029 122.84 0. 27.0577 0.911	6934 0.031 2 25.	127.49 0.5128 9354 0.9198	0.032 132.20 25.0169	0 0.5318 0.9273	0.033	136.97 718 0.9	0.5503 /340
	(-10.0) (626.055) (0.7736)	0.021 125.95 0.4 30.8331 0.889	912 0.022 29.0 29.0	130.77 0.5105 9067 0.9003	0.024 135.64 29.1785	6 0.5293 0.9095	0.025 28.63	140.55 882 0.9	0.5477
206	(0.0) (841.800) (0.8030)	0.016 128.83 0.4 34.2800 0.866	893 0.017 5 33.9	133.85 0.5086 5469 0.8789	0.018 138.90 33.0070) 0.5274 0.8898	0.018 32.64	143.98 95 0.8	0.5457 996
	(10.0) (1108.437) (0.8324)	0.012 131.44 0.4 37.4190 0.8412	875 0.013 2 36.0	136.70 0.5070 . 3795 0.8557	0.013 141.99 36.5282	0.5259 0.8685	0.014 36.39	147.22 640 0.8	0.5442
	(20.0) (1432.938) (0.8618)	0.009 133.70 0.4 40.2593 0.8144	855 0.010 39.0	139.25 0.5054 7185 0.8310	0.010 144.78 39.7600	0.5244 0.8458	0.011 39.77	150.24 22 0.8	0.5429 591
	(30.0) (1822.663) (0.891 <u>2</u>)	0.007 135.51 0.4 42.7999 0.7863	- 631 0.008 6 42.0	141.46 0.5035 6697 0.8053	0.008 147.27 42.7129	0.5230 0.8221	0.009 42.91	153.02 86 0.8	0.5416
	(40.0) (2285.418) (0.9206)	0.005 136.78 0.4 45.0268 0.7572	798 0.006 45.1	143.25 0.5013 1312 0.7787	0.006 149.40 45.3926	0.5213 0.7977	0.007	155.52)36 0.8	0.5403
	(50.0) (2829.509) (0.9500)	0.004 137.33 0.4 46.9125 0.7277	754 0.005 47.2	144.56 0.4984 2959 0.7518	0.005 151.28 47.8023	0.5193 0.7729	0.005	157.73 67 0.7	0.5389 917
	(60.0) (3463.855) (0.9794)	0.003 137.04 0.4 48.4200 0.4983	693 0.00 4	145.37 0.4949 597 0.7250	0.004 152.74	0.5170	0.004	159.65 11 0.7	0.5373 690

	SATURATED	PROPERTIES	OF CHLORO	DIFLUOROMETHANE	REF	RIGERANT 22	(CHCLF2)	TABLE HD	
т	P	D(F)	V(G)	 H(F)	 H(FG)	H(G)	S(F)	5(6)	Т	Ť
DEG C	PA	KG/H3	M3/KG	KJ/KG	KJ/KG	KJ/KG	KJ/KG.K	KJ/KG.K	DEG K	DEG C
-40.0	105882.9	1439.6	0.2054	0.00	232.54	232.54	.0000	0.9973	233.2	-40.0
-30.0	164249.9	1409.4	0.1364	10.45	226.97	237.43	0.0478	0.9812	243.2	-30.0
-20.0	245194.5	1377.0	0.0936	21.20	220.96	242.16	0.0947	0.9675	253.2	-20.0
-10.0	354029.5	1342.0	D.0661	32.28	214.41	246.69	0.1410	0.9558	263.2	-10.0
° 0.0	191519.7	1304.1	0.0478	43.73	207.22	250.95	0.1870	0.9456	273.2	0.0
40.0	478972 4	1263.0	0.0352	55.62	199.26	254.88	0.2326	0.9363	283.2	10.0
20.0	07077211	1218.1	0.0263	. 68.01	190.36	258.37	0.2783	0.9277	293.2	20.0
20.0	4400425 8	1168.8	0.0199	81.00	180.34	261.33	0.3243	0.9192	303.2	30.0
40.0	1533914.5	1114.1	0.0152	94.71	168.92	263.62	0.3710	0.9104	313.2	40.0
50.0	4044247 4	1052.7	0.0117	109.33	155.71	265.04	0.4187	0.9006	323.2	50.0
50.0	1770273+1	087 8	0.0089	125.15	140.14	265.29	0.4684	0.8891	333.2	60.0
20.0	2933730.0	901.0	0.0068	142.66	121.17	263.85	0.5212	0.8744	343.2	70.0
70.0	7480004 8	800.6	0.0050	162.91	96.75	259.66	0.5799	0.8538	353.2	80.0
90.0	4458872.0	661.4	0.0035	187.12	60.25	249.37	0.6527	0.8186	363.2	90.0
	15/07/5 0	442 7	0.0033	192.49	55.08	247.57	0.6619	0.8131	364.2	91.0
91.0	4542/05.U //0787/ 5	472 3	0.0032	196.13	49.32	245.45	0.6718	0.8068	365.2	92.0
72.0	402/030.3	C00 C	0.0051	200.16	42.74	242.90	0.6827	0.7994	366.2	93.0
93.0	4/14100.0	377.3 E77 4	0.0028	204.77	34.87	239.64	0.6951	0.7901	367.2	94 - 0
94.0	4001501.0	5/3.1	0.0026	210.39	24.41	234.80	0.7102	0.7765	368.2	95.0
75.U	4870231.0	7.046	9.0020							
95 4	180044 0 5	536.6	0.0026	211.04	23.07	234.11	0.7120	0.7746	368.3	95.1
95.2	4908110.0	532.6	0.0025	211.72	21.62	233.34	0.7138	0.7725	368.4	95.2

				7(5)	7(5)		FC(6)	B(F)	B(G)	T
Т	TR	P	РК	2177	2(0)	1 9 11 /		KJ/KG	KJ/KG	DEG
DEG C		PA								
		405992 0	0.0213	0.0033	0.9700	0.9708	0.9708	0.77E+02	0.13E+02	-40.0
-40.0	0.6316	103002.7	0.0210	0.0050	0.9579	0.9595	0.9595	0.73E+02	0.23E+02	-30.1
-30.0	0.6587	104247.7	0.0330	0 0073	0.9429	0.9458	0.9458	0.70E+02	0.32E+02	-20.6
-20.0	0.6858	245174.5	0.0711	0.0104	0.9247	0.9296	0.9296	0.67E+02	0.40E+02	-10.0
-10.0	0.7129	354027.5	0.0711	0.0145	0 9030	0.9110	0.9110	0.65E+02	0.47E+02	0.0
0.0	0.7400	496549.7	0.0770	. 0.0145	01/000	•••••				
			0 4744	0.0497	0 8774	°° n. 8900	0.8900	0.63E+02	0.53E+02	10.1
10.0	0.7670	678972.1	U.1304	0.0177	0.0//0	0 8669	0.8669	0.62E+02	0.60E+02	20.1
20.0	0.7941	907921.8	0.1824	0.0204	0.0401	n #448	0.8418	0.61E+02	0.65E+02	30./
30.0	0.8212	1190425.8	0.2392	0.0347	0.0141	0.0410	0.8149	0.61E+02	0.70E+02	40./
40.0	0.8483	1533914.5	0.3082	0.0457	0.7753	0.0177	0.0111	0.62E+02	0.74E+02	50.0
50.0	0.8754	1946243.1	0.3910	0.0595	0.7308	U. (003	0.1005			
2			1			0 75/7	0 7547	0.63E+02	0.78E+02	60./
60.0	0.9025	2435736.8	0.4894	0.0774	0.6/95	0.7507	0.7259	0.655+02	0.81E+02	70.
70.0	0,9296	3011242.0	0.6050	0.1013	0.6195	U. /259	0.1237	0.050.02	0.83E+02	80.
80.0	0.9567	3682226.8	0.7398	0.1354	0.5465	0.6943	0 4420	0 725+02	0.83E+02	90.
90.0	0.9837	4458872.0	0.8958	0.1931	0.4467	0.6621	0.0020	0 775+02	0.83E+02	91.
91.0	0.9865	4542765.0	0.9127	0.2019	0.4334	0.6588	0.000	0.132702	01002.02	
				× 2	0 ((00		0 4555	0.73E+02	0.83E+02	92.
92.0	0.9892	4627836.5	0.9298	0.2118	U.4100		0.6533	0.748+02	0.82E+02	93.
93.0	0.9919	4714100.0	0.9471	0.2233	0.4024	0.6523	0.4323	0 755+02	0.82E+02	94.
94.0	0.9946	4801561.0	0.9647	0.2373	0.3829	0.6490	0.0970	0 745+02	0.81E+02	95.
95.0	0.9973	4890231.0	0.9825	0.2556	0.3566	U.6458	0.0930	0.746+02	D.81E+02	95.
95.1	0.9976	4899160.5	0.9843	0.2579	0.3531	0.6454	0.0934	0.702.02	AINIT . AT	
95.2	0.9978	4908110.0	0.9861	0.2602	0.3492	0.6451	0.6451	0.76E+02	0.81E+02	95.

SATURATED PROPERTIES OF CHCLF2 F5 CONTINUED

DEGREES OF SUPERHEAT SATD COND 10K 20K 30K H(G) S(G) V(G) H(G) S(G) V(G) H(G) S(G) V(G) 40K (DEG C) H(G) S(6) V(G) (KPA) M3/KG KJ/KG KJ/KG.K M3/KG KJ/KG KJ/KG.K M3/KG KJ/KG KJ/KG.K M3/KG KJ/KG KJ/KG.K (TR) B(G) FC(6) B(G) FC(G) B(G) FC(G) FC(6) B(6) KJ/KG KJ/KG KJ/KG KJ/KG (-40.0)0.243 256.58 1.1076 0.234 250.40 1.0810 0.215 238.38 1.0259 0.224 244.33 1.0538 (105.883) 4.2928 0.9809 6.0273 0.9788 8.0512 0.9765 10.3787 0.9738 (0.6316) (-30.0)0.155 255.79 1.0636 0.161 262.13 1.0898 0.149 249.56 1.0367 (164.250) 0.143 243.44 1.0093 16.5992 0.9704 15.1385 0.9731 18.3389 0.9672 20.3708 0.9636 (0.6587) (-20.0)0.078 248.36 0.7953 0.102 254.67 1.0224 0.106 261.07 1.0489 0.111 267.58 1.0748 (245.195) 25.0541 0.9636 29.4619 0.9511 27.7144 0.9558 26.2493 0.9600 (0.6858) (-10.0)0.067 253.10 0.7833 0.072 257.60 1.0101 0.075 266.20 1.0364 0.078 272.89 1.0620 (354,030) 34.1528 0.9523 35.0892 0.9476 37.7589 0.9364 36.2872 0.9423 N (0.7129) 20 (0.0) 0.050 257.60 0.9729 0.052 264.32 0.9996 0.055 271.13 1.0256 0.057 278.02 1.0511 (496.550) 42.5264 0.9392 43.2080 0.9333 45.3461 0.9193 44.1440 0.9267 (0.7400)(10.0)(678.972) · 0.037 261.80 D.9637 0.039 268.77 0.9903 0.040 275.81 1.0163 0.042 282.93 1.0417 51.3509 0.9091 50.6746 0.9171 50.2457 0.9243 52.2864 0.9001 (0.7670) (20.0)0.029 272.89 0.9819 0.030 280.20 1.0079 0.032 287.57 1.0333 0.028 265.62 0.9552 (907.922) 57.3660 0.9077 57.5412 0.8992 57.9570 0.8896 58.6253 0.8789 (0.7941) (30.0) 0.024 291.89 1.0258 0.023 284.23 1.0003 0.022 276.60 0.9741 0.021 268.98 0.9471 (1190.426) 63.9288 0.8896 63.8460 0.8796 63.9965 0.8684 64.3921 0.8558 (0.8212)(40.0)0.018 287.84 0.9930 0.019 295.83 1.0187 0.017 279.83 0.9665 0.016 271.78 0.9390 (1533.915)69.9638 0.8702 69.6145 0.8586 69.4894 0.8456 69.6000 0.8311 (0.8483) (50.0) 0.015 299.35 1.0120 0.014 290.96 0.9859 0.013 273.89 0.9305 0.013 282.49 0.9588 (1946.243) 75.4915 0.8497 74.8610 0.8364 74.4423 0.8216 74.2449 0.8051 (0.8754)
·	-6 TABLE	=2) F	(CCL2F-CCL	IGERANT 113	HANE REF	DTRIFLUOROET	S OF TRICHLOR	TED PROPERTIE	CATHRA	
Т	Т	 S(G)	 S(F)		H(FG)	н(Е)				
DEG	DEG K	KJ/KG.K	KJ/KG.K	KJ/KG	KJ/KG	KJ/KG	M3/KG	KG/M3	P PA	T DEG C
-40.	233.2	0.7053	0.0000	164.45	164.45	0.00	 4.4873	4706 0	4582.7	
-30.	243.2	0.7009	0.0334	167.81	162.30	7.51	Z' 5284	4770 7	1572.7	740.0
-20.	253.2	0.6986	0.0661	175.30	160.13	15 17	2 0293	1777.3	3050.7	30.0
-10.	263.2	0.6982	0.0980	180.91	157.92	22.99	1.2258	1741.4	5515.U 9470.4	10.0
ο.	273.2	0.6993	0.1294	186.63	155.66	30.96	0 7770	4704 4	15517 0	
10.	283.2	0.7018	0,1602	192.44	153.34	20.00	0.7730	1/21.1	15543.2	0.0
20,	293.2	0.7054	0.1905	198.32	150.94	47 29	0.3002	1077.7	24508.0	10.0
30.	303.2	0.7100	0.2203	204.28	418 11	SE 84	0.3720	16//.3	37299.5	20.0
40.	. 313.2	0.7154	0.2497	210.28	145.82	55.04	0.1707	1653.6 1628.7	55001.6 78856.6	30.0
50,	323.2	0.7214	0.2787	216 32	4/7 04	77 05				
60.	333.2	0.7279	0.3073	222 24	143.00	(3.25	0.1246	1602.4	110250.5	50.0
70.	343.2	0.7349	0.3356	228.00	477 04	02.22	0.0930	1574.5	150711.3	60.0
80.	353.2	0.7422	0.3636	220.71	137.04	91.37	0.0705	1545.1	201896.2	70.0
90.	363.2	0.7497	0.3913	240.40	133.71	110.26	0.0543	1513.8 1480.5	265586.8 343681.2	80.0 90.0
100.	373.2	0.7573	0.4188	266 30	424 28	420.02	0.077/	-		
110.	383.2	0.7649	0.4462	252 11	120.27	120.02	0.0334	1444.9	438188.3	00.0
120.	393.2	0.7724	0 4735	202.11	122.10	130.01	0.0266	1406.9	551223.0	10.0
130,	403.2	0.7798	0.4705	23/./9	117.53	140.25	0.0214	1366.1	685006.8	20.0
140.	413.2	0.7870	0.5281	268.58	112.51 106.97	150.77	0.0172	1322.0	841865.8	30.0
150.	423.2	0 7937	0 5555	077 (0				12/702	1024237.0	40.0
160	433.2	0.7708	0.3333	2/3.60	100.78	172.82	0.0114	1222.0	1234667.1	50.0
170.	443 2	0 8052	0.3033	2/8.2/	93.80	184.46	0.0093	1164.5	1475829.9	60.0
180.	453.2	0.0052	0.0115	282.47	85.82	196.65	0.0075	1100.5	1750532.8	70.0
190.	463.2	0.0074	0.0407	286.02	76.48	209.54	0.0061	1027.7	2061728.8	80.0
	40012	0.0117	0.0/12	288.62	65.17	223.45	0.0048	942.8	2412540.3	90.0
200.	473.2	0.8113	0.7045	289.56	50.52	239.04	0.0038	837.9	2806277.0	0.0.0
210	483.Z	0.8024	0.7453	286.28	27.57	258.71	0.0027	685.7	3246474 . R	10.0
211.	484.2	0.8001	0.7506	285.27	23.98	261.28	0.0026	663.5	3293191.5	11.0
212	485.2	0.7971	0.7565	283.87	19.72	264.15	0.0024	638.1	3340414.0	12.0
213.	486.2	0.7923	0.7633	281.60	14.10	267.50	0.0022	607.2	3388143.3	13.0

	 TD	р	PR	Z(F)	Z(G)	FC(F)	FC(G)	B(F)	B(G)	Т
DEG C	I R	PA	••••					KJ/KG 	KJ/KG	DEG (
		4500 7	0 0005	n_0001	0.9987	0.9980	0.9987	0.94E+01	-0.36E+02	-40.0
-40.0	0.4785	1372.7	0.0005	0.0002	0.9977	0.9977	0.9977	0.70E+01	-0.29E+02	-30.0
-30.0	0.4990	3050.7	0.0007	0,0002	0.9963	0.9963	0.9963	0.49E+01	-0.23E+02	-20.0
-20.0	0.5195	5515.0	0.0010	0.0005	0.9942	0.9943	0.9942	0.32E+01	-0.17E+02	-10.0
-10.0	0.5401	7470.4	0.0020	0 0007	0.9913	0.9914	0.9913	0.19E+01	-0.12E+02	0.0
0.0	0.5606	15543.2	0.0045	0.0001						
		0/508 0	0 0074	0.0011	0.9874	0.9877	0.9876	0.87E+00	-0.67E+01	10.
10.0	0.5811	24500.0	0.0408	0.0017	0.9824	0.9828	0.9827	0.16E+00	-0.19E+01	20.0
20.0	0.6016	37299.5	0.0108	0.0025	0.9762	0.9768	0.9767	-0.24E+00	0.27E+01	30.0
30.0	0.6222	55001.0	0.0100	0.0025	0.9685	0.9694	0.9694	-0.35E+00	0.71E+01	40.1
40.0	0.6427	78856.6	0.0227	0.0035	0.7003	0.9607	0.9607	-0.16E+00	0.11E+02	50.0
50.0	0.6632	110250.5	0.0321	0.0040	0.7572	017001				
			0 0/78	0.0045	0 9482	0,9506	0.9505	0.30E+00	0.15E+02	60.0
60.0	0.6837	150/11.3	0.0430	0.0005	0.7402	0.9390	0.9389	0.11E+01	0.19E+02	70.0
70.0	0.7043	201896.2	0.0567	0.0442	0.7353	0.9259	0.9259	0.21E+01	0.23E+02	80.
80.0	0.7248	265586.8	0.0772	0.0112	0.7204	0.9114	0.9113	0.34E+01	0.27E+02	90.0
90.0	0.7453	343681.2	0.0999	0.0144	0.7035	0.2955	0.8754	0.50E+01	0.31E+02	100.
100.0	0.7658	438188.3	0.12/4	0.0103	0.0043	010/00		8	2	
			0.4400	0.0270	0 8427	0.8782	0.8782	0.68E+01	0.34E+02	110.
110.0	0.7863	551223.0	0.1602	0.0230	0.0027	0 8597	0.8597	0.89E+01	0.38E+02	120./
120.0	0.8069	685006.8	0.1991	0.0267	0.8365	0.8400	0.8400	0.11E+02	0.41E+02	130.1
130.0	0.8274	841865.8	0.2447	0.0350	0.0110	a 1 2193	0.8193	0.14E+02	0.44E+02	140.
140.0	0.8479	1024237.0	0.2978	0.0430	0.7484	0.0170	0.7975	0.17E+02	0.47E+02	150./
150.0	0.8684	1234667.1	0.3589	0.0530	0./400	0.7770				
				0.0/50	0 7444	0 7750	0.7749	0.21E+02	0.50E+02	160.
160.0	0.8890	1475829.9	0.4290	0.0657	0./110	0.7750	0.7516	0.24E+02	0.53E+02	170.
170.0	0.9095	1750532.8	0.5089	0.0809	0.0701	0.7510	0 7276	0.29E+02	0.55E+02	180.
180.0	0.9300	2061728.8	0.5994	0.0998	0.6231	0.7074	0 7034	0.33E+02	0.57E+02	190.
190.0	0.9505	2412540.3	0.7014	0.1245	U.5665	0.7031	n 4782	0.39E+02	0.58E+02	200.
200.0	0.9710	2806277.0	0.8158	0.1595	0.5018	0.6/82	0.0/02	010/2/01		
					0 /050	0 4670	D 6529	0.47E+02	0.57E+02	210.
210.0	0.9916	3246474.8	0.9438	0.2208	0.4059	0.0530	0.6514	0.48E+02	0.57E+02	211.
211.0	0.9936	3293191.5	0.9574	0.2310	0.3916	0.6504	0.0304	0.49E+02	0.56E+02	212.
212.0	0.9957	3340414.0	0,9711	0.2432	0.3747	0.6479	0.04/7	0.505+02	0.56E+02	213.
213.0	0.9977	3388143.3	0.9850	0.2587	0.3518	0.6453	0.0423	0.302.02		

OF COLOR-COLES CONTINUED F6

	SATD COND (DEG C) (KPA) (TR)	10K V(6) H(6) S(6) M3/KG KJ/KG KJ/KG.K B(6) FC(6) K1/KG	DEGREES OF SUPERHEAT 20K V(6) H(6) S(6) M3/KG KJ/KG KJ/KG.K B(6) FC(6) KJ/KG	30K V(6) H(G) S(6) M3/KG KJ/KG KJ/KG.K B(G) FC(6) KJ/KG	40K V(6) H(6) S(6) M3/KG KJ/KG KJ/KG.K B(6) FC(6) KJ/KG
		KJ/KG			
	(-40.0) (1.593) (0.4785)	6.766 169.84 0.7295 -37.6238 0.9988	7.046 .175.38 0.7539 -39.2489 0.9989	7.325 181.06 0.7776 -40.6179 0.9990	7.604 186.87 0.8009 -41.7389 0.9991
	(-30.0) (3.051) (0.4990)	3.674 175.35 0.7250 -30.6858 0.9979	3.820 181.03 0.7487 -32.0549 0.9981	3.966 186.85 0.7720 -33.1761 0.9983	4.112 192.80 0.7950 -34.0569 0.9985
	(-20.0) (5.515) (0.5195)	2.110 180.98 0.7223 -24.2628 0.9966	2.191 186.80 0.7457 -25.3842 0.9969	2.272 192.76 0.7687 -26.2652 0.9972	2.353 198.84 0.7913 -26.9130 0.9974
212	(-10.0) (9.470) (0.5401)	1.273 186.73 0.7215 -18.2776 0.9947	1.320 192.69 0.7445 -19.1588 0.9952	1.368 198.77 D.7672 -19.8067 D.9956	1.415 204.98 0.7895 -20.2284 0.9960
	(0.0) (15.543) (0.5606)	0.802 192.59 0.7223 -12.6618 0.9921	D.831 198.68 0.7450 -13.3098 0.9928	0.860 204.89 0.7673 -13.7315 0.9934	0.887 211.22 0.7893 -13.9335 0.9939
	(10.0) (24.508) (0.5811)	-0.525 198.53 0.7245 -7.3586 0.9886	0.543 204.75 0.7468 -7.7802 0.9896	0.562 211.09 0.7689 -7.9821 0.9905	0.580 217.55 0.7905 -7.9706 0.9912
	(20.0) (37.300) (0.6016)	0.355 204.56 0.7278 -2.3188 0.9842	0.367 210.91 0.7498 -2.5203 0.9855	0.379 217.37 0.7716 -2.5082 0.9867	0.391 223.95 0.7930 -2.2889 0.9877
	(30.0) (55.002) (0.6222)	0.247 210.64 0.7321 2.4951 0.9786	0.255 217.12 0.7538 2.5080 0.9803	0.264 223.71 0.7753 2.7283 0.9819	0.272 230.41 0.7964 3.1501 0.9833
a)	(40.0) (78.857) (0.6427)	D.177 216.78 D.7372 7.1158 D.9718	0.182 223.38 0.7586 7.3377 0.9740	0,188 230.09 0.7798 7.7613 0.9761	0.194 236.91 0.8007 8.3807 0.9779
	(50.0) (110.251) (0.6632)	0.129 222.94 0.7429 11.5692 0.9638	0.133 229.67 0.7642 11.9952 0.9666	D.138 236.51 D.7851 12.6173 D.9691	0.142 243.44 0.8057 13.4296 0.9714

SUPERHEATED PROPERTIES OF TRICHLOROTRIFLUOROETHANE

REFRIGERANT R113 (CCL2F-CCLF2) -6

		8 I		8		2 II				
T DEG (Р С РА	D(F) Kg/N	V(6) 3 M3/KG	H(F) KJ/KG	н(FG) Кј/Кб	KJ/KG	KJ/KG.K	KJ/KG.K	DEG K	DEG
					7/4 84	744 84		1.6666	233.2	-40.0
-40.0	J 48794	.7 9/3.		0.00	341.74	341.74	0.0534	1 4296	243.2	-30.0
-30.0	J 79293	.1 956.2	2 0.3/61	11.42	334./1	340.13	0.0551	4 3982	253.2	-20.0
-20.0		.4 937.0	6 U.249L	23.43	327.03	350.40 754 87	0.1598	1.3713	263.2	-10.0
-10.0	J 10404/	./ 71/10	D 0.1/0-	. 30.02	510.01	034100				
0.0	267976	.2 896.0	0.1199	49.21	309.93	359.14	0.2133	1.3479	273.2	0.0
10.0	377443	.0 872.1	6 0.0864	63.02	300.27	363.29	0.2671	1.3275	283.2	10.0
20.0	518325	.2 847.	7 0.0636	77.51	289.66	367.17	0.3212	1.3092	293.2	20.0
30.0	696068	.1 820.3	3 0.0475	92.76	277.93	370.70	0.3758	1.2926	303.2	30.0
40.0	916478	.0 790.4	6 0.0360	108.88	264.85	373.73	0.4313	1.2771	313.2	40.0
50.0		0 757	0 0275	124 07	250.44	376.16	0.4881	1.2620	323.2	50.0
50.0		.0 /3/	7 0.02/3	444 44	223 22	377.73	0.5466	1.2469	333.2	60.0
60.0	J 151U32U	.6 /20.0			233.34	778 27	0.6077	1.2309	343.2	70.0
70.0	1897221	.5 6/9.4	0.0163	104.30	213.07	3/0.23	0.0077	4 2129	353.2	80.0
80.0	2353793	.8 632.1	0.0125	186.38	190.04	3//.21	0.7472	4 4 9 0 8	363 2	90.0
90.0	2887910.	.5 576.2	2 0.0095	211.32	162.56	3/3.00	0.7432	1.1700	303.2	/010
100.0	3508027.	5 506.3	s 0.0070	241.11	125.31	366.42	0.8244	1.1602	373.2	100.0
110.0	4223284	.0 401.9	9 0.0047	282.77	64.76	347.53	0.9336	1.1027	383.2	110.0
444 0	4220204	5 386.0	0.0044	288.84	54.74	343.58	0.9493	1.0918	384.2	111.0
412 0		5 367.0	0.0041	295-96	42.28	338.24	0.9676	1.0773	385.2	112.0
113.0	4457739.	.5 341.9	0.0036	305.21	22.01	327.21	0.9913	1.0483	386.2	113.0
3 3			SATURAT	ED PROPERTIES OF	CH3CHF2	CONTINU	ED			
				7(5) 7(6		EC(E)	FC(G)	B(F)	B(G)	Т
T DEG C	TR	PA	F K		,			KJ/KG	KJ/KG	DEG
						0 0929	n.9827	0.80E+02	-0.13E+02	-40.0
-40.0	0.6030	48794.7	0.0109	0.001/ 0.98	223	0.7020	0.0749	D.76E+02	0.18E+01	-30.0
-30.0	0.6289	79293.1	0.0176	0.0027 0.97	43 ° a	0.9750	0.7747	0.72E+02	0.15E+02	-20.0
-20.0	0.6547	123372.4	0.0274	0.0041 0.98	030	0.7031	0.7030	0.695+02	0.28E+02	-10.0
-10.0	0.6806	184847.7	0.0411	0.0061 0.95	507	0.7527	0.7320	0 665+02	0.39E+02	0.0
0.0	0.7065	267976.2	0.0596	0.0087 0.93	546	0.7304	0.7303	01002.02	••••	
40.0	0 7727	777447 0	0.0840	0.0121 0.91	54	0.9215	0.9215	0.64E+02	0.49E+02	10.0
10.0	0.7523	511773.0	0 1153	0.0166 0.89	127	0.9024	0.9023	0.63E+02	0.59E+02	20.0
20.0	0.7562	310343.4 494048 4	0.1133	0.0222 0.84	63	0.8810	0.8810	0.62E+02	0.67E+02	30.0
30.0	U.784U	070000.1	0.1340	0.0294 0.87	58	0.8577	0.8577	0.61E+02	0.75E+02	40.0
40.0	0.8099	9164/8.U	U.2037	0.0385 0.80	09	0.8324	0.8324	0.61E+02	0.82E+02	50.0
50.0	0.0350	1105/17.0	0.1000	010002					0.005.00	60.0
60-0	0.8616	1510320.6	0.3360	0.0500 0.76	510	0.8056	0.8055	U.62E+02	U.801+UZ	70 0
70.0	0.8875	1897221.5	0.4220	0.0646 0.71	54	0.7772	0.7772	0.64E+02	U.93E+U2	/0.0
80.0	0.9134	2353793.8	0.5236	0.0838 0.66	526	0.7476	0.7476	0.67E+02	U.Y/E+UZ	
90.0	0.9392	2887910.5	0.6424	0.1096 0.60	005	0.7171	0.7171	0.71E+02	0.10E+03	90°L
100.0	0.9651	3508027.5	0.7804	0.1475 0.52	235	0.6857	0.6857	0.77E+02	0.10E+03	100.0
	0.0007	(00708/ 0	0.0705	0 2178 D 44	04	0.6538	0.6538	0.86E+02	0.10E+03	110.0
110.0	0.9909	4223284.U	0.7373	0.21/0 0.41	225	0.6506	0.6506	0.87E+02	0.10E+03	111.0
111.0	0.9935	4300419.5	0.9566	0.2307 0.37	123	n 4472	0.6473	0.87E+02	0.99E+02	112.0
112.0	0.9961	4378615.5	0.9740	U.2461 U.3/	100	0.04/3	0 4444	0.91E+02	0.96E+02	113.0
113 0	0 0087	4457739.5	0.9916	U.2682 U.33	010	U.0441	0.0441			

		SUPERHEATED P	RUPERTIES U					
SATD COND (DEG C) (KPA) (TR)	10K V(6) H(6) M3/KG KJ/1) S(G) (G KJ/KG.K	DEG V(G) M3/KG	REES OF SUPERHEAT 20K H(G) S(G) KJ/KG KJ/KG.K	30) V(6) H(0 M3/KG KJ/	(5) 5(6) /Kg Kj/Kg.K	40K V(6) H(6) M3/KG KJ/KG	S(6) Kj/kg.k
	B(6) Kj/kg	FC(6)	B(6 KJ/KG) FC(G)	B(6) Kj/kg	FC(6)	8(G) Kj/Kg	FC(G)
(-40.0) (48.795) (0.6030)	0.618 346.9 -16.1993	95 1.4929 0.9846	0.644 -18.77	352.36 1.5197 49 0.9861	0.670 358 -21.0973	.14 1.5470 0.9875	0.697 364.25 -23.1528	1.5745 0.9887
(-30.0) (79.293) (0.6289)	0.393 351.9 -0.8080	59 1.4567 0.9775	0.409	357.42 <u>1.4841</u> 700.9797	0.426 363. -5.1969	.57 1.5118 0.9817	0.442 370.04 -6.9835	1.5375 0.9834
(- 20.0) (123.372) (0.6547)	0.260 356.3 13.1054	5 1.4259 0.9685	0.271	362.57 1.4538 89 0.9715	0.281 369. 9.2485	.09 1.4817 0.9742	0.292 375.90 7.7376	1.5097 0.9766
(-10.0) (184.848) (0.6806)	0.178 361.1 25.7418	14 1.3995 0.9574	0.185 23.94	367.74 1.4277 57 0.9614	0.192 374 22.4324	.63 1.4560 0.9649	0.200 381.79 21.2013	1.4843 0.9681
(0.0) (267.976) (0.7065)	0.125 365.0 37.2532	57 1.3766 0.9441	0.130 35.73	372.86 1.4053 64 0.9493	0.136 380 34.5057	.13 1.4339 0.9538	0.141 387.67 33.5614	1.4624 0.9579
(10.0) (377.443) (0.7323)	- 0.090 370.4 47.7636	45 1.3567 0.9287	0.094 46.53	377.86 1.3858 33 0.9351	0.098 385 45.5937	.54 1.4148 0.9408	0.102 393.49 44.9432	1.4437 0.9459
(20.0) (518.325) (0.7582)	0.067 374.0 57.3669	51 1.3391 0.9111	0.069 56.43	382.68 1.3687 42 0.9190	0.072 390. 55.7950	.81 1.3782 0.9260	0.075 399.23 55.4501	1.4277 0.9323
(30.0) (696.068) (0.7840)	0.050 378.0 66.1354	86 1.3232 0.8915	0.052 65.51	387.25 1.3535 32 0.9009	0.054 395 65.1896	.90 1.3837 0.9093	0.057 404.84 65.1678	1.4139 0.9169
(40.0) (916.478) (0.8099)	0.038 382. 74.1187	53 1.3087 0.8700	0.040 73.82	391.52 1.3399 76 0.8811	0.041 400. 73.8418	.76 1.3710 0.8911	0.043 410.32 74.1683	1.4020 0.9000
(50.0) (1185.717) (0.8358)	0.029 385.	70 1.2950 0.8468	0.031 81.41	395.41 1.3274 94 0.8597	0.032 405 81.8036	.37 1.3597 0.8713	0.033 415.65 82.5078	1.3919 0.8817

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T	P	D(F)	V(6)	 H(F)	H(FG)	H(G)	S(F)	S(G)	т	т
DEG C	PA	KG/M3	M3/KG	KJ/KG	KJ/KG	KJ/KG	KJ/KG.K	KJ/KG.K	DEG K	DEG
	570.4		220.9413	0.00	2559.09	2559.09	0.0000	9.3686	273.2	0.0
10.0	1147 9	854.5	112.2705	46.44	2531.22	2577.66	0.1838	9.1230	283.2	10.0
20.0	2264 6	849.1	60.2128	90.70	2505.55	2596.25	0.3534	8.9001	293.2	20.0
30.0	4124.6	843.6	33.9049	134.94	2479.90	2614.85	0.5173	8.6977	303.2	30.0
			40 0277	478 20	2666 22	2633.63	0.6759	8.5129	313.2	40.0
40.0	/24/.9	874 8	42 4747	223 51	2434.22	2651.97	D.8296	8.3444	323.2	50.0
50.0	12233.0	031.0	7 7050	243.31	2420.40	2670.46	0.9790	8.1904	333.2	60.0
60.0	19910.0	825.5	7.7057. F 07/4	201.07	2702.37	2670.40	1.1242	8.0494	343.2	70.0
70.0	31355.9	819.1	5.0340	312.37	23/0.40	2000.03	1.2656	7.9201	353.2	80.0
80.0	47928.5	812.3	3.3050	357.03	2350.07	2707.10	1.2030			
90.0	71297.4	805.4	2.3357	401.86	2323.32	2725.17	1.4036	7.8011	363.2	90.0
100.0	103459-2	798.1	1.6501	446.90	2296.11	2743.01	1.5384	7.6916	3/3.Z	100.0
110.0	146771.2	790.6	1.1909	492.21	2268.35	2760.56	1.6703	7.5904	383.2	110.0
120.0	203949-2	782.7	0.8763	537.82	2239.93	2777.75	1.7995	7.4968	393.2	120.0
130.0	278078.7	774.6	0.6562	583.77	2210.73	2794.50	1.9263	7.4098	403.2	130.0
		7// 4	0 4007	470 42	2180 44	2810.76	2.0509	7.3288	413.2	140.0
140.0	372614.3	767.3	0.4773	474 82	2100.04	2826.43	2.1734	7.2531	423.2	150.0
150.0	471374.4	757.2	0.3034	726 22	2147.31	2020.40	2.2943	7.1821	433.2	160.0
160.0	638537.3	741.7	0.3013	772 00	2083 67	2855 66	2.4135	7.1151	443.2	170.0
170.0	818628.3	130.2	0.2300	772:U7 820 59	2003.37	2033.00	2.5314	7.0517	453.2	180.0
180.0	1036503.6	728.1	0.1707	620.57	2090.93	2007.02	215014			
190.0	1297348.9	717.4	0.1542	867.81	2011.61	2881.42	2.6481	6.9914	463.2	190.0
200 0	1404442 1	706.3	0.1257	919.83	1972.90	2892.73	2.7639	6.9335	473.2	200.0
210.0	4070244 4	694.6	0.1033	970.75	1932.09	2902.84	2.8790	6.8778	483.2	210.0
220.0	177024144 0704478 D	682.3	0.0854	1022.67	1888.93	2911.60	2.9935	6.8238	493.2	220.0
220.0	23771/0.U 2884859.A	669.3	0.0711	1075.73	1843.13	2918.86	3.1078	6.7710	503.2	230.0
23010	200403710	00710					5		547 0	2/0 0
240.0	3448960.0	655.6	0.0595	1130.07	1794.38	2924.45	3.2222	6.7189	513.2	240.0
250.0	4093420.8	641.1	0.0500	1185.86	1742.31	2928.18	3.3369	6.6673	523.2	250.0
260.0	4825493.5	625.7	0.0422	1243.32	1686.49	2929.81	3.4522	6.6154	533.2	260.0
270.0	5652703.0	609.3	0.0358	1302.68	1626.40	2929.08	3.5686	6.5630	543.2	2/0.0
280.0	6582902.0	591.8	0.0304	1364.26	1561.40	2925.66	3.6866	6.5073	553.2	200.0
200 0	7494948 0	573.0	0.0258	1428-44	1490.70	2919.14	3.8066	6.4537	563-2	290.0
270.0	10474701U 9795744 A	552 7	0.0220	1495.73	1413.28	2909.01	3.9295	6.3953	573.2	300.0
300.0	6/03/99.V 4007/7/7 0	532.7	0.0187	1566.77	1327.79	2894.57	4,0563	6.3332	583.2	310.0
310.0	100/9/9/9/.0	5017	0.0158	1642-52	1232.35	2874.87	4.1882	6.2659	593.2	320.0
320.0	11302014.0	280 N	0.0134	1724.32	1124-21	2848.53	4.3274	6.1913	603.2	330.0
330.0	130/0/21.0	400.0	010107	1127102						
340.0	14809037.0	449.8	0.0112	1814.35	998.97	2813.32	4.4769	6.1061	613.2	340.0
350.0	16709607.0	414.8	0.0093	1916.50	848.76	2765.27	4.6425	6.0045	623.2	350.0
360.0	18789686.0	371.8	0.0074	2039.32	656.22	2695.55	4.8369	5.8733	653.2	360.0
370.0	21061154.0	309.8	0.0056	2213.22	357.66	2570.88	5.1061	5.6622	643.2	370.0
371.0	21299302.0	300.8	0.0053	2238.18	311.22	2549.40	5.1444	5.6275	644.2	3/1.U
372 0	04530508 D	290.5	0.0051	2266.66	255.99	2522.66	5.1880	5.5848	645.2	372.0
777 0	21337320.U 31791704 0	278.3	0.0047	2300.79	182.26	2483.06	5.2403	5.5223	646.2	373.0
57.5.11	21/01/00.0	<u> </u>		*******						

	T DEG C	TR	P PA	PR	Z(F)	Z(G)	FC(F)	FC(6)	B(F) Kj/kg	B(G) KJ/KG	T DEG (
	0.0	0.4220	570.6	0.0000	0.0000	0.9999	1.0003	0.9999	0.16E+02	-0.21E+03	
	10.0	0.4374	1163.9	0.0001	0.0000	0.9998	1.0000	0.9998	0.80E+01	-0.12E+03	10.0
	20.0	0.4529	2246.4	0.0001	0.0000	0.9996	0.9998	0.9996	0.19E+01	-0.32E+02	20.0
	30.0	0.4683	4124.6	0.0002	0.0000	0.9994	0.9999	0.9994	-0.26E+01	0.46E+02	30.0
	40.0	0.4838	7247.9	0.0003	0.0001	0.9990	0.9992	0.9990	-0.55E+01	0.12E+03	40.0
	50.0	0.4992	12233.0	0.0006	0.0001	0.9985	0.9986	0.9985	-0.69E+01	0.19E+03	50.0
	60.0	0.5147	19910.0	0.0009	0.0002	0.9977	0.9978	0.9977	-0.68E+01	0.25E+03	60.0
	70.0	0.5301	31355.9	0.0014	0.0002	0.9967	0.9968	0.9967	-0.55E+01	0.31E+03	70.0
	80.0	0.5456	47928.5	0.0022	0.0004	0.9953	0.9954	0.9953	-0.29E+01	0.37E+03	80.0
	90.0	0.5610	71297.4	0.0032	0.0005	0.9935	0.9936	0.9935	0.94E+00	0.42E+03	90.0
	100.0	0.5765	103459.2	0.0047	0.0008	0.9912	0.9913	0.9913	0.59E+01	0.47E+03	100.0
	110.0	0.5919	146771.2	0.0067	0.0010	0.9883	0.9885	0.9885	0.12E+02	0.52E+03	110.0
	120.0	0.6074	203949.2	0.0093	0.0014	0.9848	0.9851	0.9850	0.19E+02	0.57E+03	120.0
	130.0	0.6228	278078.7	D.0126	0.0019	0.9806	0.9810	0.9809	0.28E+02	0.61E+03	130.0
	140.0	0.6383	372614.3	0.0169	0.0026	0.9756	0.9762	0.9761	0.37E+02	0.65E+03	140.0
	150.0	0.6537	491374.4	0.0223	0.0033	0.9697	0.9706	0.9705	0.47E+02	0.69E+03	150:0
	160.0	0.6692	638539.3	0.0290	0.0043	0.9628	0.9641	0.9641	0.59E+02	0.72E+03	160.0
	170.0	0.6846	818628.3	0.0371	0.0054	0.9550	0.9568	0.9568	0.71E+02	0.76E+03	170.0
2	180.0	0.7001	1036503.6	0.0470	0.0068	0.9461	0.9487	0.9486	0.85E+02	0.79E+03	180.0
σ	190.0	0.7155	1277348.9	0.0588	0.0085	0.9360	0.9396	0.9396	0.99E+02	0.82E+03	190.0
	200.0	0.7310	1606662.1	0.0729	0.0104	0.9248	0.9297	0.9296	0.11E+03	0.85E+03	200.0
	210.0	0.7464	1970241.4	0.0894	0.0127	0.9122	0.9188	0.9188	0.13E+03	0.88E+03	210.0
	220.0	0.7619	2394178.0	0.1086	0.0154	0.8784	0.9071	0.9071	0.15E+03	0.90E+03	220.0
	230.0	0.7773	2884859.0	0.1308	0.0186	0.8831	0.8945	0.8945	0.17E+03	0.92E+03	230.0
	240.0	0.7928	3448960.0	0.1564	0.0222	0.8664	0.8811	0.8811	0.19E+03	0.94E+03	240.0
	250.0	0.8082	4093420.8	0.1857	0.0264	0.8480	0.8669	0.8669	0.21E+03	0.96E+03	250.0
	260.0	0.8237	4825493.5	0.2189	0.0313	0.8281	0.8520	0.8519	0.23E+03	0.98E+03	260.0
	270.0	0.8391	5652703.0	0.2564	0.0370	0.8064	0.8363	0.8363	0.26E+03	0.10E+04	270.0
	280.0	0.8546	6582902.0	0.2986	0.0436	0.7827	0.8199	0.8199	0.28E+03	0.10E+04	280.0
	290.0	0.8700	7624248.0	0.3458	0.0512	0.7570	0.8029	0.8029	0.31E+03	0.10E+04	290.0
	300.0	0.8855	8785244.0	0.3985	0.0601	0.7290	0.7854	0.7854	0.34E+03	0.10E+04	300.0
	310.0	0.9009	10074747.0	0.4569	0.0705	0.6984	0.7673	0.7673	0.38E+03	0.10E+04	310.0
	320.0	0.9164	11502014.0	0.5217	0.0829	0.6648	0.7488	0.7488	0.41E+03	0.10E+04	320.0
	330.0	0.9318	13076721.0	0.5931	0.0979	0.6276	0.7298	0.7278	0.45E+03	0.10E+04	330.0
	340.0	0.9473	14809037.0	0.6717	0.1163	0.5858	0.7105	0.7105	0.50E+03	0.10E+04	340.0
	350.0	0.9627	16709607.0	0.7579	0,1400	0.5376	0.6709	0.6709	0.55E+03	0.10E+04	350.0
	360.0	0.9782	18789686.0	0.8522	0.1729	0.4789	0.6710	0.6710	0.62E+03	0.97E+03	360.0
	370.0	0.9936	21061154.0	0.9552	0.2290	0.3944	0.6509	0.6509	0.71E+03	0.90E+03	370.0
	371.0	0.9951	21299302.0	0.9660	0.2382	0.3817	0.6489	0.6489	0.73E+03	0.89E+03	371.0
	372.0	0,9967	21539528.0	0.9769	0.2490	0.3665	0.6468	D.6468	0.74E+03	0.88E+03	372.0

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			SU	PERHEATED	PROPERTIES	OF WATE	R 		REFRIGE	RANT R718	(H20	, E 8	3
	SATD COND (deg C) (kpa)	V(G)	10K H(g)	S(G)	0 V(6)	EGREES 0 20k H(G)	F SUPERHEAT	V(G)	30K H(6)	5(6)	V(6)	40K H(6) K1/K6	S(G)
	(TR)	M3/KG	KJ/KG	KJ/KG.K	M3/KG	KJ/KG	KJ/KG.K	M3/KG	KJ/KG	KJ/KG.K			NJ/NG.1
		8(6 Kj/k	5) (G	FC(6)	8 KJ/	(G) KG	FC(6)	KJ/	B(G) 'KG		KJ/	KG	
	(0.0) (0.571) (0.4220)	229.032 -214.9	2577.70 257 0	9.4521 .9999	237.122 -220.	2596.34 2704	9.5329 0.9999	245.212 -224.	2615.03 8078 0	9.6110 .9999	253.303 -228.	_2633.75 5940 (° 9.6868 3.9999
	(10.0) (1.164) (0.4374)	116.237 -122.5	2596. 31 105 0	9.2038 .9998	120.204 -127.	2615.00 0499	9.2819 0.9998	124.171 -130.	2633.72 8343 0	9.3577 .9998	128.137 -133.	2652.49 9079 (9.4312 .9999
	(20.0) (2.246) (0.4529)	62.269 -36.8	2614.94 809 0	8.9783 .9997	64.324 -40.	2633.67 6651	9.0541 0.9997	66.380 -43.	2652.44 7387 0	9.1276 .9997	68.435 -46.	2671.25 1428 (9.1990 .9997
217	(30.0) (4.125) (0.4683)	35.025 42.6	2633.58 359 0	8.7734 .9995	36.145 39.	2652.35 5625	8.8470 0.9995	37.265 37.	2671.17 1587 0	8.9184 .9995	38.385 35.	2690.03 3880 (8.9878).9996
	(40.0) (7.248) (0.4838)	20.561 116.8	2652.21 211 0	8.5865 .9991	21.199 114.	2671.03 4183 (8.6579 0.9992	21.836 112.	2689.89 6481 0	8.7274 .9992	* 22.474 111.	2708.80 4768 (8.7949).9993
	(50.0) (12.233) (0.4992)	-12.553 186.1	2670.80 071 0	8.4159 .9986	12.931 184.	2689.68 3382 (8.4854 0.9987	13.309 183.	2708.59 1681 0	8.5530 .9988	13.687 182.	`2727.55 5668 (8.6188].9989
	(60.0) (19.910) (0.5147)	7.939 250.9	268 9.34 892 0	8.2600 .9979	8.171 249.	2708.27 8209	8.3276 D.9981	8.404 249.	2727.25 2218 0	8.3935 .9982	8.636 249.	2746.26 1626 (8.4576 .9983
	(70.0) (31.356) (0.5301)	5.183 311.8	2707 .79 857 0	8.1171 .9969	5.330 311.	2726.79 2894 (8.1830 0.9972	5.478 311.	2745.82 2339 0	8.2473 .9974	5.626 311.	2764.89 6914 (8.3099 .9975
	(80.0) (47.928) (0.5456)	3.482 369.1	2726.12 585 O	7.9860 .9957	3.579 369.	2745.18 1076 (8.0504 0.9960	3.676 369.	2764.28 5705 0	8.1131 .9962	3.773 370.	2783.42 5239 (8.1743).9965
	(90.0) (71.297)	2.401	2744.27	7.8655	2.466	2763.40	7.9283	2.532	2782.58	7.9896	2.597	2801.79 9829 (8.0495

- 1 - P	ter :						-			
		×	19 1 967	к –	a — 1	10 De			1 ×	
DEG C	PA	KG/M3	M3/KG	KJ/K	G KJ/KG	KJ/KG	KJ/KG.K	KJ/KG.K	DEG K	DEG C
-40 0	77827.9	1667.8	0.2578	0.0	208.23	208.23	0.0000	0.8931	233.2	-40.0
-70.0	445442 7	1449.4	0.1698	10.8	8 203.65	214.53	0.0491	0.8866	243.2	-30.0
-30.0	47/4/9 3	4780 7	0.1158	21.9	3 198.73	220.66	0.0968	0.8818	253.2	-20.0
-10.0	253433.2	1356.9	0.0814	33.1	7 193.41	226.60	0.1435	0.8784	263.2	-10.0
				•			6 4893	0 8740	277 2	0.0
0.0	357979.2	1322.0	0.0586	44.7	187.61	232.31	U.1872	0.0700	213.2	10.0
10.0	492611.9	1284.4	0.0431	56.53	3 181.22	237.75	U.2344	0.0799	203.2	20.0
20.0	662 486.0	1243.5	0.0322	68.75	174.16	242.87	0.2/91	U.8/32	293.2	20.0
30.0	873062.2	1199.0	0.0244	81.33	3 166.26	247.59	0.3237	0.8721	303.2	30.0
40.0	1130115.0	1150.0	0.0187	94.43	7 157.38	251.84	0.3683	0.8708	313.2	40.0
FO O	4/70704 0	1005 8	1446	108.24	447.25	255.49	0.4134	0.8670	323.2	50.0
50.0	1437720.0	4075 4	0.0144	400 81	435 54	258.38	0.4593	0.8662	333.2	60.0
60.0	1808314.8	1035.1	0.0111	478 //	404 74	260.00	0.5067	0.8616	343.2	70.0
70.0	2242666.3	765.7	0.0000	130.40		200.23	0 5548	0.8540	353.2	80.0
80.0	2749985.8	884.9	0.0066	155.67	2 104.70	200.30	÷ 0.5500	D 8411	363.2	90.0
90.0	3337956.8	785.4	0.0049	1/5.22	63.24	230.40	0.0117	0.0411	00012	,010
			0 0074	200 31	50 38	250.70	0.6798	0.8148	373.2	100.0
100.0	4014831.8	072.7		200.37		249 17	П. 6883	0.8103	374.2	101.0
101.0	4087745.8	- 020.7	0.0033	203.5		247.11	0 4974	0.8049	375.2	102.0
102.0	4161649.5	605.9	0.0031	207.05	9 4U.ZO	247.32	0.0770	0.001/	376.2	103.0
103.0	4236546.5	582.1	0.0029	210.95	5 34.04	245.00	0.7077	0.7995	377 2	104.0
104.0	4312453.5	553.7	0.0027	215.51	26.29	241.80	U./170	0.7073	3//12	10110
			SATURAT	ED PROPERTIE	S OF R12 & F	152A CONTINL	IED			
					3/6)	EC(E)	FC(6)	B(F)	B(G)	т
т	TR	P	PK	2(F)	2(0)	E VAL 2		KJ/KG	KJ/KG	DEG C
DEG C		PA								
	A 1154	77807 0	0.0147	0 0026	0.9751	0.9757	0.9756	0.61E+02	0.40E+01	-40.0
-40.0	0.6158	13021.7	0.0107	0.0010	0 9667	0.9659	0.9658	0.57E+02	0.12E+02	-30.0
-30.0	0.6422	115642.3	0.0201	0.0040	0.0548	0.9538	0.9538	0.54E+02	0.20E+02	-20.0
-20.0	0.6686	174148.3	0.0373	0.0037	0.7510	0 9395	0.9395	0.52E+02	0.27E+02	-10.0
-10.0	0.6950	253433.2	0.0573	0.0005	0.7327	0.000	B. 9228	0.50E+02	0.33E+02	0.0
0.0	0.7214	357979.2	0.0809	0.0118	0.7167	0.7227	0.,			
40.0	0 7/70	100444 0	0 4443	0.0162	0.8946	0.9040	0.9039	0.48E+02	0.39E+02	10.0
10.0	0.7476	472011.7	0.1110	0.0247	0.8686	0.8829	0.8829	0.47E+02	0.45E+02	20.0
20.0	0.7742	662486.0	0.1477	0.0217	D 8788	0.8599	0.8599	0.46E+02	0.50E+02	30.0
30.0	0.8006	873062.2	0.1973	0.0275	0.0300	0.0377	0.8350	0.46E+02 *	0.54E+02	40.0
40.0	0.8270	1130115.0	0.2554	0.03/3	0.0040	0.0000	0 8086	0.47E+02	0.58E+02	50.0
50.0	0.8534	1439726.0	0,3253	0.0486	U./65/	0.0000	0.0000			
60-0	0.8798	1808314-6	0.4086	0.0626	0.7213	0.7808	0.7807	0.47E+02	0.62E+02	60.0 70 0
70.0	0.9062	2242666.3	0.5067	0.0808	0.6703	0.7517	0.7517	0.476+02	0.056402	20.0
80.0	0 0327	2749985 .8	0.6214	0.1051	0.6107	0.7218	0.7218	0.51E+02	U.68E+UZ	00.0
00.0	D.7327	3337954_8	0.7542	0.1398	0.5381	0.6911	0.6911	0.55E+02	0.70E+02	70.0
100.0	0.9855	4014831.8	0.9072	0.1989	0.4378	0.6578	0.6598	0.59E+02	0.70E+02	100.0
				0.0094	0 /2/4	0 4547	0.6567	0.60E+02	0.70E+02	101.0
101.0	0.9881	4087745.8	0.9236	0.2081	0.4271	0.6507	0.6535	0.61E+02	0.69E+02	102.0
102.0	0.9908	4161649.5	0.9403	0.2187	0.4070	0.0333	0.000	0.625+02	0.69E+02	103.0
103.0	0.9934	4236546.5	0.9573	0.2311	0.3715	0.0504	0.6304	D 63E+02	0.68E+02	104.0
104.0	0.9960	4312453.5	0.9744	0.2466	0.3698	0.6472	U.07/2	0.000-02	ataan an	

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SUPERHEATED PROPERTIES OF 73.8% CCL2F2 26.2% CH3CHF2 REFRIGERANT R500 (R12 & R152A) F9

	SATD COND	4.0%	DEGREES OF SUPERHEAT	30K	40K
	(DEG C) (KPA) (TR)	V(G) H(G) S(G) M3/KG KJ/KG KJ/KG.K	V(6) H(G) S(G) M3/KG KJ/KG KJ/KG.K	V(6) H(G) S(G) M3/KG KJ/KG KJ/KG.K	V(G) H(G) 5(G) M3/KG KJ/KG KJ/KG.K
		B(G) FC(G) KJ/KG	B(6) FC(6) KJ/KG	8(G) FC(G) Kj/kg	B(G) FC(G) KJ/KG
	(-40.0) (73.828)	0.270 215.22 0.9260	0.281 222.26 0.9577	0.293 229.34 0.9884 -3.2198 0.9823	0.305 236.48 1.0181 -4.9208 0.9840
~	(0.6158) (-30.0) (115.642) (0.6422)	1.2088 0.9782 0.178 221.60 0.9185 9.8110 0.9693	0.185 228.72 0.9493 7.7708 0.9723	0.193 235.90 0.9792 6.0660 0.9749	0.200 243.14 1.0082 4.6760 0.9773
50	(-20.0) (174.148) (0.6686)	0.121 227.84 0.9129 17.7113 0.9583	0.126 235.06 0.9429 16.0009 0.9623	0.131 242.36 0.9722 14.6077 0.9658	0.136 249.72 1.0006 13.5137 0.9689
219	(-10.0) (253.433) (0.6950)	0.085 233.90 0.9088 24.9927 0.9452	0.089 241.26 0.9383 23.5947 0.9503	0.092 248.69 0.9669 22.4988 0.9548	0.096 256.19 0.9949 21.6892 0.9588
	(0.0) (357.979) (0.7214)	0.061 239.77 0.9059 31.7236 0.9300	0.064 247.29 0.9349 30.6248 0.9364	0.067 254.88 0.9631 29.8154 0.9420	0.069 262.53 0.9907 29.2808 0.9470
	(10.0) (472.612) (0.7478)	° 0.045 245.41 0.9039 37.9558 0.9127	0.047 253.11 0.9325 37.1468 0.9204	0.049 260.88 0.9604 36.6160 0.9273	0.051 268.71 0.9877 36.3504 0.9335
	(20.0) (662.486) (0.7742)	0.034 250.75 0.9024 43.7281 0.8933	0.035 258.68 0.9309 43.2029 0.9026	0.037 266.65 0.9586 42.9465 0.9109 -	0.038 274.68 0.9856 42.9465 0.9184
	(30.0) (873.062) (0.8006)	0.026 255.77 0.9013 49.0662 0.8721	0.027 263.95 0.9297 48.8236 0.8831	0.028 272.17 0.9573 48.8409 0.8929	0.029 280.42 0.9842 49.1056 0.9017
	(40.0) (1130.115) (0.8270)	0.020 260.37 0.9003 53.9848 0.8492	0.021 268.87 0.9287 54.0288 0.8619	0.022 277.37 0.9564 54.3225 0.8733	0.023 285.88 0.9832 54.8544 0.8836
	(50.0) (1439.726) (0.8534)	0.015 264.49 0.8990 58.4859 0.8249	0.016 273.37 0.9277 58.8267 0.8394	0.017 282.21 0.9555 59.4053 0.8525	0.018 291.01 0.9825 60.2103 0.8643

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T	P	D(F)	V(6)	H(F)	H(FG)	H(6) K(/KG	S(F) KJ/KG.K	S(6) Kj/kg.k	T DEG K	T DEG
DEG C	PA	KG/M3	n3/KG							
-40.0	22.9	1808.2	511.6261	0.00	252.01	252.01	0.0000	1.0854	233.2	-40.0
-30.0	59.8	1796.9	203.8285	7,79	249.52	257.31	0.0349	1.0614	243.2	-20.0
-20.0	134.4	1785.4	94.4237	15.63	247.05	262.68	0.0685	1.0445	233.2	-10.0
-10.0	281.0	1772.1	46.9550	23.68	244.44	268.12	0.1017	1.0306	203.2	-10.0
0.0	552 4	1760.3	24.7974	31.59	242.04	273.64	0.1331	1.0192	273.2	0.0
10.0	1029-9	1748.0	13.7762	39.58	239.64	279.22	0.1636	1.0098	283.2	10.0
20.0	4975 4	1735.4	8.0450	47.63	237.24	284.87	0.1933	1.0025	293.2	20.0
20.0	7400 7	4722 3	4.8937	55.75	234.82	290.57	0.2222	0.9967	303.2	30.0
40.0	5070.1	1708.6	3.0888	63.95	232.39	296.33	0.2504	0.9925	313.2	40.0
			0.0455	72 24	220 07	302.14	0.2780	0.9895	323.2	50.0
50.0	8008.1	1694.5	2.0155	/2.21	227.73	202.14	0 3049	0.9876	333.2	60.0
60.0	12258.2	1679.9	1.3552	80.55	227.43	347.77	0.3047	D. 9866	343.2	70.0
70.0	18238.4	1664.6	0.9362	88.97	224.90	313.0/	0.3313	0.7000	353.2	80.0
80.0	26443.9	1648.8	0.6627	97.47	222.31	317.70	2 0 7824	0.7000	767 2	90.0
90.0	37458.6	1632.4	0.4795	106.05	219.66	325.71	U. 3824	0.7013	303.2	7010
400.0	E4070 L	1615 3	0.3539	114.71	216.93	331.65	0.4073	0.9887	373.2	100.1
100.0	70470 7	4597.5	0.2659	123.47	214.12	337.58	0.4318	0.9906	383.2	110.
110.0	70030.3	4678 9	0.2030	132.31	211.21	343.51	0.4558	0.9930	393.2	120.
120.0	77333.7	13/0.7	0.1573	141.24	208.18	349.42	0.4794	0.9958	403.2	130.0
N 130.0	124017.4	1227.0	0 1235	150.28	205.02	355.29	0.5027	0.9990	413.2	140.0
$\overset{\text{N}}{\circ}$ 140.0	160544.1	1537.4	0.1233	130120	÷					450.0
150.0	205144.9	1518.2	0.0781	159.42	201.71	361.13	0.5257	1.0024	423.2	150.0
160.0	258773.1	1496.1	0.0788	168.67	198.24	366.91	0.5484	1.0061	433.2	100.0
170.0	322672.6	1473.0	0.0639	178.03	194.58	372.62	0.5709	1.0099	443.2	170.0
480.0	709088 0	1448.7	0.0523	187.52	190.72	378.24	0.5930	1.0139	453.2	180.0
190.0	486325.9	1423.2	0.0431	197.15	186.63	383.78	0.6150	1.0180	463.2	190.0
	_		0 0757	201 02	482.28	789 20	П. 6368	1.0220	473.2	200.
200.0	588753.3	1396.3	0.0357	200.72	102.20	70/ 40	0 6586	1.0261	483.2	210.
210.0	706783.9	1368.0	0.0298	210.05	1//.04	279.97	0.0304	1 0301	493.2	220.
220.0	841898.6	1338.1	0.0250	226.94	1/2.69	377.03	0.0777	1 0340	503.2	230.0
230.0	995626.7	1306.4	0.0211	237.23	167.36	404.60	0.7014	4 0277	513.2	240.1
240.0	1169556.3	1272.7	0.0178	247.73	161.63	409.36	U. / 220	1.0377	31012	
250.0	1345334.9	1236.8	0.0151	258.48	155.42	413.90	0.7441	1.0412	523.2	250.
260.0	1584665.0	1198.4	0.0129	269.49	148.67	418.16	0.7656	1.0445	533.2	200.
270.0	4820749 8	1157.1	0.0109	280.82	141.28	422.11	0.7872	1.0473	543.2	270.
280.0	2404432 0	1112.4	0.0093	292.54	133.13	425.67	0.8071	1.0497	553.2	280.
290.0	2402014.0	1063.6	0.0079	304.71	124.05	428.76	0.8313	1.0516	563.2	290.1
			0.00/7	747 /4	117 80	631 36	0.8540	1.0526	573.2	300.
300.0	2733959.5	1007.8	0.006/	31/.40	404 00	432.20	0.8776	1.0525	583.2	310.0
310.0	3099044.5	949.5	0.0056	330.70	101.77	736.71 177 CL	0.0176	1.0509	593.2	320.
320.0	3499441.5	880.2	0.0047	345.50	0/.70	123.30	0.7020	1.0444	603.2	330.
330.0	3937433.3	796.8	0.0038	361.93	70.42	432.33	0.7277 N 0131	1.0365	613.2	340.
340.0	4415421.0	684.5	0.0029	381.78	45.29	421.21	9.7020	1.0303		
341.0	4465514-5	669.9	0.0028	384.42	41.87	426.29	0.9665	1.0347	614.2	341.
342 0	4514040 0	654.2	0.0027	387.01	38.13	425.14	0.9707	1.0327	615.4	374+

T	 TR	P	PR	Z(F)	Z(6)	FC(F)	FC(G)	B(F) KJ/KG	KJ/KG	DEG C
DEG C		PA								
			0.0000	0.0000	1.0000	1.0939	1.0000	0.10E+02	-0.61E+02	-40.0
-40.0	0.3761	2417 FO 8	0.0000	0.0000	0.9999	1.0086	0.9999	0.74E+01	-U.40ETUZ	-30.0
-30.0	0.3922	57.0	0.0000	0.0000	0.9999	1.0024	0.9999	0.52E+01	-U.38E+UZ	-20.0
-20.0	0.4083	134.4	0.0000	0.0000	0.9998	1.0005	0.9998	0.34E+01	-0.28E+02	-10.0
-10.0	0.4245	281.0	0.0001	0.0000	0 9996	0.9999	0.9996	0.20E+01	-0.19E+02	0.0
0.0	0.4406	552.1	0.0001	0.0000	0.7770			×		
				0.0000	0 0007	n. 9975	0.9993	0.94E+00	-0.11E+02	10.0
10.0	0.4567	1029.9	0.0002	0.0000	0.7773	0 0083	0.9989	0.17E+00	-0.31E+01	20.0
20.0	0.4728	1825.1	0.0004	0.0001	U.7707	0.0084	0.9982	-0.31E+00	0.43E+01	30.0
30.0	0.4890	3100.7	0.0007	0.0001	0.9962	0.7701	0.9973	-0.50E+00	0.11E+02	40.0
40.0	0.5051	5070.1	0.0011	0.0002	0.9973	0.77/7	0 0964	-0.42E+00	0.18E+02	50.0
50.0	0.5212	8008.1	0.0017	0.0003	0.9961	U.9962	0.7701			
2010		+(0 0045	-0.83E-01	0.24E+02	60.0
40.0	0.5374	12258.2	0.0026	0.0004	0.9945	0.9946	0.7773	0 505+00	0.31E+02	70.0
70 0	0.5074	18238.4	0.0038	0.0006	0.9924	0.9925	0.7729	0.175+01	0.37E+02	80.0
/0.0	0.5101	26663.9	0.0056	0.0009	0.9897	0.9899	0.9090	0.2/5+04	0 425+02	90.0
60.0	0.3070	77458 6	0.0079	0.0013	0.9864	0.9866	0.9866	0.242401	0 (85+02	100-0
90.0	0.5657	54979 4	0.0109	0.0017	0.9824	0.9827	0.9827	U.36E+U1	0.402402	10010
100.0	0.6019	51737.4	0.0107	1920 A					0.075+00	440.0
			0.0448	0.0023	0.9775	0.9781	0.9780	0.51E+01	U.53E+UZ	400.0
110.0	0.6180	70630.3	0.0140	0.0020	0 9719	0.9726	0.9726	0.68E+01	0.58E+U2	120.0
120.0	0.6341	94355 - 9	0.0198	0.0030	0.0452	0.9663	0.9663	0.87E+01	0.63E+D2	130.0
130.0	0.6503	124019.4	0.0260	0.0037	0.7034	n 9592	0.9592	0.11E+02	0.68E+02	140.0
N 140.0	0.6664	160599.1	0.0337	0.0050	U.75/0	0.0542	0.9511	0.13E+02	0.73E+02	150.0
N 150.0	0.6825	205144.9	0.0431	0.0064	0.9488	0.7512	91/244			
						0.0/07	0 9422	0.16E+02	0.78E+02	160.0
160.0	0.6986	258773.1	0.0543	0.0080	0.9390	0.9423	0.0724	0.18E+02	0.82E+02	170.0
470.0	0.7148	322672.6	0.0677	0.0077	0.9279	0.9324	0.7324	D 21E+02	0.87E+02	180.0
490.0	0 7700	398088-0	0.0836	0.0121	0.9156	0.9217	0.9210	0.216.02	0.91E+02	190.0
160.0		486325.9	0.1021	0.0147	0.9019	0.9101	0.9100	0.245+02	0.955+02	200.0
190.0	0.7470	F98757 7	0.1236	0.0178	0.8868	0.8976	0.8975	U.202702	0.732.02	
200.0	0.7632	200/23.3	0.1100			÷			0 405407	210 0
		70/787 8	0 4484	0.0213	0.8703	0.8843	0.8842	0.31E+02	0.100+03	220.0
210.0	0.7793	106/03.7	0.1707	0 0254	0 8523	0.8702	0.8701	0.35E+02	0.10E+U3	220.0
220.0	0.7954	841898.6	0.1/6/	0.0237	0 8724	0.8553	0.8553	0.39E+02	0.11E+03	230.0
230.0	0.8115	995626.7	0.2090	0.0302	0.0320	0 8397	0.8397	0.43E+02	0.11E+03	240.0
240.0	0.8277	1169556.3	0.2455	0.0357	0.0112	0.00775	0.8235	0.47E+02	0.11E+03	250.0
250.0	0.8438	1365334.9	0.2866	0.0421	0.7000	0.0200				
						0 0044	0 8066	0.52E+02	0.12E+03	260.0
260.0	0.8599	1584665.0	0.3326	0.0495	0.7627	0.0000	0.0000	0.57E+02	0.12E+03	270.0
270.0	D.8761	1829319.8	0.3840	0.0580	0.7352	0.7892	0.7072	0.62E+02	0.12E+03	280.0
280 0	0 8922	2101132.0	0.4410	0.0681	0.7053	0.7713	0.7539	0 485+02	0.13E+03	290.0
200.0		2402014-0	0.5042	0.0800	0.6726	0.7529	0.7527	0.002.02	0.13E+03	300.0
270.0		2722059 5	0.5739	0.0942	0.6364	0.7342	0.7342	0.742402	01102.00	
300.0	U.9245	2/33/3/19							0 475+07	310.0
		70000// 5	0 4505	0.1116	0.5961	0.7151	0.7151	0.806+02	0.132+03	320.0
310.0	0.9406	3077094.3	0.0305	0.1337	0.5500	0.6957	0.6956	0.87E+02	U.13E+U3	770 0
320.0	0.9567	3499441.5	0./340	0.1337	0 4954	0.6760	0.6760	0.96E+02	0.13E+03	330.0
330.0	0.9728	3937433.3	0.8265	0.1034	0 4047	0.4542	0.6562	0.11E+03	0.13E+03	340.0
340.0	0.9890	4415421.0	0.9268	U.2098	U.421/	0.0302	0.6542	0,11E+03	0.13E+03	341.0
341.0	0.9906	4465514.5	0.9373	0.2165	0.4121	0.0342	010076			
		1541010 0	0 9480	0,2238	0.4017	0.6522	0.6522	0.11E+03	0.13E+03	342.0
342.0) 0.9922	4516040.0	0.7400			Territoria de la compañía de la comp				

SATD COND (deg C) (kpa) (tr)	10K V(G) H(G) S(G) M3/KG KJ/KG KJ/KG.K	DEGREES OF SUPERHEAT 20K V(G) H(G) S(G) M3/KG KJ/KG KJ/KG.K	30K V(6) H(G) S(G) M3/KG KJ/KG KJ/KG.K	40K V(G) H(G) S(G) M3/KG KJ/KG KJ/KG.K
	B(G) FC(G) KJ/KG	B(G) FC(G) Kj/kg	B(G) FC(G) KJ/KG	B(G) FC(G) KJ/KG
(-40.0) (0.023) (0.3761)	533.571 257.31 1.1097 -62.4855 1.0000	555.515 262.68 1.1334 -64.1475 1.0000	577.459 268.13 1.1564 -65.5483 1.0000	599.404 273.65 1.1789 -66.7017 1.0000
(-30.0) (0.060) (0.3922)	212.212 262.68 1.0851 -49.8098 0.9999	220.596 268.13 1.1082 -51.2106 1.0000	228.979 273.65 1.1306 -52.3640 1.0000	237.363 279.24 1.1525 -53.2823 1.0000
(-20.0) (0.134) (0.4083)	98.155 268.12 1.0676 -39.1462 0.9999	101.885 273.65 1.0900 -40.2997 0.9999	105.616 279.24 1.1119 -41.2179 0.9999	109.347 284.90 1.1333 -41.9123 0.9999
(-10.0) (0.281) (0.4245)	48.740 273.64 1.0530 -29.3161 0.9998	50.525 279.24 1.0750 -30.2343 0.9998	52.311 284.90 1.0964 -30.9287 0.9998	54.096 290.63 1.1173 -31.4096 0.9998
(0.0) (0.552) (0.4406)	25.706 279.23 1.0411 -20.1707 0.9996	26.615 284.90 1.0625 -20.8651 0.9997	27.523 290.63 1.0834 -21.3460 0.9997	28.432 296.43 1.1038 -21.6229 0.9997
(^{**} 10.0) (1.030) (0.4567)	-14.263 284.88 1.0312 -11.5797 0.9994	14.751 290.62 1.0521 -12.0606 0.9994	15.238 296.42 1.0725 -12.3375 0.9995	15.725 302.28 1.0926 -12.4191 0.9995
(20.0) (1.825) (0.4728)	8.320 290.60 1.0234 -3.5425 0.9990	8.595 296.40 1.0438 -3.8194 0.9990	8.870 302.26 1.0638 -3.9010 0.9991	9.146 308.19 1.0834 -3.7955 0.9992
(30.0) (3.101) - (0.4890)	5.056 296.37 1.0172 4.0663 0.9984	5.218 302.24 1.0372 3.9848 0.9985	5.380 308.16 1.0568 4.0904 0.9986	5.542 314.15 1.0760 4.3754 0.9987
(40.0) (5.070) (0.5051)	3.188 302.20 1.0125 11.2943 0.9975	3.287 308.13 1.0321 11.4000 0.9977	3.387 314.12 1.0513 11.6851 0.9979	3.486 320.16 1.0701 12.1429 0.9980
(50.0) (8.008) (0.5212)	2.079 308.07 1.0091 18.1849 0.9964	2.141 314.06 1.0283 18.4703 0.9967	2.204 320.11 1.0471 18.9283 0.9969	2.267 326.22 1.0655 19.5522 0.9971
(60.0) (12.258)	1.396 313.98 1.0068	1.438 320.04 1.0256 25.2353 0.9953	1.479 326.15 1.0441 25.8597 0.9956	1.520 332.31 1.0622 26.6436 0.9959

APPENDIX G

ENTHALPY CONCENTRATION DIAGRAMS

NOTE: All isotherm temperatures are deg C.



ENTHALPY CONC DIAG R 22 -R 11 (PRESSURE= 578878.4 PA)

MASS CONCENTRATION OF R 22

224

ENTHALPY KJ/KG



MASS CONCENTRATION OF R 22





ENTHALPY CONC DIAG R 22 -R 11 (PRESSURE= 1451000.0 PA)

MASS CONCENTRATION OF R 22

226

ENTHALPY KJ/KG



ENTHALPY CONC DIAG R 12 -R 113 (PRESSURE= 739000.0 PA)

MASS CONCENTRATION OF R 12

ENTHALPY KJ/KG

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ENTHALPY CONC DIAG R 12 -R 113 (PRESSURE= 1190124.9 PA)

MASS CONCENTRATION OF R 12

228

ENTHALPY KJ/KG

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APPENDIX H

COMPUTER MODEL SAMPLE OUTPUTS

Figure	Referenced from section
H1 H2 H3 H4 H5 H6 H7 H8 H7 H8 H9 H10 H11 H12	3.2.1 3.2.2 3.2.3 3.2.4 3.2.5 3.2.6 3.2.7 3.2.8 3.2.9 3.2.9 3.2.10 3.3.1 3.4

***********	*******	************	************	*****			***********	
* R 11 -R * R 11 C * R 22 T * CONCENTRATI * RATIO OF FE * PROPORTION * DISTILLATE * FEED SUBCOO	22 ABSO HLORODIFI RICHLORO ON OF DI ED TO DI OF FEED SUPERHEA LING AT	RPTION REFRIG LUOROMETHANE FLUOROMETHANE STILLATE= 0. STILLATE= 4. FLOW THROUGH T AT EVAPORAT ABSORBER OUT=	ERATION SYSTEM (CHCLF2 (CCL3F 990 41 FEED PREHEATER= OR OUT= 0.0 0.0) +) + + + + 1 + 1 + 1 + 1 + 1 + 1 + 1 + 1 + 1 + 1	DATA AND UNITS T=TEMPERATURE DEG C P=PRESSURE KPA H=ENTHALPY KJ/KG B=AVAILABILITY KJ/KG	I T = 5.0 <<< I EVAP IN I IT = 5.0 I	* * * * * * * * * * * * * * * * * * *	<<< T= 24.0 I EVAP OUT I IT= 5.8 I
* DISTILLATE * APPROACH FL *APPROACH CON * ALL CONCENT *ENTROPY AND	SUBCOOLIN UID OFF N DENSER AN RATIONS N ENTHALPY	NG AT CONDENS EVAP TO EVAP ND ABSORBER= ARE MASS CONC DATUM TEMPER	ER OUT= 0.0 IN = 0.0 DE 0.0 0.0 DEG S OF R 11 ATURE = -40.0	DEG + 6 + + 10EG C+	IEXPANSION INI	P= 578878.4 H= 80.71 B= 63.81 STATE=2 PHA 	/\ \/ /\ \/ *** \/ *E* \/	P= 578878.4 H= 252.98 B= 50.62 STATE=SATDV
*AVBTY DATUMS	: TEMP=	30.00EG C :P	RESSURE= 101.	3 KPA+	P= 1183366.0 H= 80.71 B= 65.56 STATE=SATDL	2		
	0.0	L COND OUT						
	•	IT= 30.0	i					
		IP= 1183366 IH= 80.71	.01					
		18= 65.56	1		÷			
		ISTATE=SATD						
8 I		>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>	>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>	·>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>	››››››››››››››››››››››››››››››››››››››	››››››››››››››››››››››››››››››››››››››		*
T= 30.0 >>	> ====	*** ND= >>>	14 E			B	V.	2
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I COLUMN FE	ED1 #	* *******	****		IT= 29.9 I		V	
IT= 29.9	1 ++++	*** * FEE	D +	uuu	IP= 1183366.01			21
IP= 1183366 IH= 62.05	.UI +FE	ED#<<<;# =BOO ### # HEAT	ER *	· · · · ·	IB= 9.41 i			

V HT= 29.9 77 1 \mathbf{N} IP= 1183366.01 $\boldsymbol{\Lambda}$ I BOT IN EXP I ം 🗸 IH= 62.05 I ----- /\ MIX IN |T= 101.1 | ~ / / IFEED OUT P/HI /\ 18= 9.41 I \mathbf{V} IT= 5.8 ISTATE=SUBCL I IP= 1183366.01 IT= 29.9 1/\ IP= 578878.41 ×7 IH= 124.41 IP= 1183366.01 /\ -----IH= 252.98 18= 5.82 ~// - 1 [H= 62.05 1 /\ ****** 1B= 50.62 +<<<<<<<<<<<<< \mathbf{N} IB= 9.41 | <<<+ ******** ISTATE=SATDV I /\ -----+FEED + ISTATE=SUBCL | . *** *PRE * - 22 · · · · · · 1 *** • -********* - 7 \ ****** 1 ----------**** ---- /\ _____ I MIX OUT IBOT OUT EXP I V I BOT OUT P/HI /\ 1 BOT IN P/H I Λ IT= 74.0 1 - \/ 11= 52.2 IT= 101.1 // 1T= 101.1 I 1P= 578878.41 1P= 578878.41 \/ |P= 1183366.01 /\ IP= 1183366.01 1 IH= 153.55 [H= 124.41 1 - \/ 1H= 124.41 | /\ iH= 124.41 - \/ IB= 9.67 |B= 6.89 ____L 18= 5.82 -118= 5.82 I 1 ISTATE=2 PHA I ISTATE=2 PHA I - \ / ISTATE=SATDL / // ISTATE=SATDL | 1 14 ----------- \mathbf{N} ------1 Λ \mathbf{V} \mathbf{X} ----------- \mathbf{V} Λ $\overline{\Lambda}$ I ABSORBER INI \mathbf{N} 1 7 IT= 52.2 I ~/ 1 IP= 578878.41 **\/** 71 1 IH= 153.55 \mathbf{N} 7 \/ 1B= 9.67 - 1 1 \mathbf{V} ISTATE=2 PHA I 1 -----------×7 1 I BOTTOM OUT I ************ 1 ****** Λ IT= 101.1 1 . 1 IP= 1183366.01 +BOIL+ -73 T= 30.0 >>> * ABSORBER * >>> Λ Λ IH= 124.41 ***** *TSAT= 30.0* IB= 5.82 $\mathbf{V} \approx$ - 1 ********** Λ \mathbf{V} 'ISTATE=SATDL | IABSORBER OUTI \mathbf{V} 1 >>>>>>> 1T = 30.0 \sim \sim 1P= 578878.41 \mathbf{V} 1 1H= 62.05 I \mathbf{N} ************************* \mathbf{V} 1B= 8.98 *COLUMN PLATE NUMBERS: TOTAL=13: RECTIFYING= 7: STRIPPING= 3 * 1 ISTATE=SATDL | \mathbf{V} *RECTIFYING HX TOTAL= 0.000KW: STRIPPING HX TOTAL= 0.000KW * $\mathbf{\Lambda}$ \mathbf{N} *FEEDPLATE: TEMP= 56.30 DEG C: LIQUID CONC= 0.322: VAP CONC= 0.739* 1 \mathbf{V} 1 *ABSOLUTE REFLUX FLOW TO DISTILLATE FLOW RATIO= 1.70 \mathbf{N} 1 *ACTUAL REFLUX FLOW TO MINIMUM REFLUX FLOW RATIO= 1.17 I PUMP IN \mathbf{V} IT= 30.0 \mathbf{V} I EVAPORATOR I CONDENSER I ABSORBER I BOILER I PUMP I BOOST* 1 IP= 578878.41 \mathbf{N} 1 --------IH= 62.05 \mathbf{V} 123.434 131.306 | 0.1071 0.000+ 117.872 KW | 10.000 18= 8.98 \mathbf{X} Λ \mathbf{V} ISTATE=SATDL $\mathbf{\Lambda}$ FEED 1 . 1 ******** \mathbf{V} 1--------------# PUMP # IDISTILLATEI BOTTOMS | P/HEAT | COLUMN/HXI TOTAL * \sim I TUO 9MU9 I ******** ------- Λ IT= 29.9 | 0.0000 | 0.0000 | 0.2561* Λ \mathbf{V} 1 0.0580 I 0.1981 *FLOWS KG/S \mathbf{N} [P= 1183366.0] Λ ■CONCENTRATION 1 0.990 1 0.050 1 0.263 IH= 62.05 I \mathbf{X} Λ 1B= 9.41 * MINIMUM REFLUX FLOW (KG/S) = 0.0349 ISTATE=SUBCL I ********************************** *********** 5.0, 30.0 & 101.1 DEG C 2.114 BASED ON *CARNOT COP= PUMP KW AS %= 0.3 +COP= 0.318 COPHEAT= 1.315

FIGURE H1

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NO	I TEMP I DEG C	і к і у/х	IHEAT TEI I KW I	FEED FLOW KG/S	IIA/BY CH I KJ/S	I PHAS	EI MASS I CONC	IENTHALP	YIENTROPY IKJ/KG K	IAVAIL/T	YIFUG COEF	IFUG COEF	I FLOW I KG/S
1	1 30.0 1	1 1.01	17.872	0.0000	1 -0.006	I LIQ	10.990	80.71 261.28	0.327 0.000	1 65.56	1 0.847 1 0.000	1 0.001 1 0.000	I 0.0990
2	30.7	1.03	0.000	0.0000	-0.013	I LIQ I VAP	1 10.962 10.990	80.75 261.31	 0.334 0.922	 61.85 65.62	1 1 0.859 1 0.844	 0.187 0.720	 0.0410 0.0990
3	31.7	1.07	1 0.0001	0.0000	 -0.012 	LIQ VAP	 0.917 0.978	 80.82 261.19	 0.342 0.925	1 56.69 64.21	1 1 0.881 1 0.845	 0.185 0.731	 0.0408 0.0990
4	33.5	 1.13 	1 0.0001 1 1	0.0000	 -0.014	LIQ VAP	 0.846 0.960	 80.98 261.50	 0.352 0.929	 49.54 62.21	 0.921 0.848	0.183	0.0406
5 1	36.4	1 1.26	1 0.0001	0.0000	 -0,019 	LIQ VAP	 0.740 0.931	 81.37 262.01	1 1 0.363 1 0.934	 40.14 59.42	1 1 0.994 1 0.852	I I 0.183 I 0.738	0.0402
6	40.9	1 1.47	0.0001	0.0000	 -0.024		1 10.604 10.888	 82.42 262.79	1 0.373	1 29.43	1 1.123	0.188	0.0378
7	46.7	1 1.77	 0.000	0.0000	-0.024		I I0.470	1 1 84.64	I I 0.381	20.20	1 1.315	0.203	0.0397
8 I 1 8	52.3	1 2.08	1 0.0001	0.0000	-0.014		I 10.375	1 87.52		1 14.46	1 1.519		0.0399
91	56.3	1 2.30	0.000	0.2561	1 -0.7341		I 10.322	87.89	0.750	40.28	1 1.671	0.760	0.3367
0 i	59.1	2.45	0.000	0.0000	-0.068	LIQ	10.734	91.67	0.396	46.02	0.878 1.781	0.766 0.259	0.0980
1	66.7	2.85	0.0001	0.0000	-0.4161	LIQ	10.711 1 10.221	266.44	0.953 0.406	44.52 7.16	0.882 2.075	0.769 0.306	0.1386 0.3475
2	82.3	3.58	1 0.0001	0.0000	-1.106	VAP	10.629 10.125	267.70	0.952 0.428	40.74 4.87	0.891 2.771	0.778 	0.1410
3 1	101.1	4.22	 -31.306	0.0000	4.9981	VAP	10.447 0.050	270.06	0.939 0.457	34.97		0.793	0.1495
1					1	VAP	10.211 	273.25	0.900	32.89	0.939	0.811	0.1717

FIGURE H1 CONTD

SECOND LAW ANALYSIS	KJ/S	PERCENT OF	AVAILABILITY	LOSS
		2 9		
PIPEWORK	0.000	0.000		
CONDENSER DUI TO FRECOULER IN	0.000	0.000		
PRECOULER LIQ OUT TO EXPANSION IN	0.000	0.000	13	
EVAPORATOR OUT TO PRECODER IN	0.000	0.000		
PRECOULER VAP OUT TO MIXING IN	0.000	0.000		
MIXING OUT TO ADSURDER IN	0,000	0.000		
ABSORBER OUT TO FORF IN	0,000	0.000		
PUMP OUT TO PECU SPLIT	0.000	0.000		
FEED SPLIT TO FREMENICK IN	0.000	0.000		
PREMEATER OUT TO FEED JOIN	0.000	0.000		
FEED SPLIT TO RECTIFIER HA IN	0.000	0.000		
RECITFIER HX OUT TO FEED JOIN	0.000	0.000		
BOILOW OUL OF COLUMN TO HA IN	0.000	0.000		
BOTTOM HX OUT TO EXPANSION IN	0.000	0.000		
		÷.		
EXPANSION				
DISTILLATE EXPANSION	-0.101	1.6/4		
BOTTOMS EXPANSION	0.212	-3.499		
PUMP	0.109			
ANTERNAL MEAT EXCUANCERS		-		
INTERNAL HEAT EXCHANGERS	0.000	0.000		
PRECODLER	0.000	0.000		
PREHEATER		-	24	
EXTERNAL HEAT EXCHANGERS			9	
EVAPORATOR	0.102	-1.689		
EVAP REFRIGERANT STREAMS ONLY -0.766				÷.
EVAP CARNOT -0.868		14.328		
ABSORBER	-2.003	33.060		
CADNOT COM AUERAGE TEMP 0.000				
CARNOT FROM AVERAGE TELL OTOGO			÷ .	
COLUMN				
	5,949			
COLUMN ONLY = 210007	-3.394	56.028		
(MINIMUM SEPERATION WORK= 2.549)				
CONDENSER	0.000	0.000		
COND CARNOT	0.000	0.000		
CONDENSER STREAMS ONLY	-0.006	- 0.077		
	0.000			
		27		

FIGURE H1 CONTD

 R 11 -R 22 R 11 CHLO R 22 TRIC CONCENTRATION RATIO OF FEED PROPORTION OF DISTILLATE SUB APPROACH FLUID *APPROACH CONDEN: *ALL CONCENTRAT 	ABSORPTION REFRIGERATION SYSTEM RODIFLUOROMETHANE (CHCLF2) HLOROFLUOROMETHANE (CCL3F) OF DISTILLATE= 0.990 TO DISTILLATE= 4.41 FEED FLOW THROUGH FEED PREHEATER= 1.000 G AT ABSORBER OUT= 0.0 DEG G AT ABSORBER OUT= 0.0 DEG COOLING AT CONDENSER OUT= 0.0 DEG OFF EVAP TO EVAP IN = 0.0 DEG SER AND ABSORBER= 0.0 0.0 DEG HAND ABSORBER= 0.0 0.0 DEG ABSOR AT IN TEMPERATURE = -60 0000 G	I DATA AND UNITS I IT=TEMPERATURE DEG C I IP=PRESSURE KPA I IH=ENTHALPY KJ/KG I IB=AVAILABILITY KJ/KGI	T= 5.0 <<< I EVAP IN I IT= 5.0 I IP= 578878.41 IH= 80.71 I IB= 63.81 I ISTATE=2 PHA I	* * * * * * * * * * * * * * * * * * *	<<< T= 24.0 I EVAP OUT I IT= 5.8 IP= 578878.41 IH= 252.98 IB= 50.62 ISTATE=SATDV I
*AVBTY DATUMS: TI	EMP= 30.0DEG C :PRESSURE= 101.3 KPA	IP= 1183366.01 IH⇒ 80.71 I IB= 65.56 I ISTATE≠SATDL I		**** \/ /\ \/ /\ \/ /\ \/	
	I COND OUT I IT= 30.0 I IP= 1183366.0I IH= 80.71 I IE= 65.56 I ISTATE=SATDL I		8		
5	>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>		·····		8 11
T= 30.0 >>>	*COND* >>> ****** * * * *	e.	ю		
					, ,
	· · · · · · · · · · · · · · · · · · ·				
I COLUMN FEEDI IT= 66.4 I IP= 1193366.01 IH= 113.42	<pre>* * * * * * * * * * * * * * * * * * *</pre>	I FEED I= 66.4 P= 1183366.0 H= 113.48 () B= 11.46			0.5
18= 113.46 18= 11.46 1STATE=2 FHA	****** * OEMIGG * * * *#******* * *	//ISTATE=2 PHA 1 //		\/ \/	

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234

+>

IP= 1183366.01 /\ |H= 113.48 | /\ ******* IB= 11.46 | <<<+ ISTATE=2 PHA 1 -----------* $\mathbf{\Lambda}$. \mathbf{N} ٠ 1 BOT IN P/H I 1 . |T= 101.1 | 4 \sim |P= 1183366.01 Λ ٠ |H= 124.41 | $\overline{}$. 18≓ 5.82 I $\overline{}$ ISTATE=SATDL I Λ ----- Λ $\overline{}$ Λ $\overline{}$ 73 7 Λ -7 _____ I BOTTOM OUT I 1 1 IT= 101.1 | ****** 1P= 1183366.01 +BOIL+ 1 [H= 124.41 ! ###### 1 VZ. \mathbf{N} 18= 5.82 1 1 ISTATE=SATDL | \mathbf{V} _____ >>>>>> ************************* *COLUMN PLATE NUMBERS: TOTAL=12: RECTIFYING= 7: STRIPPING= 2 * *RECTIFYING HX TOTAL= 0.000KW: STRIPPING HX TOTAL= 0.000KW * *FEEDPLATE: TEMP= 68.30 DEG C: LIQUID CONC= 0.209: VAP CONC= 0.611* *ABSOLUTE REFLUX FLOW TO DISTILLATE FLOW RATIO= 1.99 +ACTUAL REFLUX FLOW TO MINIMUM REFLUX FLOW RATIO= 1.15

120.913

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*CONCENTRATION 1 0.990 1 0.050

*COP= 0.470 COPHEAT= 1.465

* MINIMUM REFLUX FLOW (KG/S) = 0.0501

I EVAPORATOR I CONDENSER | ABSORBER | BOILER | PUMP | BOOST+

1

IDISTILLATE: BOTTOMS | P/HEAT | COLUMN/HXI TOTAL +

1

PUMP KW AS %= 0.5

110.263 121.178 | 0.107| 0.000+

FEED

|-----

0.263

FLOWS KG/S / 0.0580 / 0.1981 / 0.2561 / 0.0000 / 0.2561

*CARNOT COP= 2.114 BASED ON 5.0, 30.0 & 101.1 DEG C

I BOT IN EXP I 17 IH= 62.05 I I MIX IN I IB= 9.41 I IT= 29.9 17 IT= 5.8 I IP= 1183366.01 \mathbf{V} ISTATE=SUBCL I IP= 578878.41 11 -----IH= 57.91 | \mathbf{V} IH= 252.98 I 18= 2.88 - E *<<<<<<<<I>STATE=SUBCL1 \Z 18= 50.62 ********* ISTATE=SATDV I /\ ------*** - 73 *** . - 44 Λ ******** ---- // _____ ×7 I TUO X1M I IBOT OUT EXP I \/ I BOT OUT P/HI /\ IT= 39.1 17= 30.1 I \/ IT= 29.9 1/\ 1P= 578878.41 \/ 1P= 578878.41 IP= 1183366.01 /\ IH= 57.91 | \/ IH= 102.12 | IH= 57.91 | /\ 18= 8.25 18= 2.48 | \/ - 1 IB= 2.88 I/\ ISTATE=SUBCL I \/ ISTATE=2 PHA I ISTATE=SUBCL | /\ -----------×7 ---- // ŃZ Λ 1 \mathbf{V} 11 ABSORBER INI 1 IT= 39.1 I \mathbf{X} 1 IP= 578878.41 ~// \mathbf{V} IH= 102.12 | 73 18= 8.25 1 Λ \mathbf{V} ISTATE=2 PHA 1 \mathbf{V} 7 1/ 2 77 *********** \mathbf{N} Λ T= 30.0 >>> + ABSORBER + >>> 1 *TSAT= 30.0* 1 ************ \sim IABSORBER OUTI \mathbf{V} 1 XZ IT= 30.0 I $\langle \Lambda \rangle$ ĪΛ. |P= 578878.41 $\overline{}$ IH= 62.05 Λ \mathbf{N} 1B= 8.98 \mathbf{V} $\langle \Lambda \rangle$ ISTATE=SATOL I Λ \mathbf{V} ----- \mathbf{V} 1 \mathbf{V} 1 1 \mathbf{V} I PUMP IN \mathbf{V} ≈ / \ IT= 30.0 \mathbf{N} 1P= 578878.41 Λ \mathbf{V} \mathbf{V} IH= 62.05 1 IB= 8.98 M 1 ISTATE=SATDL I \mathbf{V} Λ Λ ******** PUMP + \sim BARRANA I PUMP OUT I \sim IT= 29.9 Λ \mathbf{V} \mathbf{M} IF= 1183356.01 $\langle \Lambda \rangle$ \mathbf{V} IH= 62.05 I Λ 18= 9.41 - 1 ISTATE=SUBCL | -----

11- 1100000.01

1.1

*FEED *

.

*PRE *

.

---- //

IFEED OUT P/HI /\

IT= 66.4 1 /\

235

* KW | 10.000

FIGURE H2

		0.0000 0.0000 0.0000 0.0000	-0.007 -0.020 -0.028 -0.028	LIQ VAP LIQ VAP LIQ VAP	10.990 10.997 10.962 10.990 10.908 10.908	80.71 261.28 80.75 261.31 80.84 261.43	0.327 0.000 0.334 0.922 0.344	65.56 0.00 61.85 65.62	1 0.847 1 0.000 1 0.859 1 0.844 1 0.886	0.001 0.000 0.187 0.720	0.1158 0.0000 0.0577 0.1158
	0.000 0.000	0.0000 0.0000 0.0000	-0.020	LIQ VAP LIQ VAP	10.962 10.990 1 10.908 10.976	80.75 261.31 80.84 261.43	1 0.334 1 0.922 1 1 0.344	61.85 65.62	0.859 0.844	0.187	0.0573 0.1158
	0.0001	0.0000 0.0000	-0.028 -0.047	LIQ VAP	10.908 10.976	1 1 80.84 1 261.43	0.344	55.72	0.886	0.185	1
5 1.18 5 1.41		0.0000		1	1		I U1740	63.94	1 0.846	0.731	0.1157
1 3 1.41	1 1		1	LIQ VAP	10.806 10.949	 81.10 261.61	 0.357 0.931	45.83 61.13	 0.946 0.849	0.183	 0.0570 0.1150
E	1 0.0001	0.0000	-0.085 -0.085	LIQ VAP	 0.636 0.899	 82.10 262.70	 0.371 0.939	31.83 56.67	 1.088 0.857	 0.186 0.743	0.0561
i 1.87		0.0000	 -0.115 	LIQ VAP	 0.437 0.816	 85.48 264.59	 0.383 0.948	18.12 50.62	 1.377 0.868	1 0.209 1 1 0.755 1	0.0561
1 2.41	1 0.000	0.0000	 -0.076 	LIQ VAP	1 10.298 10.718	 91.20 266.32	1 1 0.395 1 0.953	10.46	1 1.752 0.881	0.255 0.768	0.0571
2.76	1 0.0001	0.0000	-0.030 -0.030	LIQ VAP	 0.234 0.647	 95.76 267.55	I I 0.403 I 0.952	7.66	1 1 2.024 1 0.889	 0.295 0.776	0.0584
1 2.93	0.0001	0.2561	-0.275		10.209	 98.11 267.86	 0.408 0.951	6.74	I I 2.161 I 0.893	0.316 0.779	0.2916
3.21	0.0001	0.0000	-0.192	LIQ	1 10.169 10.544	1 102.47	1 1 0.416 1 0.948	5.61	1 2.410	0.357 0.785	0.2982
i i 3.71	0.0001	0.0000	-0.5391		10.109	 111.55 270.53	1 0.433 1 1 0.933 1	4.80	1 2.921 0.916	0.457	0.3133
4.22	-21.178	0.0000	3.4391		10.050	 124.42 273.24		5.82	1 3.630	0.617	0.1981
	5 1.87 5 2.41 1 2.76 1 2.76 1 3.21 1 3.71 1 4.22 CH is	5 1.87 0.000 2.41 0.000 2.76 0.000 3.21 0.000 3.21 0.000 3.71 0.000 4.22 -21.178 4.22 -21.178	1.87 0.000 0.0000 2.41 0.000 0.0000 2.76 0.000 0.0000 2.76 0.000 0.0000 2.76 0.000 0.0000 3.21 0.000 0.0000 3.71 0.0000 0.0000 4.22 -21.178 0.0000 4.22 -21.178 0.0000	1.87 0.000 0.0000 -0.115 2.41 0.000 0.0000 -0.076 2.76 0.000 0.0000 -0.030 2.76 0.000 0.0000 -0.030 3.21 0.000 0.2561 -0.275 3.71 0.000 0.0000 -0.192 4.22 -21.178 0.0000 3.439	1.87 0.000 0.0000 -0.115 LIQ 2.41 0.000 0.0000 -0.076 LIQ 2.76 0.000 0.0000 -0.030 LIQ 2.76 0.000 0.0000 -0.030 LIQ 3.21 0.000 0.0000 -0.172 LIQ 3.71 0.000 0.0000 -0.5391 LIQ 4.22 -21.178 0.0000 3.439 LIQ VAP VAP VAP VAP VAP	1.87 0.000 0.0000 -0.1151 LIQ 0.437 2.41 0.000 0.0000 -0.0761 LIQ 0.278 2.76 0.000 0.0000 -0.030 LIQ 0.234 2.76 0.000 0.0000 -0.030 LIQ 0.234 2.76 0.000 0.2561 -0.2751 LIQ 0.209 3.21 0.0001 0.0000 -0.1921 LIQ 0.167 3.71 0.0001 0.0000 -0.5391 LIQ 0.1647 4.22 -21.1781 0.0000 3.4391 LIQ 0.050 VAP 0.211 1 0.0200 3.4391 LIQ 0.050	1.87 0.000 0.0000 -0.115 LIQ 10.437 85.48 2.41 0.000 0.0000 -0.076 LIQ 10.278 91.20 2.76 0.000 0.0000 -0.030 LIQ 10.234 95.76 2.76 0.000 0.0000 -0.030 LIQ 10.234 95.76 2.73 0.000 0.2561 -0.275 LIQ 10.209 98.11 VAP 10.647 267.55 10.000 0.0000 -0.192 LIQ 10.209 98.11 VAP 10.611 267.86 10.001 0.0000 -0.192 LIQ 10.247 VAP 10.611 267.86 10.000 -0.5391 LIQ 102.47 VAP 10.0001 0.0000 -0.5391 LIQ 10.169 102.47 VAP 10.0001 0.0000 -0.5391 LIQ 10.109 111.55 14.22 -21.1781 0.0000 3.4391 LIQ 10.050 124.42 14.22 -21.1781 0.0000 3.4391 LIQ 1	1.87 0.0001 0.0000 -0.1151 LIQ 10.437 85.48 0.383 2.41 0.0001 0.0000 -0.0761 LIQ 10.278 91.20 0.395 2.76 0.0001 0.0000 -0.0761 LIQ 10.278 91.20 0.395 2.76 0.0001 0.0000 -0.0301 LIQ 10.234 95.76 0.403 2.76 0.0001 0.0000 -0.0301 LIQ 10.2091 98.11 0.403 2.73 0.0001 0.2561 -0.2751 LIQ 10.2091 98.11 0.408 3.21 0.0001 0.0000 -0.1921 LIQ 10.1691 102.471 0.416 3.71 0.0001 0.0000 -0.5391 LIQ 10.1091 111.55 0.433 4.22 -21.178 0.0000 3.4391 LIQ 10.0501 124.421 0.457 VAP 10.2111 273.24 0.900 124.421 0.457 10.900	1.87 0.000 0.0000 -0.1151 LIQ 10.437 85.48 0.383 18.12 2.41 0.000 0.0000 -0.0761 LIQ 10.298 91.20 0.395 10.46 2.76 0.000 0.0000 -0.0761 LIQ 10.234 95.76 0.403 7.66 2.76 0.000 0.0000 -0.0751 LIQ 10.209 98.11 0.403 7.66 2.73 0.000 0.2561 -0.2751 LIQ 10.209 98.11 0.408 6.74 VAP 10.647 1267.86 0.951 40.03 3.21 0.0001 0.0000 -0.1921 LIQ 10.167 102.47 0.416 5.61 3.71 0.0001 0.0000 -0.5391 LIQ 10.107 111.55 0.433 4.80 4.22 -21.1781 0.0000 3.4391 LIQ 10.0501 124.42 0.4571 5.82 VAP 10.211 273.24 0.900 32.89 14.14 14.22 14.571 5.82	1.87 0.000 -0.1151 LIQ 10.437 85.48 0.383 18.12 1.377 8 2.41 0.000 0.0000 -0.0761 LIQ 10.298 91.20 0.395 10.46 1.752 9 2.41 0.000 0.0000 -0.0761 LIQ 10.298 91.20 0.395 10.46 1.752 9 2.76 0.0001 0.0000 -0.0301 LIQ 10.234 95.76 0.403 7.66 2.024 9 10.647 267.55 0.952 41.52 0.889 1 2.73 0.000 0.2561 -0.2751 LIQ 10.207 98.11 0.408 6.74 2.161 1 VAP 10.647 267.55 0.952 41.52 0.889 3.21 0.0001 0.2561 -0.2751 LIQ 10.247 0.416 5.61 2.410 1 0.0000 -0.1921 LIQ 10.169 112.47 0.416 5.61 2.410 1 0.0000 -0.5391 LIQ 10.109 <	1.87 0.000 0.0000 -0.1151 LIQ 10.437 85.48 0.383 18.12 1.377 0.209 2.41 0.000 0.0000 -0.0761 LIQ 10.298 91.20 0.395 10.461 1.752 0.255 2.41 0.000 0.0000 -0.0761 LIQ 10.298 91.20 0.395 10.461 1.752 0.255 2.41 0.0001 0.0000 -0.0761 LIQ 10.298 91.20 0.395 10.461 1.752 0.255 2.76 0.0001 0.0000 -0.0301 LIQ 10.234 95.76 0.4033 7.66 2.024 0.295 3.273 0.0001 0.2561 -0.2751 LIQ 10.207 98.11 0.408 6.74 2.161 0.316 VAP 10.611 267.86 0.951 40.03 0.893 0.779 3.21 0.0001 0.0000 -0.1921 LIQ 10.167 102.47 0.416 5.61 2.410 0.359 3.71 0.0001 0.0000 -0.5391 <t< td=""></t<>

FIGURE H2 CONTD

ŧá	SECOND LAW ANALY	'SIS K	J/S	PERCENT OF	AVAILABILITY	LOSS
PTPEWORK				0.000		
CONDENS	ER OUT TO PRECOOLER IN		0.000	0.000		
PRECOOL	ER LIQ OUT TO EXPANSION	I IN	0.000	0.000		
FVAPORA	TOR OUT TO PRECOOLER IN	1	0.000	0.000		
PRECOOL	ER VAP OUT TO MIXING IN	1	0.000	0.000		
MIXING	OUT TO ABSORBER IN		0.000	0.000		
ABSORBE	R OUT TO PUMP IN		0.000	0.000		
PUMP OU	T TO FEED SPLIT		0.000	0.000		
FEED SP	LIT TO PREHEATER IN		0.000	0.000		
PREHEAT	ER OUT TO FEED JOIN		0.000	0.000		
FEED SP	LIT TO RECTIFIER HX IN		0.000	0.000		
RECTIFI	ER HX OUT TO FEED JOIN		0.000	0.000		
BOTTOM	OUT OF COLUMN TO HX IN		0.000	0.000		
ROTTOM	HY OUT TO PREHEATER IN		0.000	0.000		
PREHEAT	ER OUT TO EXPANSION IN		0.000	0.000		
FIPANSION		•		2 454		
DISTILL	ATE EXPANSION		-0.101	4 544		
BOTTOMS	EXPANSION		-0.080	1.744		
			0.109			τi.
	AT SYCHANGERS					
INTERNAL HE	AI ELCHANGERS	10 m	0.000	0.000		
PRECOOL	ER		-0.057	1.379		
CATERNAL HE	AT FICHANGERS					
EVAPORATOR			0,102	-2.475		
EVAP RE	FRIGERANT STREAMS ONLY	-0.766		20 999		
EVAP CA	RNOT -0.868			201777		
		5				8
ABSORBER	AND MIXING REFRIGERA	NT STREAMS ONLY	-1:129	27.308		
CARNOT	FROM AVERAGE TEMP O	.000		8		
COLUMN						
CARNOT	ONLY		4.024			
(STREAN COLUMN (MINIMU	IS ONLY = 2.031 J ONLY=CARNOT-STREAMS IM SEPERATION WORK=	2.024)	-1.993	48.224		
CONDENSER			0.000	0.000	0	
COND C	ARNOT		-0.007	0.167		
CONDEN	SER SIREANS UNLT					
		a antista anna 2016 anna 2016 anna 2016 anna 2016 anna 2016 anna 2016 anna 2016 anna 2016 anna 2016 anna 2016 a	0.000			
		-				

FIGURE H2 CONTD

<pre>* R 11 -R 22 * R 11 CHLOR * R 11 CHLOR * R 22 TRICH * CONCENTRATION OF * RATIO OF FEED T * PROPORTION OF F * DISTILLATE SUPE * FEED SUBCOOLING * DISTILLATE SUBC * APPROACH FLUID * APPROACH FLUID * APPROACH CONDENS * ALL CONCENTRATI * ENTROPY AND ENTH * AVBTY DATUMS: TE</pre>	ABSORPTION REFRIGERATION SYSTEM RODIFLUOROMETHANE (CHCLF2 ALOROFLUOROMETHANE (CCL3F OF DISTILLATE= 0.990 TO DISTILLATE= 4.41 FEED FLOW THROUGH FEED PREHEATER= RHEAT AT EVAPORATOR OUT= 0.0 COLING AT CONDENSER OUT= 0.0 OFF EVAP TO EVAP IN = 0.0 DEG SER AND ABSORBER= 0.0 0.0 DEG ONS ARE MASS CONCS OF R 11 TALPY DATUM TEMPERATURE = -40.0D MP= 30.0DEG C :PRESSURE= 101.3	<pre></pre>	I T= 5.0 (((I T= 5.0 (((I T= 5.0 I I T= 5.0 I I F= 579255.21 I H= 64.22 I I B= 66.38 I I STATE=2 PHA I	<pre>EVAPORATOR TEA(= 5.8 /\ \/ /\ \/ /\ \/ *** \/ *E* \/ *** \/</pre>	<pre><< T= 24.0 i EVAF OUT i IT= 5.8 IP= 579255.2 IH= 252.99 IB= 50.64 ISTATE=SATDV I </pre>
******	I COND OUT I IT= 30.0 I IP= 1183366.01 IH= 80.71 I IB= 65.56 I ISTATE=SATOL I	IN- 04.22 1 IB= 67.17 I ISTATE=SUBCL I	IP/COOL L OUTI IT= 17.2 I IP= 1183366.01 IH= 64.22 I IB= 67.17 I ISTATE=SUBCL I	/\ \/ /\ \/ * * * * * PRECOOLER * * *	IP/COOL V IN I IT= 5.8 I IP= 579255.21 IH= 252.99 I IB= 50.64 I ISTATE=SATDV I
T= 30.0 >>>		*****	<pre>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>></pre>		IP/COOL V OUTI IT= 30.0 1 IP= 579255.21 IH= 269.47 1 IB= 47.56 1 ISTATE=SUPHV 1
1 COLUMN FEEDI IT= 66.4 I IP= 1183366.01 IH= 113.46 I	* * * * * * * * * * * * * * * * * * *	FEED 			

	• • • • • •	8	IFEED OUT P/HI /	T= P= H= B=	27.7 1183366.0 62.06 7.42	BOT IN T= 29.	I EXP I 9 I		I MLX IN I 17= 30-0	
			IT= 66.4 1 // IP= 1183366.01 // IH= 113.46 1 //		ATE=SUBCL T	iH= 57. (B= 2.	91 I 88 I	Ň.	1P= 579255.21 1H= 269.47	
	* *		IB= 11.47 <<< ISTATE=2 PHA	* *< *FEED * *PRE *		/\ /\ -++		*******	ISTATE=SUPHV I	
	• • >>	>>>>>>>>	·>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>	+HEAT +>	>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>	/\>>>>#E	*>>>>>>>>>	+ XIN +<<<		
	* * /\			÷ ÷	2.	<u>, , , , , , , , , , , , , , , , , , , </u>	i i i	* *		
				******				\/		
	* * /\		I BOT IN P/H I	1	BOT OUT P/HI	<u></u>	IBOT OUT	EXP 1 V/	I MIX OUT I	
	+ + /\		IT= 101.1 l	1	T= 29.9 I	/\	IT= 30.1	1 I V	1T= 40.1 1	
	+ + /\		IP= 1183366.01	1	P= 1183366.01	Λ	1P= 5792	255.21 \/	P= 5/9255+2 H= 105 93	
			H= 124.41 B= 5.82	1	H= 57.71 B= 2.88		18= 2.4	48 \/	18= 8.26 I	
			ISTATE=SATDL I	~ i	STATE=SUBCL I		ISTATE=SU	UBCL I V/	ISTATE=2 PHA I	
,	• • A					Λ		\/		
:	* * /\				*	Λ				
								Ň.	ABSORBER INI	
•						X		V/	IT= 40.1 I	
	• • · · · ·					Λ		M	IP= 579255.21	
	• • /\							N/	1H= 185.93 I	
+	► + /\							10	ISTATE=2 PHA I	
								Ň.		
IT= 101.1	* * /\					Λ.		***********	H# 39	
(P= 1183366.01	-/\ #J108					<u>//</u>		*	*	
H= 124.41 } + 8≃ 5.82	****** /\ \/ /\					/\ ≖ 3 /\	0.0 ///	+TSAT= 30.0	+ 	
ISTATE=SATDL I	\/ /\					Λ		*********	LABSORBER OUT	
	>>>>>>							Ň	IT= 30.0 I	
x.						~		17	IP= 579255.21	
***************	**********	*******	***************	*******	+ ¥	<u>//</u>		N/	1H= 62.06 I	
+COLUMN PLATE NUM	BERS: TOTAL=	12: REC	TIFYING= 7: STRIP	PING= 2	*	Λ			ISTATE=SATDL	
RECTIFYING HX TO)TAL= 0.0001	W: STRIP	PING HX IVIAL= U.	NC= 0.61	1	$\overline{\Lambda}$		Ň		
FEEDPLAIE: IEMP *ABSOLUTE REFILIY	FLOW TO DIST	ILLATE FL	OW RATIO= 1.99		+	~		\Z ₹		
#ACTUAL REFLUX FI	OW TO MINIMUN	N REFLUX	FLOW RATIO= 1.15		+ 1	\wedge		M		
*						Λ	<i>2</i>	No.	1 PUMP 10 1	
I EVAPORAT	TOR I CONDENSE	R IABSC	KREK I BOILER I PUN		-*	Λ		Ň	IP= 579255.21	
+ KW 10.000	119.076	110.2	40 19.319 0.0	981 0.00	D*			17	IH= 62.06 I	
***************	***********	********	****************	*******	**	/\		- V	1B= 9.60 I	
+	1 I		I FEED		*	Λ			ISTATE-SATUL I	
•	L INTETILIATE -	OTTOME	L P/HEAT & COLUMN/H	XI TOTAL	- *	$\overline{\Lambda}$		+ PUMP +		
*	INTRATECHICI C				- +	1		********	I PUMP COT I	
•	0.0530 1 (0.1804	1 0.2334 1 0.0000	1 0.233	4 *	\wedge		N/	1T= 29.9 1	
*FLOWS KG/S						(Λ)		×7	16- TT00700*01	
*FLOWS KG/S •CONCENTRATION	0.990 10	0.050	1 0.255			11		17	(H= 52-06 1	

FIGURE H3

NO I	TEMP	I K I Y/X	IHEAT TEI I KW I	FEED FLOW KG/S	IA/BY CH I KJ/S	I PHASE	I MASS I CONC	IENTHALP	YIENTROPY IKJ/KG K	IAVAIL/TY	IFUG COEF	IFUG COEFI	FLOW KG/S
1 1	30.0	1 1.01	1 19.0761	0.0000	-0.006 	LIQ VAP	10.990	80.71 261.28	1 0.327	65.56 0.00	1 0.847 1 0.000	0.001 0.000	0.1056
21	30.7	1.03		0.0000	-0.018	LIQ VAP	10.962	80.75 261.31	0.334 0.922	61.85	I I 0.857 I 0.844	0.187 0.720	0.0526
3 1	32.0	1 1 1.08	0.0001	0.0000	-0.026 	LIQ VAP	1 10.908 10.976	1 80.84 261.45	l 0.344 0.926	55.72 63.74	1 1 0.886 1 0.846	 0.185 0.731	0.0525
4 1	34.6	1 1.18	0.0001	0.0000	 -0.043 	LIQ VAP	 0.806 0.949	 81.10 261.61	 0.357 0.931	45.85	1 1 0.946 1 0.849	 0.183 0.735	0.0520
5	39.7	1.41	0.0001	0.0000	-0.078 -0.078	LIO	1 - 10.637 10.899	 82.07 262.69	 0.371 0.939	31.86	1 1.087		0.0512
6 	48.5	 1.87		0.0000	-0.105 -0.105		10.437	I I 85.47 I 264.58	I 0.383 I	18.14	1 1.377	0.209	0.0511
7 1	58.3	i 2.41	0.0001	a.0000	-0.0691	LIG	1 10:299	1 91.18	1 0.395 1	10.48	1	0.755 - 0.255-	0.1041
8 1	65.0	2.76	0.0001	0.0000	-0.027	FI6	10.235	1 200.31 1 1 95.74	1 0.403 I	7.67	U.881 I I 2.023	0.294	0.1041
9 I	68.3	2.73	. 0.000	0.2334	-0.251	LIQ	10.647 1 10.209	1 267.55 1 1 98.09	0.953 0.408	41.54 6.75	0.889 1 2.159	0.776 0.316	0.1050
1	74.1	3.21		ו ו 0.0000	-0.1751	VAP	10.611 10.170	267.86 102.45	1 0.951 0.616	40.05	0.893		0.1062
1	85.6	3 71		 	-0.4971	VAP	0.545	1 268.90	0.948 	37.65	0.700	0.785	0.0853
ļ	03.0	3.71		1	-0.9731	VAP	0.406	111.54 1 270.53	1 0.433 1 1 0.934 1 1 1	4.80 34.14	2.921	0.457 0.796 	0.2855 0.0913
2	101.1	4.22	-19.319 	0.0000 	3.1371	Q1L VAP	0.050	124.42	1 0.457 0.900	5.82 32.89	3.630 0.939	0.617 0.811	0.1804 0.1051

FIGURE H3 CONTD

KJ/S	PERCENT OF	AVAILABILITY LOSS
	0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000	
0.000	0.000	
-0.042 -0.073	1.114 1.941	
0.099		8
-0.078 -0.052	2.059	e e e e e e e e e e e e e e e e e e e
0.033	-0.883	
2	23.000	6 92 CAN 4
2		5 101 (1 8005185
-0.867	22.990	
3.671		4 _{0.1}
-1.818	48.232	385
0.000 -0.006	0.000	н ж
	0.000 0.0052 0.033 0.033 0.033 0.033 0.033 0.0352 0.0867 1.818	0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.001 0.000 0.002 1.114 -0.052 1.340 0.033 -0.883 23.000 -0.867 22.970 3.671 -1.818 48.232

FIGURE H3 CONTD

****************	******	***********	*************	*****			*************	
* R 11 -R 22	ABSORP	TION REFRIGE	RATION SYSTEM				•	
* R 11 CHLOF	RODIFLU	OROMETHANE	(CHCLF2) +	I DATA AND UNITS			
R 22 TRICH	ILOROFL	JOROMETHANE	(CCL3F	> +	IT=TEMPERATURE DEG C		*	(() T- 0(D
+ CONCENTRATION C	DF DIST	LLLATE= 0.9	40		IP=PRESSURE KPA	1 1= 2*0 <<<	TSATS E 4	K K K I≡ 24+U S
* RATIO OF FEED 1	TO DIST.	LLLAILE 5.4	CEN DOCUEATED~	* • 4 00*	INSENTHALPT NJ/NG		* ISHI- 5.0	
* PROPORIION OF F	FOUEAT A	JW INRUUGH FI	EEV FRENEAIER= 9 OUT=	1.00¥	B-AVAILABILIII KJ/KG	I EVAP IN I		I EVAP OUT I
+ FEED SUBCOOLING	S AT ARS	SORBER OUT=		DEG +	2000 Contraction (1990)	IT= 5.0 [**********	IT= 5.8 I
* DISTILLATE SUBC	COOLING	AT CONDENSE	ROUT= 0.0	DEG +		P= 579255.21	$\Lambda \rightarrow \Lambda \Lambda$	IP= 579255.21
+ APPROACH FLUID	OFF EVA	AP TO EVAP I	N = 0.0 DE	G +		1H= 64.22 1	/\ × \/	IH≖ 252.99 I
*APPROACH CONDENS	SER AND	ABSORBER=	0.0 0.0 DEG			1B= 66.38 I	/\ \/	IB= 50.64 I
* ALL CONCENTRATI	ONS ARE	E MASS CONCS	OF R 11	+	TEXPANSION INT	ISTATE=2 PHA I	+++ \/	ISTATE=SATDV I
*ENTROPY AND ENTH	ALPY DA	TUM TEMPERAT	TURE = -40.0	DEG C+	IT= 17.2 I		*E* \/	
#AVBTY DATUMS: TE	EMP= 30	D.ODEG C :PRE	ESSURE= 101.	3 KPA+	IP= 1183366.01		*** \/	
**************	*******	**********	************	******				
					(5= 0/.1/ ISTATE⇒SUP(1 1	IR/COOL I OUTI		LP/COOL V IN I
					ISTATE-SUDGE T	IT= 17.2	*************	
		TUO QNOD	1			IP= 1183356.01		IP= 579255.21
	i	T= 30.0	1			IH= 64.22 I		IH= 252.99 I
		P= 1183366.0	ונ			1B= 67.17 I	* PRECOOLER *	18= 50.64 I
	l	H= 80.71	1	2		ISTATE=SUBCL I		ISTATE=SATDV I
	1	8= 65.56	1					
(+)	1	STATE=SATDL	1				************	
							<u> </u>	
2	>>	·>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>	»»»»»»»»»»»»»»»»»»»»»»»»»»»»»»»»»»»»»»	>>>>>>	>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>	·····		9
		TSAT=	30.0					
	******							IR/COOL Y OUT I
1= 30.0 >>>	*COND*							1T= 30.0 I
						1P= 1183366.01	N/	1P= 579255.21
		•			81	1H= 80.71	Ň.	1H= 267.47 I
		±1				18= 65.56 I	N/	18= 47.56 I
	÷ .=					ISTATE=SATDL I		ISTATE=SUPHV I
	- ¥ - ¥						1/	
20							V	
66	* *							
	* *							
	* *							
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	* *				10 G	2		
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	- ¥ - ¥						. V	
	+ +							
	• _•						V	
	* *						V.	
2	• •							
	* *				L EEED /			
L COLUMN EECH	- + + 	********			1 FEEV 1		N/ N/	
	* *	5 EEED	•		1P= 1193364-01		1	
1P= 11ARRUALD1	#FEED4	- FELD (((# R0097	Accesses	au	[H= 89.52]		1	
(He Bard) :	4+5+**	HEATER	*	/	NIB= 3.75		V.	
18= 8.75 I	4 #	********	***		VISTATE=SUBCL I			
DEALER THE PLAN				,	· · · · · · · · · · · · · · · · · · ·		X.7	

	•	/ \	IFEED	IN P/H I		\ \	/
*	•		(T= 2	9.9		· · · · · · · · · · · · · · · · · · ·	/
		1	19= 11	87744 01			
*	-		11-11	000000	I BAT TH		
· •	*	//	. IH= 6	2.06	I BOL IN	EXP I A	
*	# - ₂₀	IFEED OUT P/HI /\	18=	9.42 1	11= 29.9	P 1 2 ⁰ A	/ I MIXIN I
	•	LT= 57.9 L/\	T ISTATE	SUBCL I	IP = 11833	166.0I \	/ IT= 30.0 I
		10= 1197744 D1 /\			14- 67 0	14 1 X	/ 19= 579255 21
	_	11- 1103300.01 //			10- 37.7		
*	•	TH= 84.25 1 //	******		IB= 2.8	8 I X.	/ IN= 269.47 I
+	+	18= 8.75 I K	<<# #<<<<	<<<<<<!--</b-->	\ STATE=S	UBCLI V.	/ IB= 47.56 I
	+ 1	STATE=SUBCL I	#FEED #	1			STATE=SUPHY I
L BOT HY OUT I							*
	•		FRE F		``		
[T= 69.5 +	*>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>	>>>>>>>>>> >>>>>>>>>>>>>>>>>>>>>>>>>>	>>+HEAT + >>>	·>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>	\>>>>>>*E*	->>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>	X +
IP= 1183366.01 €	*		+ +	. /		+	+
			*******		182		
10= 73.44 1 4	•			/	1		
18= 2.71 *	•			/	N 3		
ISTATE=SUBCE I +	F	I BOT IN P/H	L L BO	T OUT PZHI Z	N 1	BOT OUT EXP 1 . V	I TUO XIM I
I STATE-SCOVE T		17- 49 6	17-			T- 70 4 4 1	
	•	11- 07.3	11-	27.7 17	`	1- 30.1 1 1	11- 40.1
· · · · · · · · · · · · · · · · · · ·	•	IP= 1183366.0	I IP=	1183366.01 /	\ I	P= 579255.21 \/	/ IP= 579255.21
	•	18= 93.44	1 18=	57.71 1 /	ν I	H= 57.91 W	(IH= 105.93 I
6.6	-	10- 0.74		0.99 / //			(10- 4.24 1
*		16= 2.71	1 16=	2.00 17	V F	8= 2.40 l \/	16= 0.20 1
		ISTATE=SUBCL	I ISTA	TE=SUBCL /	\ I	STATE=SUBCL I \/	/ ISTATE#2 PHA I
J DOT HY THE LA				/			
IT= 101.1 + 3	•	2			1		
IP= 1183366.01 +	F. Contraction of the second se				\	14	
					-	, i i i i i i i i i i i i i i i i i i i	ARSORRER INI
10= 124.41 1 *					77 77		I HOGORDER INT
[B= 5.82 ≠ 3	E)			/`		14	/ IT= 40.1 I
ISTATE=SATDE 1 + 3	•	G					IP= 579255.21
			1.				
				/	`	· · · · · ·	10-103.73 1
* 1	•<<<<				N	\/	/ IB= 8.26 I
		SML			. E U - E		ISTATE=2 PHA I
					•		
I BOTTOM OUT I * *	• 7N			1	L	~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~	
BOTTOM OUT * +					N	\/ *******	*****
BOTTOM OUT * *						\/ ********	*****
I BOTTOM OUT I * IT= 101.1 ****** IP= 1183366.01 *BOIL*			•			*	****
BOTTOM OUT * * T= 101.1 ****** P= 1183366.01 *BOIL; H= 124.41 *****			e 		ν ν ν Τ= 30	* * • ABSORE	****** * SER * >>>
BOTTOM OUT * * * T= 101.1 ****** P= 1183364.0 *BOIL* H= 124.41 ****** B= 5.82 \/		a			``. `	********* * .0 >>> * ABSORE *TSAT=	****** * SER * >>> 30.0*
BOTTOM OUT * * T= 101.1 ****** P= 1183366.01 *BOIL* H= 124.41 ****** B= 5.82 \/		a	د ۲		. T= 30	.0 >>> + ABSORE *TSAT=	 ****** * SER * >>> 30.0*
BOTTOM OUT * * T= 101.1 ****** P= 1183366.01 *BOIL* H= 124.41 ****** B= 5.82 \/ STATE=SATDL \/					``T=`30	******** * •O >>> * ABSORE *TSAT= ********	BER >>> 30.0+
BOTTOM OUT * * * IT= 101.1 ****** IP= 1183366.01 *BOIL* IH= 124.41 ****** IB= 5.82 // ISTATE=SATDL //			•		` T= `30	******** * .0 >>> * ABSORE *TSAT= ********	
I BOTTOM OUT I * IT= 101.1 ****** IP= 1183366.01 *801L3 IH= 124.41 ****** IB= 5.82 \/ ISTATE=SATDL I \/	· /\ • /\ • /\ • /\ · /\ · /\	a	• • •		T= 30	********* * • O >>> * ABSORE *TSAT= *********	***** BER * >>> 30.0* IABSORBER OUTI IT= 30.0 J
I BOTTOM OUT I * IT= 101.1 I ****** IP= 1183366.01 *BOIL3 IH= 124.41 ****** IB= 5.82 \/ ISTATE=SATDL I \/			• • •		T= 30	******** * .0 >>> * ABSORE *TSAT= ********	BER + >>> 30.0* IABSORBER OUTI IT= 30.0 I
BOTTOM OUT * * IT= 101.1 ****** IP= 1183366.01 *BOIL IH= 124.41 ****** IB= 5.82 // ISTATE=SATDL //			* * *		T= 30	******* * • O >>> * ABSORE *TSAT= ******** \/ \/	BER + >>> 30.0* IABSORBER OUTI IT= 30.0 J IP= 579255.21
BOTTOM OUT * * * * * * * * * * * * * * * * * *					T= 30	********* * • O >>> * ABSORE *TSAT= ******** \/ \/ \/	<pre>##### BER # >>> 30.0* IABSORBER OUTI IT= 30.0 J IP= 579255.21 IH= 62.06 i</pre>
BOTTOM OUT * * * * * * * * * * * * * * * * * *		**************************************	PPING= 2 4		. T= 30	******** * • • • • • • • • • • • • • •	HABSORBER OUTI IABSORBER OUTI IT= 30.0 J IP= 579255.21 IH= 62.06 I IB= 9.00 I
BOTTOM OUT * * * * * * * * * * * * * * * * * *	/\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\	IFYING= 7: STRI	PPING= 2 +		T= 30	******** * • O >>> * ABSORE *TSAT= ******** \/ \/ \/	IABSORBER OUTI IT= 30.0 J IP= 579255.21 IH= 62.06 IB= 9.00 J
BOTTOM OUT * * * * * * * * * * * * * * * * * *	<pre>/\ /\\ /\\ /\\\ /\\\\ /\\\ /\\\\ /\\\\ /\\\\ /\\\\ /\\\\ /\\\\ /\\\\ /\\\ /\\\\ /\\\\ /\\\\ /\\\\ /\\\\ /\\\\ /\\\\ /\\\\ /\\\\ /\\\\ /\\\\ /\\\\\ /\\\\\ /\\\\\ /\\\\\\\\</pre>	TIFYING= 7: STRI PING HX TOTAL= 5	PPING= 2 + .587KW +		, T= 30	********* * .0 >>> * ABSORE *TSAT= ******** \/ \/ \/ \/ \/	ABSORBER OUTI IABSORBER OUTI IT= 30.0 I IP= 579255.21 IH= 62.06 I IB= 9.00 I ISTATE=SATOL I
BOTTOM OUT * * * * * * * * * * * * * * * * * *	<pre>/\ /\\ /\\\ /\\ /\\ /\\ /\\ /\\ /\\\ /\\ /\\ /\\\ /\\\ /\\\ /\\\ /\\\ /\\\ /\\\ /\\\</pre>	TIFYING= 7: STRI Sing HX Total= 5 Conc= 0.239: VAP C	PPING= 2 + .587KW + .0NC= 0.453+		T= 30	**************************************	<pre> ###### # BER # >>> 30.0# #################################</pre>
BOTTOM OUT * * * * * * * * * * * * * * * * * *	<pre>/\ /\ /\ /\ /\ /\ /\ /\</pre>	TIFYING= 7: STRI Sing HX Total= 5 Conc= 0.239: VAP C	PPING= 2 + .587KW + :0NC= 0.653+		T= 30	******** * ******* *TSAT= ******* \/ \/ \/ \/	<pre> ##### # ############################</pre>
BOTTOM OUT * * * * * * * * * * * * * * * * * *	<pre>/\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /</pre>	TIFYING= 7: STRI SCONC= 0.237: VAP C DW RATIO= 1.89	PPING= 2 + .587KW + :0NC= 0.653+		. T= 30	******** * • • • • • • • • • • • • • •	<pre>##### BER >>> 30.0* IABSORBER OUTI IT= 30.0 I IP= 579255.21 IH= 62.06 I IB= 9.00 I ISTATE=SATDL I</pre>
BOTTOM OUT * * * IT= 101.1 ****** IP= 1183364.01 *BOIL IH= 124.41 ****** IB= 5.82 // ISTATE=SATDL // >> *COLUMN PLATE NUMBERS: *RECTIFYING HX TOTAL= *FEEDPLATE: TEMP= 64.5 *ABSCLUTE REFLUX FLOW *ACTUAL REFLUX FLOW TO	<pre>/\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /</pre>	TIFYING= 7: STRI TIFYING= 7: STRI SING HX TOTAL= 5 CONC= 0.239: VAP C OW RATIO= 1.87 FLOW RATIO= 1.24	PPING= 2 + .587KW + CONC= 0.653+ *		T= 30	******** * * * * * * * * * * * * * * *	BER + >>> 30.0* IABSORBER OUTI IT= 30.0 I IP= 579255.21 IH= 62.06 i IB= 9.00 I ISTATE=SATOL I
BOTTOM OUT * * * * * * * * * * * * * * * * * *	// // // // // // // // // // // // //	TIFYING= 7: STRI SING HX TOTAL= S CONC= 0.239: VAP C DW RATIO= 1.89 LOW RATIO= 1.24	PPING= 2 + .587KW + CONC= 0.653+		T= 30	******** * * * * * * * * * * * * * * *	<pre></pre>
BOTTOM OUT * * * * * * * * * * * * * * * * * *	<pre>/\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /</pre>	TIFYING= 7: STRI SING HX TOTAL= S CONC= 0.239: VAP C W RATIO= 1.89 LOW RATIO= 1.24	PPING= 2 + .587KW + :ONC= 0.653*		T= 30	******** * .0 >>> * ABSORE *TSAT= ******** \/ \/ \/ \/ \/ \/ \/ \/ \/	<pre> ##### # BER >>> 30.0# IABSORBER OUTI IT= 30.0 I IP= 579255.21 IH= 62.06 IB= 9.00 I ISTATE=SATDL I ISTATE=SATDL I IT= 30.0</pre>
BOTTOM OUT * * * * * * * * * * * * * * * * * *	<pre>/\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /</pre>	TIFYING= 7: STRI TIFYING= 7: STRI TING HX TOTAL= 5 CONC= 0.239: VAP C W RATIO= 1.89 TLOW RATIO= 1.24 RER 1 BOILER 1 PU	PPING= 2 + .587KW + CONC= 0.653+ * MP I BOOST*		T= 30	**************************************	<pre></pre>
I BOTTOM OUT I * IT= 101.1 I ****** IP= 1183366.01 *BOIL IH= 124.41 I ***** IB= 5.82 I // ISTATE=SATDL I // ISTATE=SATDL I // STATE=SATDL I // STATESATDL I // STATESATDL I //	/\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /	TIFYING= 7: STRI PING HX TOTAL= S CONC= 0.239: VAP C DW RATIO= 1.89 FLOW RATIO= 1.24 RBER I BOILER I PU	PPING= 2 + 587KW + ONC= 0.653+ * MP I BOOST+		T= 30	******** * .0 >>> * ABSORE *TSAT= ******** \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/	<pre></pre>
BOTTOM OUT * * * * * * * * * * * * * * * * * *	/\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /	TIFYING= 7: STRI TIFYING= 7: STRI TIG HX TOTAL= 5 CONC= 0.237: VAP C W RATIO= 1.87 TLOW RATIO= 1.24 RBER 1 BOILER 1 PU	PPING= 2 4 .587KW + :ONC= 0.453* # MP I BOOST* 0781 0.000*		T= 30	******** * .0 >>> * ABSORE *TSAT= ******** \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/	<pre> ##### # BER >>> 30.0# #### I ABSORBER OUTI IT= 30.0 I IP= 579255.21 IH= 62.06 I ISTATE=SATDL I IT= 30.0 I IT= 30.0 I IP= 579255.21 IH= 62.06 I </pre>
BOTTOM OUT * * * * * * * * * * * * * * * * * *	/\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /	TIFYING= 7: STRI FING HX TOTAL= S CONC= 0.239: VAP C OW RATIO= 1.89 FLOW RATIO= 1.24 RBER I BOILER I PU	PPING= 2 + .587KW + CONC= 0.453+ # MP I BOOST+ 0781 0.000+		T= 30	******** * ABSORE *TSAT= ******** // // // // // // //	<pre></pre>
BOTTOM OUT * * * * * * * * * * * * * * * * * *	/\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /	TIFYING= 7: STRI PING HX TOTAL= S CONC= 0.239: VAP C OW RATIO= 1.89 COW RATIO= 1.24 RBER 1 BOILER 1 PU O 118.297 1 0.	PPING= 2 + 587KW + 0NC= 0.653+ # MP I BOOST+ 0781 0.000+		T= 30	******* * * * * * * * * * * *	<pre> ##### BER # >>> 30.0# IABSORBER OUTI IT= 30.0 I IP= 579255.21 IH= 62.06 i IB= 9.00 I ISTATE=SATDL I IT= 30.0 I IP= 579255.21 IH= 62.06 I IB= 9.00 I IB= 9.00 I ID= 579255.21 IH= 62.06 I IB= 9.00 I ID= 9.0</pre>
BOTTOM OUT * * * * * * * * * * * * * * * * * *	/\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /	TIFYING= 7: STRI TING HX TOTAL= S CONC= 0.237: VAP C DW RATIO= 1.89 FLOW RATIO= 1.24 RBER I BOILER I PU O I18.297 I 0. FEED	PPING= 2 4 .587KW * :ONC= 0.453* # MP I BOOST* 0781 0.000*		T= 30	**************************************	<pre> ##### # BER # IABSORBER OUTi IT= 30.0 IP= 579255.21 IH= 62.06 IB= 9.00 ISTATE=SATDL I IT= 30.0 IP= 579255.21 IH= 62.06 IB= 7.00 ISTATE=SATDL I ISTATE=SATDL I </pre>
BOTTOM OUT * * * IT= 101.1 ****** IP= 1183366.01 *BOIL IH= 124.41 ****** IB= 5.82 // ISTATE=SATOL // >: *COLUMN PLATE NUMBERS: *RECTIFYING HX TOTAL= *FEEDPLATE: TEMP= 64.5 *ABSCLUTE REFLUX FLOW TO ** EVAPORATOR ** EVAPORATOR ** *	/\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /	TIFYING= 7: STRI FING HX TOTAL= S CONC= 0.239: VAP C OW RATIO= 1.89 FLOW RATIO= 1.24 RBER I BOILER I PU CO 118.297 I 0. FEED	PPING= 2 + .587KW + CONC= 0.453+ # MP I BOOST+ 0781 0.000+		T= 30	**************************************	<pre></pre>
BOTTOM OUT * * * * * * * * * * * * * * * * * *	/\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /	TIFYING= 7: STRI SING HX TOTAL= S CONC= 0.239: VAP C W RATIO= 1.89 LOW RATIO= 1.24 RBER I BOILER I PU O I18.297 I 0. FEED PUBLIC CONTINUE	PPING= 2 + .587KW + :0NC= 0.653+ # MP I BOOST+ 0781 0.000+		T= 30	**************************************	<pre> ###### # BER # >>> 30.0# ##### I ABSORBER OUTI I I T= 30.0 I I P= 579255.21 I H= 62.06 I B= 9.00 I I STATE=SATDL I I T= 30.0 I I P= 579255.21 I H= 62.06 I I B= 9.00 I I STATE=SATDL I #### ###############################</pre>
BOTTOM OUT * * * * * * * * * * * * * * * * * *	/\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /	TIFYING= 7: STRI TING HX TOTAL= S CONC= 0.239: VAP OW RATIO= 1.89 TLOW RATIO= 1.24 REER I BOILER I PU O I18.297 I 0. FEED P/HEAT I COLUMN/	PPING= 2 + .587KW + CONC= 0.653+ 		T= 30	**************************************	<pre></pre>
I BOTTOM OUT I * IT= 101.1 I ****** IP= 1183366.01 *BOIL IH= 124.41 I ***** IB= 5.82 I // ISTATE=SATDL I // 	/\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /	TIFYING= 7: STRI PING HX TOTAL= S CONC= 0.239: VAP C OW RATIO= 1.89 FLOW RATIO= 1.24 REER I BOILER I PU O I18.297 I 0. FEED P/HEAT I COLUMN/	PPING= 2 + 5587KW + ONC= 0.653+ * MP I BOOST+ 0781 0.000+ * HXI TOTAL +		T= 30	****** * ******** *TSAT= ******** // // // // // // // // // //	<pre> ***** BER >>> 30.0* IABSORBER OUTi IT= 30.0 I IP= 579255.21 IH= 62.06 I ISTATE=SATDL I IT= 30.0 I IT= 30.0 I IP= 579255.21 IH= 62.06 I IB= 7.00 I ISTATE=SATDL I *** P** I PUMP OUT I </pre>
I BOTTOM OUT I ******* IT= 101.1 I ****** IP= 1183366.01 *BOIL3 IH= 124.41 ****** IB= 5.82 \/ ISTATE=SATDL I \/ ISTATE=SATDL I \/ *COLUMN PLATE NUMBERS: * *RECTIFYING HX TOTAL= * *FEEDPLATE: TEMP= 64.5 * *ABSQLUTE REFLUX FLOW * * I EVAPORATOR I * * I EVAPORATOR I * I DISTI * I DISTI	/\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /	TIFYING= 7: STRI TIFYING= 7: STRI TING HX TOTAL= 5 CONC= 0.237: VAP C OW RATIO= 1.87 TLOW RATIO= 1.24 RBER 1 BOILER 1 PU O 118.277 1 D. FEED P/HEAT 1 COLUMN/	PPING= 2 4 .587KW + :ONC= 0.653* * MP I BOOST* 0781 0.000* * HXI TOTAL *		T= 30	******** *****************************	<pre> ***** * SER * >>> 30.0* ***** IABSORBER OUTI IT= 30.0 I IP= 579255.21 IH= 62.06 I IB= 9.00 I ISTATE=SATDL I IT= 30.0 I IP= 579255.21 IH= 62.06 I IB= 7.00 I ISTATE=SATDL I *** P** I PUMP OUT I IT= 29.9 I </pre>
BOTTOM OUT * * * * * * * * * * * * * * * * * *	/\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /	TIFYING= 7: STRI TING HX TOTAL= S CONC= 0.239: VAP C OW RATIO= 1.89 TLOW RATIO= 1.24 REER I BOILER I PU O I18.297 I 0. FEED P/HEAT I COLUMN/ 0.2334 I 0.0000	PPING= 2 * .587KW * ONC= 0.653* MP I BOOST* 0781 0.000* * HXI TOTAL * I 0.2334*		T= 30	****** * * * * * * * * * * *	<pre></pre>
BOTTOM OUT * * * T= 101.1 ****** P= 1183366.01 *BOIL H= 124.41 ****** B= 5.82 \/ STATE=SATDL \/ >: *COLUMN PLATE NUMBERS: *COLUMN PLATE NUMBERS	/\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /	TIFYING= 7: STRI SING HX TOTAL= S CONC= 0.239: VAP C W RATIO= 1.89 LOW RATIO= 1.24 RBER I BOILER I PU O I18.297 I 0. FEED P/HEAT I COLUMN/ 0.2334 I 0.0000 0.263	PPING= 2 + 587KW + 0NC= 0.553+ * MP I BOOST+ 0781 0.000+ + HXI TOTAL + I 0.2334+		T= 30	******* * ******** *TSAT= ******** // // // // // // // // // //	<pre> ***** * BER * 30.0* **** IABSORBER OUTi IT= 30.0 I IP= 579255.21 IH= 62.06 IB= 9.00 I ISTATE=SATDL I IT= 30.0 I IP= 579255.21 IH= 62.06 I IB= 9.00 I ISTATE=SATDL I *** P** P** I PUMP OUT I IT= 29.9 I IP= 1183366.0I </pre>
BOTTOM OUT * * * T= 101.1 ****** P= 1183364.01 *BOIL H= 124.41 ****** B= 5.82 \/ STATE=SATDL \/ 	/\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /	TIFYING= 7: STRI TING HX TOTAL= S CONC= 0.237: VAP C DW RATIO= 1.89 FLOW RATIO= 1.24 REER I BOILER I PU O I18.297 I 0. FEED P/HEAT I COLUMN/ 0.2334 I 0.0000 0.263	PPING= 2 4 .587KW * ONC= 0.653* * MP I BOOST* 0781 0.000* * HXI TOTAL * I 0.2334*		T= 30	******* ******************************	<pre> ##### # BER >>> 30.0# #### I ABSORBER OUTI IT= 30.0 J IP= 579255.21 IH= 62.06 I ISTATE=SATDL I IT= 30.0 I IT= 30.0 I IP= 579255.21 IH= 62.06 I IB= 7.00 I ISTATE=SATDL I ### 62.06 I ISTATE=SATDL I ### 1 PUMP OUT I IT= 29.9 I IP= 1183366.0I IH= 62.06 I </pre>
I BOTTOM OUT I ****** IT= 101.1 I ****** IP= 1183366.01 *BOIL3 IH= 124.41 ****** IB= 5.82 \/ ISTATE=SATDL I // 'STATE=SATDL I // *COLUMN PLATE NUMBERS: * *ABSGLUTE REFLUX FLOW * *ACTUAL REFLUX FLOW TO * * I EVAPORATOR I * I DISTI	/\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /	TIFYING= 7: STRI PING HX TOTAL= S CONC= 0.239: VAP C OW RATIO= 1.89 FLOW RATIO= 1.24 REER I BOILER I PU O I18.297 I 0. FEED P/HEAT I COLUMN/ 0.2334 I 0.0000 0.263	PPING= 2 + 587KW + ONC= 0.653+ PPI BOOST+ 0781 0.000+ HX1 TOTAL + 1 0.2334+		T= 30	****** * * * * * * * * * * *	<pre> ***** BER * >>> 30.0* IABSORBER OUTI IT= 30.0 I IP= 579255.21 IH= 62.06 I ISTATE=SATDL I IT= 30.0 I ISTATE=SATDL I IT= 30.0 I ISTATE=SATDL I ISTATE=SATDL I ISTATE=SATDL I ISTATE=SATDL I ISTATE=SATDL I IT= 29.9 I IF= 1183366.0 IH= 62.06 I IE= 7.00 I IE= 7.0</pre>
BOTTOM OUT * * * * * * * * * * * * * * * * * *	/\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /	TIFYING= 7: STRI SING HX TOTAL= S CONC= 0.239: VAP C OW RATIO= 1.89 LOW RATIO= 1.24 RBER 1 BOILER 1 PU O 118.297 1 0. FEED P/HEAT 1 COLUMN/ 0.2334 1 0.0000 0.263	PPING= 2 + .587KW + .0NC= 0.653* * IMP I BOOST* 078I 0.000* * HXI TOTAL * (0.2334*		T= 30	**************************************	<pre> ***** * BER >>> 30.0* **** IABSORBER OUT! IT= 30.0 I IP= 579255.21 IH= 62.06 IB= 9.00 I ISTATE=SATDL I IT= 30.0 I IP= 579255.21 IH= 62.06 I IB= 7.00 I ISTATE=SATDL I *** P** I PUMP OUT I IT= 29.9 I IP= 1183366.01 IH= 62.06 I IB= 7.42 I </pre>
BOTTOM OUT * * * * * * * * * * * * * * * * * *	/\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /	TIFYING= 7: STRI TING HX TOTAL= S CONC= 0.239: VAP C OW RATIO= 1.89 TLOW RATIO= 1.24 REER 1 BOILER 1 PU O 118.297 1 0. FEED P/HEAT 1 COLUMN/ 0.2334 1 0.0000 0.263	PPING= 2 + .587KW + CONC= 0.453 HP I BOOST+ 0781 0.000+ HXI TOTAL + I 0.2334+ +		T= 30	* • • • • • • • • • • • • •	<pre></pre>
I BOTTOM OUT I * IT= 101.1 I ****** IP= 1183366.01 *BOIL' IH= 124.41 ****** IB= 5.82 \/ ISTATE=SATDL I \/ ISTATE=SATDL I \/ *COLUMN PLATE NUMBERS: * *COLUMN PLATE NUMBERS: * *COLUMN PLATE NUMBERS: * *FEEDPLATE: TEMP= 64.5 * *ABSGLUTE REFLUX FLOW * *ACTUAL REFLUX FLOW * * I EVAPORATOR I * * I	/\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /	TIFYING= 7: STRI SING HX TOTAL= S CONC= 0.239: VAP C W RATIO= 1.89 LOW RATIO= 1.24 RBER 1 BOILER 1 PU O 118.297 1 0. FEED P/HEAT 1 COLUMN/ 0.2334 1 0.0000 0.263	PPING= 2 + 5587KW + 0NC= 0.553+ * MP I BOOST+ 0781 0.000+ + HXI TOTAL + I 0.2334+		T= 30	**************************************	<pre> ##### BER >>> 30.0# IABSORBER OUTI IT= 30.0 I IP= 579255.21 IH= 62.06 IB= 9.00 I ISTATE=SATDL I F PUMP IN I IT= 30.0 I IP= 579255.21 IH= 62.06 I IB= 7.00 I ISTATE=SATDL I ### FUMP OUT I IT= 29.9 I IP= 1183366.0I IH= 62.06 I IB= 7.42 I ISTATE=SUBCL I </pre>
BOTTOM OUT * * * IT= 101.1 ****** IP= 1183366.01 *BOIL IH= 124.41 ****** IB= 5.82 \/ ISTATE=SATDL \/ STATE=SATDL \/ *COLUMN PLATE NUMBERS: *RECTIFYING HX TOTAL= *FEEDPLATE: TEMP= 64.5 *ABSOLUTE REFLUX FLOW *ACTUAL REFLUX FLOW * I EVAPORATOR * I EVAPORATOR * I EVAPORATOR * I DISTI * KW 10.000 11 * * I DISTI * LOWS KG/S 0.05 * CONCENTRATION 0.97 * MINIMUM REFLUX FLOW * CAENOT COP* 2.114	/\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /	TIFYING= 7: STRI TIFYING= 7: STRI TING HX TOTAL= 5 CONC= 0.237: VAP C OW RATIO= 1.87 TLOW RATIO= 1.24 RBER 1 BOILER 1 PU O 118.297 1 0. FEED P/HEAT 1 COLUMN/ 0.2334 1 0.0000 0.263	PPING= 2 + .587KW + :ONC= 0.453+ * MP I BOOST* 078I 0.000+ * HXI TOTAL * I 0.2334+ *		T= 30	**************************************	<pre> ##### # BER >>> 30.0# #### IT= 30.0 I IP= 579255.21 IH= 62.06 IB= 9.00 I ISTATE=SATDL I IT= 30.0 I IP= 579255.21 IH= 62.06 I IB= 7.00 I ISTATE=SATDL I ### P # I PUMP OUT I IT= 29.9 I IP= 1183366.0 IH= 62.06 I IB= 7.42 I ISTATE=SUBCL I </pre>
I BOTTOM OUT I * IT= 101.1 I ****** IP= 1183366.01 *BOIL IH= 124.41 I ****** IB= 5.82 I // ISTATE=SATDL I // 	/\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /\ /	TIFYING= 7: STRI PING HX TOTAL= S CONC= 0.239: VAP C OW RATIO= 1.89 FLOW RATIO= 1.24 REER I BOILER I PU O I18.297 I 0. FEED P/HEAT I COLUMN/ 0.2334 I 0.0000 0.263 30.0 & 121.1 D23 W AS %= 0.5	PPING= 2 + 587KW + ONC= 0.653+ PPI BOOST+ 0781 0.000+ HXI TOTAL + 1 0.2334+ C +		T= 30	**************************************	<pre></pre>

FIGURE H4

NO	I TEMP	I K	THEAT THE	FEED FLOW	IA/BY CH	I PHASE	I MASS	IENTHALP	YIENTROPY	AVAIL/T	IFUG COEF	IFUG COEF	I FLOW
	1 DEG C	1 1/1	1 1/10 1	KG/5	1 KJ75	1	I CONC	I KJ/KG	IKJ/KG K	KJ/KG	ICOMPT 1	ICOMPT 2	K6/S
1	1 30.0 1	1 1.01	18.050 	0.0000	1 -0.006	I LIQ I VAP	10.990	1 80.71 1 261.28	1 0.327 1	65.56 0.00	1 0.847	I 0.001	0.0999
2	1 30.7	1 1.03	1 0.0001	0.0000	 -0.016	I LIQ VAP	10.962	1 1 80.75 1 261.31		61.85	1 0.857	0.187	0.0469
3	31.9	1 1.07	0.000	0.0000	 -0.020	LIQ	10.911	1 80.83	I I I I I I I I I I I I I I I I I I I	56.05	1 0.885	0.185	0.0469
4	34.2	1 1.16	0.0001	0.0000	-0.029	LIQ	10.977	1 261.44 1 81.05		64.03 47.11	1 0.845 1 1 0.937	0.731 0.183	0.0999
-	79 5	1		0 0000		VAP	10.953	261.52	1 0.930 1	61.51	0.849	1 0.734 1	0.0999
1	30.5	1		0.0000	-0.049 	VAP	10.873	81.79 262.40	1 0.368 1 1 0.937 1	34.66 57.65	1.051	0.184 0.741	0.0458 0.0994
6 1	45.7	1 1.72	1 0.0001	0.0000	-0.069	Q11 VAP	10.491 10.843	84.17 263.89	i 0.380 i i 0.945 i	21.56 52.44	1.279 0.865	0.200 i 0.751 i	0.0455 0.0788
7	54.3	1 2.19	0.000	0.0000	-0.054 	LIQ Vap	10.347 10.759	1 88.67 1 265.55	I 0.390 I I 0.951 I	12.95 47.14	1 1 1.594 1 0.875	0.234 0.763	0:0461
8 1	61.0	1 2.55	1 0.0001	0.0000	-0.022 -0.022	LIQ	10.271	1 1 92.92	 0.378 0.953	9.20	1 1.856		0.0468
9 1	64.5	2.74	0.0001	0.2334	-0.1331	LIQ	10.239	1 95.39	1 0.403 I	7.84	1 2.002	0.291	0.2879
	73.2	3,17	-2.794	0.0000	0.0201		10.653 1 10.175	1 267.33		41.78 5.75	0.888		0.0978
1		I		1	1	VAP	10.555	268.78	1 0.948 1	38.01	0.879	0.784	0.1075
1 1	86.1	3.73 	1 -2.7941	0.0000	-0.1191	LIQ VAP	0.107 0.400	111.9 4 270.62	1 0.434 1 1 0.933 1	4.80 34.04	2.943 0.915	0.461 0.797	0.2799 0.1006
2	101.2	4.22	1-18.2971	a.aaaa i	2.9801	LIQ VAP	0.050	124.44 273.19	1 0.457 1 1 0.900 1	5.92 32.89	3.631 0.939	0.517 0.811	0.1804
I	A A PY CL			i nalv	i			1.0117	i - i	. 1	01737	0.011	0.077

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FIGURE H4 CONTD

SECOND LAW ANALYSIS	KJ/S	PERCENT OF AVAILAB	ILITY LOSS
PIPEWORK			
CONDENSER OUT TO PRECOOLER IN	0.000	0.000	
PRECOOLER LIQ OUT TO EXPANSION IN	0.000	0.000	
EVAPORATOR OUT TO PRECOOLER IN	0.000	° 0.000 *****	100
PRECOOLER VAP OUT TO MIXING IN	0.000	0.000	
MIXING OUT TO ABSORBER IN	0.000	0.000	
ABSORBER OUT TO PUMP IN	0.000	0.000	
PUMP OUT TO FEED SPLIT	0.000	0.000	
FEED SPLIT TO PREHEATER IN	0.000	0.000	
PREHEATER OUT TO FEED JOIN	0.000	0.000	
FEED SPLIT TO RECTIFIER HX IN	0.060	0.000	
RECTIFIER HX OUT TO FEED JOIN	0.000	0.000	()
BOTTOM OUT OF COLUMN TO HX IN	0.000	0.000	
BOTTOM HX OUT TO PREHEATER IN	0.000	0.000	
PREHEATER OUT TO EXPANSION IN	0.000	0.000	
EXPANSION		-	
DISTILLATE EXPANSION	-0.042	1.174	
BOTTOMS EXPANSION	-0.073	2.047	
PUMP	0.099	-	
INTERNAL HEAT EXCHANGEDS		-	
PRECOOLER	-0.078	2 4 7 9	8
PREHEATER	-0.078	2.170	
	-9.127	3.330	
EXTERNAL HEAT EXCHANGERS		×.	
EVAPURATUR	0.033	-0.931	· · · · ·
EVAP REFRIGERANT STREAMS ONLY -0.834			
EVAP CARNUT -0.867		24.249	
	-		
ABSORBER			
ABSORBER AND MIXING REFRIGERANT STREAMS ONLY	-0.867	24.238	
CARNOT FROM AVERAGE TEMP 0.000			
COLUKN			
CARNOT ONLY	3.477		
(STREAMS ONLY = 1.927)			
COLUMN ONLY=CARNOT-STREAMS	-1.549	43 331	
(MINIMUM SEPERATION WORK= 2.482)		401001	
CONDENSER			
COND CARNOT	0.000	0 000	
CONDENSER STREAMS ONLY	-0.000	0.000	
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FIGURE H4 CONTD

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| * R 11 -R 22 /
* R 11 CHLOR(
* R 22 TRICH
* CONCENTRATION OF | ABSORPTION REFRIGERATION
DDIFLUOROMETHANE (C
OROFLUOROMETHANE (C
F DISTILLATE= 0.990 | SYSTEM # I DAT
HCLF2) # I DAT
CL3F) # IT=TEM
IP=PRE
IH=ENT | A AND UNITS I
PERATURE DEG C I
SSURE KPA I
HALPY KJ/KG | T= 5.0 <<< | +
EVAPORATOR
TSAT= 5.8 + | <<< T= 24.0 |
| - RATIO OF FEED TO | DISTILLATE= 4.41 | ENEATER= 0.99% IB=AVA | ILABILITY KJ/KGI | 4 | · · · · · | |
| * PROPORTION OF FI | ED FLOW THROUGH FEED FA | 0.0 DEG * | | EVAP IN | + + | EVAP OUT |
| · DISTILLATE SUPER | AT ABCORBER OUTS | 0.0 DEG * | E. | T= 5.0 | *********** | 11= 3+0 F |
| + FEED SUBCOULING | OI ING AT CONDENSER OUT= | 0.0 DEG * | 1 | P= 579255.21 | | IN= 252.99 L |
| - APPROACH FILLD | OFF EVAP TO EVAP IN = | 0.0 DEG * | | H= 64.22 | | IB= 50.64 |
| APPROACH CONDENSI | ER AND ABSORBER= 0.0 | 0.0 DEG * | | B= 66.38
State-2 BHA | *** | ISTATE=SATDV I |
| # ALL CONCENTRATIO | ONS ARE MASS CONCS OF | R 11 * | EXPANSION INT | STATE=2 PRA I | *F* \/ | |
| *ENTROPY AND ENTH | ALPY DATUM TEMPERATURE = | -40.0DEG C* | T= 17.2 F | | +++ V | |
| AVBTY DATUMS: TEN | 1P= 30.0DEG C :PRESSURE | = 101.3 KPA* | P = 1103300.01 | | /\ \/ | |
| ************** | ********************* | *************************************** | R = 67.22 | | /\ \/ | |
| | | i | STATE=SUBCL 1 | P/COOL L OUTI | // // | IP/COOL V IN I |
| | | | | T= 17.2 1 4 | *********** | IT= 5.8 I |
| | | | | P= 1183366.01 4 | . + | 1P= 579255.21 |
| | | 3 | 1 | H= 64.22 4 | * | IH= 252.99 |
| | 11= 30,0 · | | 1 | B= 67.17 4 | PRECOOLER | 18= 50.04 I |
| | | | 1 | STATE=SUBCL 1 4 | • | ISTALE=SALUV I |
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| T- 30 8 >>> | +COND+ >>> | | 1 | | ÿ | IT= 30.0 I |
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| | * * | | | STATE=SATDL | 1/ | ISTATE=SUPHV I |
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| I RECT HX IN 1 | * * | | | | | |
| IT= 29.9 | + *
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| P= 1183366.0 | 1 | ~~~~~~~~~~~~~~~~~~~~~~ | <<<<<<<< | << | 2 <u>V</u> | |
| 1H= 62.06 | | | | \wedge | | |
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| 17= 40.2 | +>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>> | ·>>>>>>>>> | | | N/ | |
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| I COLUMN FEEDI | | \/1P= 1 | 183366.01 | /\ IP= 1183366 | · · · · · · · · · · · · · · · · · · · | |
| IT= 53.0 I | ACCEDACCA ROOST 444 | <<<<<<< + H= | 84.60 i | / IH= 62.06</td <td>1 <u>K</u></td> <td></td> | 1 <u>K</u> | |
| IP= 1183366.01 | ARALLA & WEATER # | /\ IB= | 8.68 1 | V /\ 18= 9.42 | | |
| 1H= 64.60 1 | * * ******* | /\ IST/ | TE=SUBCL I | V /\ ISTATE=SUBCI | | |

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H= 124.41 H= 124.41 IB= 5.82 ISTATE=SATDL *COLUMN PLATE NU *COLUMN PLAT	Image: Second state Image: Second state Image: Second state Image: Second state <td>: RECTIFYING= 7: STRIPPIN : STRIPPING HX TOTAL= 6.805 LIQUID CONC= 0.271: VAP CONC= LATE FLOW RATIO= 1.66 REFLUX FLOW RATIO= 0.75 IABSORBER BOILER PUMP I10.240 16.110 0.098! FEED FEED FEED I FEED I FEED I 0.2311 0.0023 050 0.263 0.0367</td> <td>G= 3 + KW * 0.690+ * BOOST+ * 0.000+ ****** TOTAL * 0.2334+ *</td> <td></td> <td>= 30.D >>></td> <td>* ABSORBER *TSAT= 30.0 V V V V V V V V V V V V V</td> <td>IABSORBER OUTI IT= 30.0 IP= 579255.21 IH= 62.06 IB= 9.00 ISTATE=SATDL IT= 30.0 IP= 579255.21 IH= 62.06 ISTATE=SATDL IT= 30.0 ISTATE=SATDL IB= 9.00 ISTATE=SATDL IP= 1183366.01 IH= 62.06 IB= 9.42</td>	: RECTIFYING= 7: STRIPPIN : STRIPPING HX TOTAL= 6.805 LIQUID CONC= 0.271: VAP CONC= LATE FLOW RATIO= 1.66 REFLUX FLOW RATIO= 0.75 IABSORBER BOILER PUMP I10.240 16.110 0.098! FEED FEED FEED I FEED I FEED I 0.2311 0.0023 050 0.263 0.0367	G= 3 + KW * 0.690+ * BOOST+ * 0.000+ ****** TOTAL * 0.2334+ *		= 30.D >>>	* ABSORBER *TSAT= 30.0 V V V V V V V V V V V V V	IABSORBER OUTI IT= 30.0 IP= 579255.21 IH= 62.06 IB= 9.00 ISTATE=SATDL IT= 30.0 IP= 579255.21 IH= 62.06 ISTATE=SATDL IT= 30.0 ISTATE=SATDL IB= 9.00 ISTATE=SATDL IP= 1183366.01 IH= 62.06 IB= 9.42
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 NO I	TEMP	I К	IHEAT TEL	FEED FLOW	A/BY CHI	PHASE	I MASS	IENTHALP	IENTROPYI	AVAIL/TY KJ/KG	IFUG COEF	IFUG COEFI	FLOW KG/S
	DEG C	Y/X	KW 15,857	0.0000	1 -0.0051	LIQ	10.990	1 80.71	1 0.327 1	65.56	1 0.847	0.001	0.0878
2	30.7	1	1 1 1 1 1 0.0101	0.0000	 -0.007	LIQ	10.997	1 201.20 1 1 80.75		61.85	0.859	0.187 0.720	0.0349
	31.7	1	1 1 1 1 1 0.0101	0.0000	 -0.001	LIQ	10.990	80.82	0.342	56.69	1 1 0.881 1 0.845	1 0.185 0.731	 0.0348 0.0879
		1 4 44		0.0000	 0.006	LIQ	10.978 1 10.843	1 261.17 1 80.98	0.352	49.34	1 1 0.922	I I 0.183 I 0.733	 0.0344 0.0878
4	33.0			0.0000	 0.014	VAP	10.959 1 10.729	261.48 81.43	0.929	39.22	1 1.002	I I 0.183	1 1 0.0338 1 0.0876
5	36.8 	1 1.27		0.0000	1	VAP	10.927 I 10.575	1 262.06 1 1 82.77	1 0.934 1 1 0.375	27.34	1 1.157	0.190	I I 0.0328
6	42.0 	1.52 		0.0000		VAP	10.877	1 263.12	1 0.941 1 1 0.384	54.98 1 1 17.31	I 0.860	I I 0.212	0.0319
7	49.2	1 1.91	0.010 	0.0000	 	VAP	10.809	1 264.59	1 0.948	50.15	1 0.869 1 1 1.674	1 0.754 1 1 0.244	0.0865 0.0318
8	56.4	2.30	0.010	0.0000	1 0.016 1	I LIQ VAP	10.321 10.738	1 265.92	0.952	45.97	0.878	1 0.766 1 1 0.279	1 0.0862 0.2759
9	61.0	1 2.55	0.000	0.2334	-0.107 	I LIQ I VAP	10.271	1 92.96 1 266.59 1	1 0.398 1 0.953 1	43.51	1 0.884	0.771	1 0.0865
10	 65.9 	 2.81 	-2.268	0.0000	1 0.095	I LIQ I VAP	10.227	1 96.37 1 267.68	0.405 0.952 	1 7.40 1 41.13 1	1 2.060 1 0.890	0.777	0.0975
11	1 1 74.0	 3.21	1 -2.268	0.0000	-0.011	LIQ VAP	10.170	102.37	1 D.416 1 D.948	1 5.63 1 37.70	1 2.404 1 0.900 1	0.358 0.785	1 0.2636 1 0.0893 1
12	1 86.4	1 1 3.74	 -2.268 	0.000	 -0.153 	I LIQ I VAP	10.106	112.14	0.435	4.80 33.99	1 2.954 1 0.917	0.463 0.797	1 0.2655 1 0.0847
13	101.2	4.22	 -16.110	0.0000	1 1 2.598	I LIQ I VAP	10.050 10.210	124.47	0.457	5.83 32.89	1 3.633 1 0.939	0.617 0.811	0.1805 0.0857
	1	i i	i		1	1	1	1	1			•	

NOTE; A/BY CH is for streams only.

FIGURE H5 CONTD

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SECOND LAW ANALYSIS	KJ/S	PERCENT	0F	AVAILABILITY	LOSS
PIPEWORK	*********				
CONDENSER OUT TO PRECOOLER IN	0.000	0.000		2	
PRECOOLER LIQ OUT TO EXPANSION IN	0.000	0.000			
EVAPORATOR OUT TO PRECOOLER IN	0.000	0.000			
PRECOOLER VAP OUT TO MIXING IN	0.000	0.000			
MIXING OUT TO ABSORBER IN	0.000	0.000			
ABSORBER OUT TO PUMP IN	0.000	0.000			
PUMP OUT TO FEED SPLIT	0.000	0.000			
FEED SPLIT TO PREHEATER IN	0.000	0.000			
PREHEATER OUT TO FEED JOIN	0.000	-0.003			
FEED SPLIT TO RECTIFIER HX IN	0.000	0.000			
RECTIFIER HX OUT TO FEED JOIN	0.000	0.009			
BOTTOM OUT OF COLUMN TO HX IN	0.000	0.000			
BOTTOM HX OUT JO PREHEATER IN	0.000	0.000			
PREHEATER OUT TO EXPANSION IN	0.000	0.000	39		
EXPANSION					
DISTILLATE EXPANSION	-0.042	1.329			
BOTTOMS EXPANSION	~0.073	2.316			
UNP	0.099	-			
NTERNAL WEAT EXCHANCEDE		-			
PRECONIER	0.070				
PREMEATER	-0.078	2.456			
	-0.000	- 2./10			
TERNAL HEAT EXCHANGERS					
VAPORATOR	0.033	-1.054			
EVAP REFRIGERANT STREAMS ONLY -0.834		8			
EVAP CARNOT -0.867		27.436			
BSORBER					
ABSORBER AND MIXING REFRIGERANT STREAMS ONLY Carnot from average temp 0.000	-0.867	27.425			
OLUMN					
CARNOT ONLY	3.044				
(STREAMS ONLY = 1.885)	0.001				
COLUMN ONLY=CARNOT-STREAMS	-1.176	37 214			
(MINIMUM SEPERATION WORK= 2.497)	*****	5/1211			
ONDENSER					
ONDENSER COND CARNOT	0 000	0 000			
ONDENSER COND CARNOT CONDENSER STREAMS ONLY	0.000	0.000			
ONDENSER Cond Carnot Condenser Streams Only	0.000 -0.005	0.000 0.165			

FIGURE H5 CONTD

* R 11 -R 22 ABSORPTION REFRIGERATION SYSTEM * R 11 -R 22 ABSORPTION REFRIGERATION SYSTEM
* R 11 __CHLORODIFLUOROMETHANE (CHCLF2 __1) * II=CDATA AND _UNITS !_
* R 22 TRICHLOROFLUOROMETHANE (CCL3F __] * II=CDATA AND _UNITS !_
CONCENTRATION OF DISTILLATE= 0.970 _____ # IP=PRESSURE KPA __II= 5.0 ______ * EVAPORATOR * ______ (CL3F __] * II=CDATA AND _UNITS !______
* RATIO OF FEED TO DISTILLATE= 0.970 ______ # IP=PRESSURE KPA __II= 5.0 ______ * EVAPORATOR * ______ (CL3F __] * II=CDATA AND _UNITS !______ * EVAPORATOR * ______ (CL3F __] * II=CDATA AND _UNITS !______ * EVAPORATOR * ______ (CL3F __] * II=CDATA AND _UNITS !______ * EVAPORATOR * ______ (CL3F __] * II=CDATA AND _UNITS !_____ * EVAPORATOR * ______ (CL3F __] * II=CDATA AND _UNITS !_____ * EVAPORATOR * ______ (CL3F __] * II=CDATA AND _UNITS !_____ * EVAPORATOR * ______ (CL3F __] * II=CDATA AND _UNITS !_____ * EVAPORATOR * ______ (CL3F __] * II=CDATA AND _UNITS !_____ * EVAPORATOR * ______ (CL3F __] * II=CDATA AND _UNITS !_____ * EVAPORATOR * ______ (CL3F __] * II=CDATA AND _UNITS !_____ * EVAPORATOR * ______ (CL3F __] * II=CDATA AND _UNITS !_____ * EVAPORATOR * ______ (CL3F __] * II=CDATA AND _UNITS !_____ * EVAPORATOR * ______ (CL3F __] * II=CDATA AND _UNITS !_____ * EVAPORATOR * ______ (CL3F __] * II=CDATA AND _UNITS !_____ * EVAPORATOR * ______ (CL3F __] * II=CDATA AND _UNITS !_____ * EVAPORATOR * ______ (CL3F __] * II=CDATA AND _UNITS !_____ * EVAPORATOR * ______ (CL3F __] * II=CDATA AND _UNITS !_____ * EVAPORATOR * ______ (CL3F __] * II=CDATA AND _UNITS !_____ * EVAPORATOR * ______ (CL3F __] * II=CDATA AND _UNITS !______ * EVAPORATOR * _______ (CL3F __] * II=CDATA AND _UNITS !______ * EVAPORATOR * _______ (CL3F __] * II=CDATA AND _UNITS !______ * EVAPORATOR * _______ (CL3F __] * II=CDATA AND _UNITS !______ * EVAPORATOR * _______ (CL3F ______ * EVAPORATOR * _______ * EVAPORATOR * _______ * EVAPORATOR * _______ * EVAPORATOR * _______ * EVAPORATOR * ______ * EVAPORATOR * _______ * *APPROACH CONDENSER AND ABSORBER= 0.0 0.0 DEG -ISTATE=SATDV L *ENTROPY AND ENTHALPY DATUM TEMPERATURE = -40.0DEG C+ IT= 21.4 | +E+ V *** *AVBTY DATUMS: TEMP= 38.0DEG CalPRESSURE= 101.3 KPA+ _____P= 1451145.81
 Hardensis
 Here
 Solution
 Image: Market and the second s ISTATE=SUBCL | IP/COOL L OUTI // IP/COOL V IN I IP= 1451145.81 I COND OUT I IH= 69.51 I + IH= 252.98 I 18= 74.84 1 * PRECOOLER * 18= 53.43 1 ISTATE=SUBCL 1 * PRECOOLER * 18TATE=SATDY L IP= 1451145.81 1H=......91.56 18= 72.62 i ************* AND THINK AND AND AND A STATE=SATOL. N. . IN THE ST /\ TSAT= 38.0 IP/COOL L IN I T= 38.0 >>> +COND+ >>> ىمەرىمەر ئىچەممىيەتەرەپەرەر بەرەۋە ۋە ۋە ۋە يەرىمەر بىرىيەت بىرىيەر بىرىيەت
 IP=
 1451145.81
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 IP=
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 IH=
 291.56
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 IH=
 12.2 At 1 IN. 291.56 A AND V the state of the second second second second second second second second second second second second second se 122 - 2 State of the second of the second of the second of the second of the second of the second of the second of the V V -----יין דער ביין בייני בייני בייני אינגביא אינגעראיג אינגעראי אינגער אינג אינגעראין אינגעראין אינגעראין אינגעראין א A COMPANY ALCONOMY -----· 이 가슴 아이는 것 같아. 1. V 100 C \mathbf{V} $\sim 10^{-10}$ \mathbf{V} \mathbf{V} --------I FEED \mathbf{V} IT= 79.3 I \mathbf{M} \approx I COLUMN FEEDI *********** IT= 79.3 | +++++ + FEED + IP= 1451145.81 IP= 1451145.81 *FEED*<<<* BOOST *<<<<<<< > IH= 108.57 | \mathbf{X} ΔI IH= 108.57 | ₹###### # HEATER ● /\IB= 11.15 | Service Servic *********** /\ISTATE=SUBCL | 18= 11.16 1 *

State and the second second second second second second second second second second second second second second	•		FEED IN P/H 1			
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			H= 67.88	FROT IN EXP.1		ALL THE ALL
	* IFI	ED OUT P/HE /	ABTATE-SUBCI	778 1451145.81	1	118 38.0
	Participation of the	1451145.81 1				1P= 579069.81
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	acatoper 18	11.16. 1. LCC	EED a	CCT STATE=SUBCLI	*******	ISTATE=SUPHY_1_
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(T= 89.3	+>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>	>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>	EAT	_/ <u>\</u>	>>>>>* MIX 4	
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	* 103575 1 1 1 2 V	IT= 89.3	IT= 37.8	I/\ IT= 38		11= 45.0 1 1P= 579069.81
		[P= 1451145.8]			-88 \/	IH= 95.57 I
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	The second second second second second second second second second second second second second second second s	ISTATE=SUBCL L	ISTATE=SUBCL	I /\ ISTATE=	SUBCL I V	ISTATE=2 PHA I
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L POTTON OUT 1 5	A CALL AND A CALL AND A CALL			A start and a start of the start of	• • • • • • • • • • • • • • • • • • •	
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LT=_112,4,****** IP= 1451145.8 *BOIL IH= 136.13	- <u>A</u>		2000 		ABSORBER STSAT= 38.0	
LT=_112.4***** IP= 1451145.8I *BOIL IH= 136.13***** IB= 11.28 I // ISTATE=SATDL I				∧	ABSORBER TSAT= 38.0	
LT=_112.4L***** IP= 1451145.8I *BOIL IH= 136.13***** IB=_11.28 I \/ ISTATE=SATDL I					**************************************	IABSORBER OUTI
IT=_112.4L_, ***** IP= 1451145.8I *BOIL IH= 136.13***** IB=_11.28 I ISTATE=SATDL I	· · · · · · · · · · · · · · · · · · ·			∧	**************************************	IABSORBER OUTI 11=38.0 I 1P= 579069.81
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PREHEATER	- 200		-0.238	5.064			
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FIGURE H6 CONTD

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* R 11 -R 22	ABSORPTION REFRIGERATION SYSTEM	*		+ *	
* R 11 CHLO	RODIFLUOROMETHANE (CHCLF2)	+ I DATA AND UNITS	1		
+ R 22 IRIG	NE DISTILLATES D.990	+ IP=PRESSURE KPA	' T∎ 5.0 <<<	+ FVAPORATOR	- <<< T= 24.0
+ RATIO OF FEED	TO DISTILLATE= 4.41	+ IH=ENTHALPY KJ/KG		+ TSAT= 5.8 .	
+ PROPORTION OF	FEED FLOW THROUGH FEED PREHEATER= 1.00	D+ IB=AVAILABILITY KJ/KG		+ *	
* DISTILLATE SUP	ERHEAT AT EVAPORATOR OUT= 0.0 DEG	*	I EVAP IN I	+ *	I EVAP OUT I
+ FEED SUBCOOLIN	AT ABSORBER OUT= 0.0 DEG	*	IT= 5.0 I	***********	IT= 5.8 I
# DISTILLATE SUB	COOLING AT CONDENSER OUT= U.U DEG	*	IM= 5/9255.21		1P= 5/9255.21
* APPROACH FLUID	CP EVAP IN EVAP IN = 0.0 DEG	• ••••••••	IB= 66.38		
+ ALL CONCENTRAT	LONS ARE MASS CONCS OF R 11	+ IEXPANSION INI	ISTATE=2 PHA I	+++ \/	ISTATE=SATDV I
SENTROPY AND ENT	HALPY DATUM TEMPERATURE = -40.00EG (C= T= 17.2		+E+ \/	
*AVBTY DATUMS: TI	EMP= 30.0DEG C :PRESSURE= 101.3 KPA	+ IP= 1183366.01		+++ \/	
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* * III# 62.06 I	
HEED OUT P/HI /\ IB= 9.42 I	
+ + IT= 58.5 I /\ ISTATE=SUBCL I	IP= 1183366.01 \/ IT= 30.0 I
IP= 1183366-01 /\	IH= 57.91 I \/ IP= 579255.21
	B= 2.88 \/ H= 269.47
	(((/)) STATE=SUBCLI \/ 18= 47.56
	A A A A A A A A A A A A A A A A A A A
+ + ISTATE=SUBCL +FEED +	
TT 70 4 +>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>	·> /\>>>>>+E+>>>>>>>+ MIX +
	/\ ++* + +
	/\ *******
IH= 94.31 I = =	- //
1B= 2.76 1 * *	
ISTATE=SUBCL I + + I BOT IN P/H I BOT OUT P/	
+ + IT= 70.4 1 1T= 29.9	
• • • !P= 1183366.0! !P= 1183366.	0 /\ P= 579255.21 \/ P= 579255.21
1H= 94.31 1 1H= 57.91	/\ H= 57.91 \/ H= 105.93
1P= 2.76 B= 2.88	/\ B= 2,48 \/ B= 8,26
	I IN ISTATESURCE I V/ ISTATE=2 PHA I
* * ISTATE=SUBCL I ISTATE=SUBCL	
1 BOT HX IN I + +	
	N V
	/\ \/
	// I ABSORBER INI
	// IT= 40.1 I
18=.5.82 i + +	// IP= 579255-21
ISTATE=SATDL I * *	
	// 18ª 8.26
	/\ \/ ISTATE=2 PHA I
	//
I BOTTOM OUT I = + //	/\
IT= 101.1 ====== /\	
IP= 1183366.01 →80IL+ //	
1H= 124.41 ###### /\	/\ T= 30.0 >>> + ABSURBER + ///
	/\ +TSAT= 30.0+
	/\ ****************
ISTATE=SALUE I V/ //	// IABSORBER OUTI
>>>>>>	/\ \/ IT= 30.0
	× × × × × × × × × × × × × × × × × × ×

ACTIVATE NUMPERS, TOTAL # 8. RECTIEVING= 4: STRIPPING= 1 *	/\ \/ B= 9.00
COLUMN PLATE NUMBERS, TOTAL OF ACTIVE AT A TATA S 6 43184	/\ \/ ISTATE=SATDL
*RECTIFYING HX TOTAL= U.UUUKWI SIRIFFING A TOTAL STORMAR	//
FEEDPLATE: TEMP= 65.25 DEG C: LIQUID CONCª 0.232: VAP CONCª 0.045	
★ABSOLUTE REFLUX FLOW TO DISTILLATE FLOW RATIO [™] 2.81 *	
ACTUAL REFLUX FLOW TO MINIMUM REFLUX FLOW RATIO= 2.51	
THAT THE PART AND A CONDENSED AREODED A BOTT FR A PIMP A BOOST	/\ \/ T= 30.0
+ I EVAPORATOR I CONDENSER TABSURGER I BUTCER I TOTTE I BOUCH	/\ \/ 1P= 579255-21
*	
* KW 10.000 26.893 10.240 27.133 0.098 0.000*	

FEED *	/\ \/ ISTATE=SATDL
	/\ +++++++
P D D D D D D D D D D D D D D D D D D D	/\ + PUMP +
* IDISTILLATER BUTTONS I P/HEAT I COLONNYBAT FORE	A SERVICE I PUMP OUT I
\$~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~	
FLOWS KG/S 0.0530 0.1804 0.2334 0.0000 0.2334	
ACONCENTRATION 0.990 0.050 0.263 *	// // //= 1183366.01
	/\ \/ {H= 62.06
	<<<<<<<>>
*.MINIMUM REFLUX FLOW (KG/S) = U.U381	ISTATE=SUBCL I

*CARNOT COP= 2.114 BASED ON 5.0, 30.0 & 101.1 DEG C *	
ACOPT 0 347 COPHEAT= 1.364 PUMP KW AS %= 0.4 *	

.

FIGURE H7

1.8

10	I TEMP I DEG C	I K I Y/X	IHEAT TEN	FEED FLOW KG/S	IA/BY CH I KJ/S	I PHAS	EI MASS	IENTHALP	YIENTROPYI	AVAIL/TY KJ/KG	IFUG COEF	IFUG COEF	I FLOW
1	1 30.0	1 1.01	1 26.8931	0.000	1 -0.009	I LIQ	10.990	1 80.71	1 0.327 1	65.56	1 0.847	1 0.001	I NG/S
1	1	.i	i i		1	I VAP I	10.997	1 261.28	1 0.000 1	0.00	0.000	1 0.000	1 0.0000
2	30.7	1 1.03	1 0.0001	0.0000	1 -0.039	LIQ	10.962	80.75	0.334	61.85	1 1 859	1 0 4 9 7	1
i		i	1 1			VAP	10.990	1 261.31	1 0.922 1	65.62	1 0.844	0.720	1 0.0958 0.1489
3 1	32.3	1 1.09	1 0.0001	0.0000	-0.085	LIQ	10.892	1 80.87	1 0.344 1	E4 40	1	1	1
- i		1			!!!	VAP	10.972	1 261.46	1 0.926 1	63.49	0.844	0.184	0.0953
4 1	36.8	1 1.27	i o.oooi	0.0000	-0.219	LIQ	10.728	1 81 44		70.07		1	011400
1		1 1			1 1	VAP	10.927	1 262.12	0.934	37.U/ 57.07	1 1.004 1 0.853		0.0938
5 1	47.9	1 1.83	i 0.000i	0.0000	-0.4751	LIQ	1	1 85 47			1	01700	0.1403
-	•	1	1 1			VAP	10.822	1 264.40	0.382 1	18.83		0.207	0.0932
6 [65.3	2.77	0.0001	0.2334	-0.6661	1 10	1 272	1	1			0.134	0.1468
1		1	!!	1	1	VAP	10.645	267.50		7.59	2.033	0.296 1	0.3379
7 İ	82.7	1 3.60	1 -5.4311	0.0000	-0 3741	1.10	1	l	I I.		0.007	0.776 1	. 8.1462
1		!	1 1	1	013/11	VAP	10.123	109.20	1 0.429 1	4.86	2.790 1	0.430 1	0;3292
1	101.1	1 4.22	1 1	0.0000	1		1			37.80	U.912	0.794	0.1574
I		Ì	1 1	0.000 1	4.3431	LIQ VAP	10.050 (124.41	0.457 1	5.82 1	3.630 i	0.617 1	0.1804
1		1	3 I I	i	1	T CIF		213.19	0.400 1	32.88	0.939	0.811	0.1487

FIGURE H7 CONTD

SECOND LAW ANALYSIS	KJ/S	PERCENT OF	AVAILABILITY	LOSS
PIPFUORK				
CONDENSER OUT TO PRECOOLER IN	0.000	0.000		
PRECOOLER LIQ OUT TO EXPANSION IN	0.000	0.000		
EVAPORATOR OUT TO PRECOOLER IN	0.000	0.000		
PRECOOLER VAP OUT TO MIXING IN	0.000	0.000		
MIXING OUT TO ABSORBER IN	0.000	0.000		
ABSORBER OUT TO PUMP IN	0.000	0.000		
PUMP OUT TO FEED SPLIT	0.000	0.000		
FEED SPLIT TO PREHEATER IN	0.000	0.000		
PREHEATER OUT TO FEED JOIN	0.000	0.000		
FEED SPLIT TO RECTIFIER HX IN	0.000	0.000		
RECTIFIER HX OUT TO FEED JOIN	0.000	0.000		
BOTTOM OUT OF COLUMN TO HX IN	0.000	0.000		
BOTTOM HX OUT TO PREHEATER IN	0.000	0.000		
PREHEATER OUT TO EXPANSION IN	0.000	8.000		
EXPANSION				
DISTILLATE EXPANSION	-0.042	0.799		
BOTTOMS EXPANSION	-0.073	1.393		
PUNP	C.079			
INTERNAL HEAT EXCHANGERS				
PRECOOLER	-0.078	1.477		
PREHEATER	-0.133	2.524		
FYTERNAL HEAT EXCHANGERS				
EVAPORATOR	0.033	-0.634		
EVAP REFRIGERANT STREAMS ONLY -0.834				
EVAP CARNOT -0.867		16.501	8	
	3 .			
ABSORBER				
ABSORBER AND MIXING REFRIGERANT STREAMS ONLY Carnot from average temp 0.000	-0.867	16.494		
COLUMN				
CARNOT ONLY	5.156			
(STREAMS ONLY = 1.936)				
COLUMN ONLY=CARNOT-STREAMS	-3.220	61.277		
(MINIMUM SEPERATION WORK= 2.478)				
CONDENSER				
COND CARNOT	0.000	0.000		
CONDENSER STREAMS ONLY	-0.007	0.159		
	a.000	-		

FIGURE H7 CONTD

*

19 J. Q

■ R 11 -R 22	ABSORPTION REFRIGERATION				
• K 11 UNL		CL3F) + IT=TEMPER	ATURE DEG C	· · · · · · · · · · · · · · · · · · ·	
+ CONCENTRATION	OF DISTILLATE= 0.990	+ IP=PRESSU	RE KPA T= 5.0 <<<	# EVAPORATOR #	<<< T= 24.0
+ RATIO OF FEED	TO DISTILLATE= 6.20	+ H=ENTHAL	PY KJ/KG I	* TSAT= 5.8 +	
+ PROPORTION OF	FEED FLOW THROUGH FEED PRE	EHEATER= 1.00+ IB=AVAILA	BILITY KJ/KGI	* *	
+ DISTILLATE SU	PERHEAT AT EVAPORATOR OUT-	0.0 DEG +	I EVAP IN I	**************************************	I EVAP OUT I
# FEED SUBCOOLI	NG AT ABSORBER OUT=	0.0 DEG +		***********	11= 3.0 I 10= 570255 21
+ DISTILLATE SU	BCOOLING AT CONDENSER OUT=	0.0 DEG +			1H= 252.99 1
+ APPROACH FLUI	D OFF EVAP TO EVAP IN	0.0 DE6 +			18= 50.64 I
#APPROACH CONDE	NSER AND ABSORBER U.U. C		ANSTON INI ISTATE=2 PHA I	*** \/	ISTATE=SATDY I
+ ALL CONCENTRA	TIONS ARE MASS CONCS OF		17.2	*E* \/	
SENTROPY AND EN	TALPT DATUM TEMPERATURE -	- 404 7 KPAs Ps 4	1183366.01	+++ V	
AVBIY DATUMSE	IEAPS SULUES C IPRESSURE		66.22		
************	************************	IB=	67.17		
		ISTA	TE=SUBCL IP/COOL L OUTI		1P/COOL V IN I
			IT= 17.2 I	*********	IT= 5.8 I
	I COND OUT 1	U	IP= 1183366.01	+ +	1P= 579255.21
	(T= 30.0 i		H= 64.22	+ +	IH= 252.99
	[P= 1183366.0]		18= 67.17 i	PRECOOLER +	1B= 50.64 I
	(H= 80.71		ISTATE=SUBCL I	+ +	ISTATE=SATDV I
	1B= 65.56 I			• 2	
	ISTATE=SATDL		12	************	
	>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~		
	/\ TSAT= 30.0			Ň	
	*****			Ň	IP/COOL V OUT
T= 30.0 >>>	+COND+ >>>		IT= 30.0 1	ÿ	1T= 30.0 I
	*****		IP= 1183366-01	ÿ	1P= 579255.21
			(H= 80.71		
	• •				IH= 269.47 I
	• •		IB= 65.56 I		H= 269.47 B= 47.56
	• •		IB= 65.56 I Istate=satdl I		H= 269.47 B= 47.56 State=Suphy
			IB= 45.56 I Istate=satdl I		IH= 269.47 IB= 47.56 ISTATE=SUPHY
			IB= 45.56 ISTATE=SATDL 		IH= 269.47 IB= 47.56 ISTATE=SUPHV
			IB= 45.56 ISTATE=SATDL 		IH= 269.47 IB= 47.56 ISTATE=SUPHV
			IB= 45.56 I Istate=satdl I 		IH= 269.47 I IB= 47.56 I ISTATE=SUPHV I
			IB= 45.56 F ISTATE=SATDL I 		IH= 269.47 I IB= 47.56 I ISTATE=SUPHV I
			IB= 45.56 F ISTATE=SATDL I		IH= 269.47 I IB= 47.56 I ISTATE=SUPHV I
			IB= 65.56 H ISTATE=SATDL I		IH= 269.47 I IB= 47.56 I ISTATE=SUPHV I
			IB= 65.56 I ISTATE=SATDL I		IH= 269.47 I IB= 47.56 I ISTATE=SUPHV I
			IB= 65.56 I ISTATE=SATDL I		IH= 269.47 I IB= 47.56 I ISTATE=SUPHV I
			IB= 65.56 I ISTATE=SATDL I		IH= 269.47 I IB= 47.56 I ISTATE=SUPHV I
			IB= 65.56 I ISTATE=SATDL I		IH= 269.47 IB= 47.56 ISTATE=SUPHV
n Naŭ			IB= 65.56 I ISTATE=SATDL I		IH= 269.47 I IB= 47.56 I ISTATE=SUPHV I
т. Ч.а. У			IB= 65.56 I ISTATE=SATDL I		IH= 269.47 I IB= 47.56 I ISTATE=SUPHV I
5			IB= 65.56 I ISTATE=SATDL I		IH= 269.47 I IB= 47.56 I ISTATE=SUPHV I
n Mai S			IB= 65.56 I ISTATE=SATDL I		IH= 269.47 I IB= 47.56 I ISTATE=SUPHV I
чай 9			IB= 65.56 I ISTATE=SATDL I		IH= 269.47 IB= 47.56 ISTATE=SUPHV
чий 19			IB= 65.56 I ISTATE=SATDL I		IH= 269.47 IB= 47.56 ISTATE=SUPHV
			IB= 65.56 I ISTATE=SATDL I		IH= 269.47 IB= 47.56 ISTATE=SUPHV
т. Чаб Ф			IB= 65.56 ISTATE=SATDL 		IH= 269.47 I IB= 47.56 I ISTATE=SUPHV I
148 1			IB= 65.56 ISTATE=SATDL 		IH= 269.47 I IB= 47.56 I ISTATE=SUPHV I
		I FEED	IB= 65.56 ISTATE=SATDL 		IH= 269.47 IB= 47.56 ISTATE=SUPHV
I COLUMN FEED		I FEED T= 60.2 P= 1183	IB= 65.56 ISTATE=SATDL 		IH= 269.47 IB= 47.56 ISTATE=SUPHV
I COLUMN FEED IT= 60.2	* * * * * * * FEED * *	I FEED T= 60.2 P= 11833	IB= .65.56 ISTATE=SATDL 		IH= 269.47 IB= 47.56 ISTATE=SUPHV

	in the second second second second second second second second second second second second second second second					
	* *		EED IN P/H 1		Ň	×.
			= 1183366.01		Ň.	
		/\ IH	= 62.06 i	I BOT IN EXP I	N/	
		IFEED OUT P/HI /\ IB	= 9.42	IT= 29.9 I	<u> </u>	I MIX IN 1
	· · · · ·	IT= 60.2 1 /\ IS	TATE=SUBCL I	IP= 1183366.01	N N	11= 30.0 I
	• •	P= 1183366.01 /\ -		IN= 59.28 I		r = 3/7233.21
		IH= 91.87 I /\ ++++++		18= 4.07 I		10- 207.47 T
	* *			A STATE-SUBCCI		ISTATE=SUPHY I
	* *	ISTATE=SUBCL I +FEEV +		() ***		
I BOT HX OUT I	* *		· · · · · · · · /	/\>>>>>+E+>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>	+ XIN +<<<	
T= 68.2	* *////////////////////////////////////	+ +	1	/\ +++	+ +	
10- 06 82 i		******	1	/\	********	
10= 79:04 I		*****			\/	
ISTATE=SUBCL I	* *	I BOT IN P/H I	I BOT OUT P/HI /	A IBOT OUT	EXP I V/	
		T= 68.2 1	IT= 29.9 /			11= 30.7 1
- 1993 1993		iP= 1183366.01	IN= 1183366.01 /		33121 \/	iH= 93.19 I
	+ +	H= 94.82	17 - 37.20 1/			18= 8.34 I
	• •	18= 3.62 I	ISTATE=SUBCL 1 /	ISTATE=SU	BCLIV	ISTATE=2 PHA 1
	+ + 	131ALE-300CL (/\	\/	
I BOT HX IN I			1	~	~ //	
T= 82.5 1	1 1		/	1	\/	
10- 400 04 1	1 1		/		V	I ABSORBER INI
19= 4.86			1		5 M (2) 🚺 🖷	
ISTATE=SATOL					N.	
					N N	
	* *<<<<				No.	ISTATES? PHA I
	* * /\				Ň	
I BOTTOM OUT I	+ + /				**********	•
IT= 82.5 I			8		•	+
P= 1183366.01	+B01L+ //			/\ T= 30.0 >>>	+ ABSORBER	+ >>>
IM= 107.00 I			1	/\ /	*TSAT= 30.0	-
ISTATE=SATDL 1	Ň Ň		/		**********	
	>>>>>					17- 70 0 1
						IP= 579255.21
					Ň	IH= 62.06 I
*************	***************************************	$\mathbf{a}_{\mathbf{a}} \mathbf{a}_{\mathbf{b}} \mathbf{a}$. /		Ň	18= 9.00 ł
+COLUMN PLATE N	UMBERS: TOTAL=11: RE	DITAL AT ALL ALL ALL ALL ALL ALL ALL ALL A	+ /	~	Ň	ISTATE=SATOL I
*RECTIFYING HX	101AL= 0.000AW1 31K1	CONC= 0.252: VAP CONC= 0.6	70+ /	1	V/	
PREEVPLAIES IEP	IN FLOW TO DISTILLATE F	LOW RATIO= 1.85	+ /	/ \	N/	
ACTINA SEFINY	FLOW TO MINIMUM REFLUX	FLOW RATIO= 1.16	· · · /		M	
-NUIGHE REFEUX						
	ATOR I CONDENSER TABS	ORBER I BOILER I PUMP I BOO	ST+ /		N N	11= 3U.U I 10= 579255 21
I EVAPOR			/			11 - 577433441
I EVAPOR	; 117.674 10.	224 117.898 1 0.1371 0.0			Ň	18= 9.00 1
• I EVAPOR		*************************	*** /	/ \	Ň	ISTATE=SATDL I
• I EVAPOR	*********************	I PPER	- /		********	
• I EVAPOR		FEED	/			
• I EVAPOF • KW I 10.000		FEED	/	/ \	+ PUMP +	
• I EVAPOR • KW I 10.000	I I I DISTILLATEI BOTTOMS	I FEED	/ L + / + /		• PUMP •	I PUMP OUT I
I EVAPOR	I I IDISTILLATEI BOTTOMS	I P/HEAT I COLUMN/HXI TOTA	/ L + / + / 85+ /		• PUMP • •••••	I PUMP OUT I IT= 29.9 I
• I EVAPOR • KW I 10.000	 DISTILLATE! BOTTOMS 0.0530 0.2755	I P/HEAT I COLUMN/HXI TOTA	// // 85* //		• PUMP • •••••••• \/ \/	I PUMP OUT I IT= 29.9 I IP= 1183366.01
<pre>FLOWS KG/S CONCENTRATION</pre>	 DISTILLATE! BOTTOMS 0.0530 0.2755 0.990 0.124	FEED P/HEAT I COLUMN/HXI TOTA 0.3285 I 0.0000 I 0.32 0.263	/ L + / / 85+ / + /		• PUHP • •••••••• \/ \/ \/	I PUMP OUT I IT= 29.9 I IP= 1183366.01 IH= 62.06 I
<pre>FLOWS KG/S CONCENTRATION FINITHUM DEEL</pre>	 101STILLATE! BOTTOMS 0.0530 0.2755 0.990 0.124	FEED P/HEAT I COLUMN/HXI TOTA 1 0.3285 I 0.0000 I 0.32 0.263	/ L + / + / 85+ / / +	/\ /\ /\ /\ /\ /\	• PUMP • •••••••• \/ \/ \/ <<<<<<<	I PUMP OUT I IT= 29.9 I IP= 1183366.01 IH= 62.06 I IB= 9.42 I

PUMP KW AS %=

*000- 0 664 COPHFAT= 1.547

0.8

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NO I	TEMP DEG C	I K I Y/X	IHEAT TEI I KW I	FEED FLOW KG/8	IA/BY CH I KJ/S	I PHASI I	EI MASS I Conc	IENTHALP	YIENTROPY IKJ/KG K	AVAIL/TY	IFUG COEF ICOMPT 1	IFUG COEF I COMPT 2	I FLOW I KG/S
1	30.0	1.01	1 17.6741	0.0000	-0.006 	LIQ VAP	10.990	1 80.71 1 261.28	1 0.327	65.56	1 0.847 1 0.000	1 0.001 1 0.000	1 0.0979
2	30.7	1.03	0.000	0.0000	 -0.015	LIQ VAP	10.962	80.75 261.31	0.334	61.85	0.859 0.844	0.187	0.0448 0.0979
3	31.8	 1.07 	0.000 	0.0000	-0.018	LIQ VAP	1 10.912 10.977	 80.83 261.44	0.343 0.925	56.18 64.07	 0.884 0.845	 0.185 0.731	 0.0448 0.0978
4	34.1	1, 1 1.16 1 '	0.000	0.0000	 -0.025 	LIQ VAP	1 10.825 10.954	81.04 261.52	0.355 0.930	47.61 61.65	0.934 0.849	0.183	 0.0444 0.0978
5 	38.1	 1.33 	0.000 	0.0000	 -0.041 	LIQ VAP	1 10.687 10.915	81.69 262.31	1 0.367 1 0.936	35.77 58.02	 1.038 0.854	0.184	 0.0439 0.0974
6	44.6	1 1.66	0.000	0.0000	 -0.057 	LIQ VAP	1 10.513 10.853	83.74 263.68	0.378 0.944	23.04 53.14	1.243 0.863	0.197	0.0435
7	52.7	 2.10 		0.0000	 **-0*:046 	LIQ- VAP	1 10.369- 10.775	 87:74 265.26	 -0-388- 0-951	14.14 48.06	 - <u>1.533</u> - 0.874	0.761	 - 0.0438 0.0965
8	59.2	 2.45 	0.000	0.0000	-0.023 	LIQ VAP	 0.289 0.709	 91.77 266.58	 0.396 0.953	10.01	 1.787 0.882	0.260	0.0446
91	63.0	 2.65 	0.000	0.3285	-0.064 -0.064	LIQ VAP	1 10.252 10.670	 94.30 266.92		8.39 42.53	 1.938 0.887	0.282 0.773	0.3776 0.0976
10 	69.3	 2.98 	-3.923 	0.0000	0.1621	LIQ VAP	1 10.201 10.600	%. 98.86 268.21	 0.409 0.951	6.50 39.59	 2.204 0.894	0.323 0.780	0.3636
11	82.5	 3.59 	 -17.898 	0.0000	2,053	LIQ VAP	 0.124 0.444	 109.07 270.08	 0.429 0.938	4.86 34.90	 2.782 0.911	0.429 0.794	0.2755
I NOTE	A/87 C	l Hitai	I I Or streams	s only.			1	1	1 1	1	ι, i	1	

FIGURE H8 CONTD

SECOND LAW ANALYSIS	KJ/8	PERCENT OF
PIPEWORK		
CONDENSER OUT TO PRECOOLER IN	0.000	0.000
PRECOOLER LIQ OUT TO EXPANSION IN	0.000	0.000
EVAPORATOR OUT TO PRECOOLER IN	0.000	0.000
PRECOOLER VAP OUT TO MIXING IN	0.000	0.000
MIXING OUT TO ABSORBER IN	0.000	0.000
ABSORBER OUT TO PUMP IN	0.000	0.000
PUMP OUT TO FEED SPLIT	0.000	0.000
FEED SPLIT TO PREHEATER IN	0.000	0.000
PREHEATER OUT TO FEED JOIN	0.000	0.000
FEED SPLIT TO RECTIFIER HX IN	0.000	0.000
RECTIFIER'HX OUT TO FEED JOIN	0.000	0.000
BOTTOM OUT OF COLUMN TO HX IN	0.000	0.000
BOTTOM HX OUT TO PREHEATER IN	0.000	0.000
PREHEATER OUT TO EXPANSION IN	0.000	0.000
EXPANSION		
DISTILLATE EXPANSION	-0.042	1 500
BOTTOMS EXPANSION	-0.114	4.080
	0.140	
INTERNAL HEAT EXCHANGERS		
PRECUVLEN	-0.078	2.788
	-0.132	4.755
EXTERNAL HEAT EXCHANGERS		
LVAPURATOR	0.033	-1.197
EVAP REFRIGERANT STREAMS ONLY -0.834		
EVAP CARNOT -0.867		31.156
	201	
NBSORBER	•	
ABSORBER AND MIXING REFRIGERANT STREAMS ONLY	-0.573	20.606
CARNOT FROM AVERAGE TEMP 0.000		
OLUMN		
CARNOT ONLY	2.643	
(STREAMS ONLY = 1.639)		
COLUMN ONLY=CARNOT-STREAMS	-1.005	34.094
(MINIMUM SEPERATION WORK= 1.920)		001074
ONDENSER		
COND CARNOT	0.000	0.000
CONDENSER STREAMS ONLY	-0.000	0.000
		0.207
	0.000	
-		

FIGURE H8 CONTD

AVAILABILITY LOSS

	2		
		************	* × 5
 R 11 -R 22 ABSORPTION REFRIGERATION SYSTEM R 11 CHLORODIFLUOROMETHANE (CHCLF2) R 22 TRICHLOROFLUOROMETHANE (CCL3F) CONCENTRATION OF DISTILLATE= 0.930 RATIO OF FEED TO DISTILLATE= 4.48 	I DATA AND UNITS I IT=TEMPERATURE DEG C I IP=PRESSURE KPA I T= 5.0 <<< IH=ENTHALPY KJ/KG I	* * * * * * EVAPORATOR * * TSAT= 10.9 *	<<< T= 24.0
 PROPORTION OF FEED FLOW THROUGH FEED PREHEATER= 1.00* DISTILLATE SUPERHEAT AT EVAPORATOR OUT= 0.0 DEG * FEED SUBCOOLING AT ABSORBER OUT= 0.0 DEG * DISTILLATE SUBCOOLING AT CONDENSER OUT= 0.0 DEG * 	IB=AVAILABILITY KJ/KGI 	* * * * * * * * * * * * * * * * * * *	EVAP OUT T= 10.9 P= 557506.1
 APPROACH FLUID OFF EVAP TO EVAP IN = 0.0 DEG * APPROACH CONDENSER AND ABSORBER= 0.0 0.0 DEG * ALL CONCENTRATIONS ARE MASS CONCS OF R 11 * ENTROPY AND ENTHALPY DATUM TEMPERATURE = -40.0DEG C* 	IH= 66.01 I IB= 58.76 I IEXPANSION INI ISTATE=2 PHA IT= 19.7 I ID= 14/0275 0	/\ \/ /\ \/ *** \/ *E* \/	IH= 253,23 B= 43.58 STATE=SATDV
**************************************	$H^{2} = 66.01$		
	ISTATE=SUBCL 1 IP/COOL L OUT		IP/COOL V IN I
I COND OUT I IT= 30.0 I IT= 1140875 DI	IP= 1140875.01 IH= 66.01 I IH= 59.64 1	* * * * * *	IP= 10.7 IP= 557506.11 IH= 253.23 IB= 43.58
IH= 78.98 I IB= 58.24 I ISTATE=SATDI I	ISTATE=SUBCL 1	* * *	ISTATE=SATDV J
>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>	······	/\ \/	
/\ TSAT≖ 30.0			
T= 30.0 >>> +COND+ >>> ++++++	17-200 L IN 1	N N	IP/COOL V OUTI IT= 30.0 I
	P= 1140875.0 H= 78.98 B= 58.24		P= 557506.1 H= 266.20 8= 41.31
	ISTATE=SATDL I		ISTATE=SUPHV I
* *			
* *			
	с. т.		
* *			
* *			
	I FEED I		
I. COLUMN FEEDI + + + +++++++++++ IT= 52.9 +++++++ + FEED + IP= 1140875.01 +FEED+<<<+ BOOST +<<<<<<<<<<< IH= 83.92 ++++++ + HEATER + / IR= 7.86 + + + +++++++++++++++++++++++++++++	II= 52.9 I IP= 1140875.01 IH= 83.92 I VIB= 7.86 I VISTATE=SUBCI I		
ISTATE=SUBCI I + +	\	Ň	

A A IFEED IN	P/H 1 V
/\ IT= 29.9	
* * /\ IP= 11408	
• • /\ H= 61.6	
- + IFEED OUT P/H1 /\ 18= 8-6	
⊕ ⊕ 17= 52.9 /\ ISTATE=SU	B(L) [P= 11406/5.0] (7 [P= 557506.1]
+ + 1H= 83,92 1 /\ ++++++	(B=2.00)
* * 1B= 7.86 <<<* *<<	ACCCCCC/TI STATE=SUBCCT
* * ISTATE=SUBCL +FEED +	
I BOT HX OUT I + +	
IT= 62.0 I + +>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>	
P= 1140875.0 + +	
IH= 86.51 I + +	// //
1B= 2.37 + *	UT P/HI // IBOT OUT EXP I // I MIX OUT I
ISTATE=SUBCL I * * I ISI IN FM I ISI IN FM I ISI ISI ISI IN FM I ISI ISI ISI ISI ISI ISI ISI ISI ISI	2 /\ T= 30.1 \/ T= 40.3
	0875-01 /\ IP= 557506.11 \/ IP= 557506.11
	-90 1 /\ 1H= 57.90 1 \/ 1H= 104.35 1
	.86 I /\ IB= 2.46 I \/ IB= 7.32 I
	SUBCL J /\ ISTATE=SUBCL J \/ ISTATE=2 PHA 1
	//
BOT HX IN I * *	
IT= 99.2 1 + +	// //
	ABSORBER INI
	// · · · · · · · · · · · · · · · · · ·
	// IP= 557506.11
ISTATE=SAIDL I + +	/\ \/ H= 104.35
	/\ \/ B= 7.32
	/\ ISTATE=2 PHA I
	// ***************
	/\ T= 30.0 >>> + ABSUKBER + ///
ISTATE=SATDL I V/ /V	A LABSORBER OUT I
>>>>>	11 = 30.0
	\/ 1P= 557506.11
	// IH= 61.69
***************************************	() $ B= 8.19$
*COLUMN PLATE NUMBERS: TOTAL= 9: RECTIFYING= 3: STRIPPING= 3 *	ISTATE=SATDL I
*RECTIFYING HX TOTAL= 0.000KW: STRIPPING HX TOTAL= 8.004KW	
*FEEDPLATE: TEMP= 60.70 DEG C: LIQUID CONCE U.258: VAP CONCE D.874	
*ABSOLUTE REFLUX FLOW TO DISTILLATE FLOW RATIO	
*ACTUAL REFLUX FLOW TO MINIMUM REFLUX FLOW RATIO- 1.2/	
*	// IT= 30.0 I
 I EVAPORATOR I CONDENSER TABSORBER I BOILER I FOIL 	// IP= 557506.11
	// H= 61.69
* KW 10.000 18.8// 10.1// 11.6/0 10.0/0 10.0/0	/\ \/ 18= 8.19 1
FEED *	/\ ISTATE=SATUL I
	/\ ********
IDISTULATEL BOTTOMS P/HEAT 1 COLUMN/HX1 TOTAL *	/\ + PUMP +
• · · · · · · · · · · · · · · · · · · ·	
AFLOUS K6/S 0.0534 0.1861 0.2395 0.0000 0.2395*	
ACONCENTRATION 0.930 0.050 0.246 *	
	/ $/$ $/$ $/$ $/$ $/$ $/$ $/$ $/$ $/$
* MINIMUM REFLUX FLOW (KG/S) = 0.0304	
	, STATE-SOUCE 1
*CARNOT COP= 2.057 BASED ON 5.0. 30.0 % 99.2 DEG C *	
*COPHE D.582 COPHEAT= 1.576 PUMP KW 48 %= 0.6	
· · · · · · · · · · · · · · · · · · ·	

TEMP DEG C	I K I Y/X	IHEAT TEI I KW I	FEED FLOW KG/S	IA/BY CHI I KJ/S I	PHASE	I MASS	IENTHALPY I KJ/KG	IENTROPYI Ikj/kg ki	AVAIL/TY KJ/KG	IFUG COEF ICOMPT 1	IFUG COEF ICOMPT 2	I FLOW I KG/S
30.0	1 1.06	1 16.8771 1	0.000	1 -0.0361	LIQ VAP	10.930	/ 78.98 / 261.01	0.334 0.000	58.24 0.00	0.878	0.007	1 0.0924
35.1	1.27		0.0000	-0.109	LIQ Vap	10.735	79.62 261.68	0.357 0.935	39.72 58.63	1.003 0.856	0.181 0.743	0.0382 0.0924
43.8	 1.70	1 0.0001	0.0000	-0.101 	LIQ VAP	10.498 10.849	1 82.27 1 263.44	0.373 0.946	22.01 52.10	1.274 0.867	0.197 0.757	0.0378 0.0914
53.8	2.27	1 0.0001 1 1	0.0000	 -0.062 	LIQ VAP	 0.331 0.751	 87.62 265.31	0.384 0.953	12.05 45.92	 1.653 0.880	0.238	 0.0385 0.0912
60.7	2,65	1 0.0001	0.2395	-0.150 	LIQ VAP	 0.256 0.679	 92.13 266.25	 0.394 0.954	8.47 42.25	1 1.934 0.888	0.276	 0.2896 0.0919
65.4	2.90	1 -2.2281	0.0000	 0.107 	LIQ VAP	1 10.216 10.628	I 1 95.49 1 267.37	 0.401 0.953	6.88 39.98	 2.136 0.894	0.306	 0.2817 0.1035
73.1	3.29	 -2.228 	0.0000	 0.021 	LIQ VAP) 10.164 10.540	 101.21 268.46	 0.412 0.948	5.32 36.79	 2.472 0.903	0.362	 0.2771 0.0956
84.8	3.81	1 -2.2281	0.0000	-0.128	LIQ	 0.104 0.397	 110.46 270.10	 0.429 0.933	4.57	3.008 3.008 0.918	0.464	0.2779 0.0910
99.2	4.30	 -17.096	0.0000	2.713		10.050	1 122.43	 0.451 0.901	5.52	1 3.689 1 1 3.689 1	D.616	0.1861
	DEG C 30.0 35.1 43.8 53.8 60.7 65.4 73.1 84.8 99.2	DEG C i Y/X 30.0 i 1.06 35.1 i 1.27 43.8 i 1.70 53.8 i 2.27 60.7 i 2.65 65.4 i 2.90 73.1 i 3.29 84.8 i 3.81 99.2 i 4.30	DEG C I Y/X I KW I 30.0 1.06 16.8771 I I 35.1 1.27 0.0001 I I 43.8 1.70 0.0001 I I 53.8 2.27 0.0001 I I 60.7 2.65 0.0001 I I 73.1 3.29 -2.2281 I I 84.8 3.81 -2.2281 I I 97.2 4.30 I 17.0961 I	DEG C Y/X KW KG/S 30.0 1.06 16.8771 0.0000 35.1 1.27 0.0001 0.0000 43.8 1.70 0.0001 0.0000 53.8 2.27 0.0001 0.0000 60.7 2.65 0.0001 0.2375 65.4 2.90 -2.2281 0.0000 73.1 3.27 -2.2281 0.0000 84.8 3.81 -2.2281 0.0000 97.2 4.30 -17.0961 0.0000	$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	DEG C Y/X KW KG/S KJ/S 30.0 1.06 16.8771 0.0000 -0.0361 LI0 35.1 1.27 0.0001 0.0000 -0.1091 LI0 35.1 1.27 0.0001 0.0000 -0.1091 LI0 43.8 1.70 0.0001 0.0000 -0.1091 LI0 53.8 2.27 0.0001 0.0000 -0.0621 LI0 60.7 2.65 0.0001 0.2395 -0.1501 LI0 65.4 2.90 -2.2281 0.0000 0.1071 LI0 73.1 3.27 -2.2281 0.0000 0.0211 LI0 84.8 3.81 -2.2281 0.0000 0.0211 LI0 79.2 4.30 -17.0961 0.0000 2.7131 LI0	DEG C i Y/X i KW i KG/S i KJ/S i CONC 30.0 i 1.06 i 16.8771 0.0000 i -0.0361 LI0 i0.930 35.1 i 1.27 0.0001 0.0000 i -0.1091 LI0 i0.735 35.1 i 1.27 0.0001 0.0000 i -0.1091 LI0 i0.735 43.8 i 1.70 0.0001 0.0000 i -0.1011 LI0 i0.478 53.8 2.27 0.0001 0.0000 i -0.0621 LI0 i0.331 60.7 2.65 0.0001 0.2395 i -0.1501 LI0 i0.256 65.4 2.90 i 2.2281 0.0000 0.1071 LI0 i0.216 73.1 3.27 i -2.2281 0.0000 0.0211 LI0 i0.104 84.8 3.81 i -2.2281 0.0000 i 0.0211 LI0 i0.104 97.2 4.30	DEG C I Y/X I KW I KG/S I KJ/S I I CONC I KJ/KG 30.0 I 1.06 I 16.8771 0.0000 -0.0361 LI0 10.930 I 78.98 35.1 I 1.27 0.0001 0.0000 I -0.1091 LI0 10.735 I 79.62 43.8 I 1.70 0.0001 0.0000 I -0.1091 LI0 10.498 82.27 43.8 I 1.70 0.0001 0.0000 I -0.1011 LI0 10.498 82.27 43.8 I 1.70 0.0001 0.0000 I -0.1011 LI0 10.498 82.27 43.8 I 1.70 0.0001 0.0000 I III0 10.498 82.27 43.8 I 1.70 0.0000 0.0000 III0 10.331 87.62 53.8 2.47 0.0000 0.2395 III0 III0 10.256 92.13 60.7 2.65	$\begin{array}{c c c c c c c c c c c c c c c c c c c $	$\begin{array}{c c c c c c c c c c c c c c c c c c c $	DEG C i Y/X i KW i K6/S i KJ/S i CONC i KJ/KG iKJ/KG i	DEG C Y/X I KW I KG/S I KJ/S I CONC I KJ/KG IKJ/KG ICOMPT 1 ICOMPT 2 30.0 I 1.06 I 16.8771 D.0000 -0.036I LIQ ID.930 I 78.98 ID.0334 S8.24 ID.0478 ID.0000 ID.00000

 $\boldsymbol{E}_{i}^{(i)}$

FIGURE H9 CONTD

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+ R 12 -R 113	ABSORPTION REFRIGERATION SYSTEM			+ +	
+ R 12 DICH	LORODIFLUOROMETHANE (CCL2F2)	+ I DATA AND UNITS		+ +	
+ R 113 TRICH	HLOROTRIFLUOROETHANE (CCL2F-CCLF2)	+ II=IEMPERATURE DEG C		+ EVAPORATOR +	<<< T= 26.0
+ PATIO OF FEED 1	TO DISTILLATE= 4.84	* IH=FNTHALPY KJ/KG		+ TSAT= 6.4 +	111 I- 411W
+ PROPORTION OF F	FEED FLOW THROUGH FEED PREHEATER= 1.00	* IB=AVAILABILITY KJ/KG		+ +	
# DISTILLATE SUP	ERHEAT AT EVAPORATOR OUT= 0.0 DEG	•	I EVAP IN I	+ +	I EVAP OUT I
+ FEED SUBCOOLING	5 AT ABSORBER OUT= 0.0 DEG	•	IT= 5.0 I	************	
+ DISTILLATE SUB	COOLING AT CONDENSER OUT= 0.0 DEG	÷	1P= 360458.51		17= 360456.51 14= 490 35 1
* APPROACH FLUID	OFF EVAP TO EVAP IN = U.U DEG				18= 26.88
+ ALL CONCENTRATI	IONS ARE MASS CONCS OF R 12	IEXPANSION INI	ISTATE=2 PHA I	+++ \/	ISTATE=SATDV I
*ENTROPY AND ENTH	HALPY DATUM TEMPERATURE = -40.0DEG C	+ IT= 15.2 I		+E+ \/	
+AVBTY DATUMS: TE	MP= 30.0DEG C :PRESSURE= 101.3 KPA	• IP= 739267.71	<i>.</i>	*** \/	
*************	***************************************	+ 1H= 49.11 I			
		1B= 39.27 J			
	•	ISTATE=SUBCL I	17/100L L 0011	// //	
			IP= 739267.71		IP= 360458.51
	1T= 30.0		IH= 49.11 I	• · •	IH= 190.35 I
	IP= 739267.71		IB= 39.27 I	+ PRECOOLER +	IB= 26.88 I
	1H= 63.66 1		ISTATE=SUBCL I	+ *	ISTATE=SATDV I
	18= 37.86 i	*		-	
	ISTATE=SATDL I				
	/\ TSAT= 30.0			V V	-
	*****			\/	
T= 30.0 >>>	+COND+ T>>>		IP/COOL L IN I	. V	IP/COOL V OUTI
	*****		IT= 30.0 1	N.	1T= 30.0 I
			17= /3726/./I		IF= 300430.31 IU= 204 90 I
			10= 00:00 1		IB= 24.62
			ISTATE=SATDL I	Ň	ISTATE=SUPHV I
10		8.1		V/	
				V	
	* *			V	
	* +				
		14 a	2		
	*: *			N/	
	* +			~ //	
	# +			\/	
	·* +			M.	
					21
				Ň	
	* *			N/	
	* +			\/	
	* *			V	
	* +		3	N N	
		I FFED I			
	* *	IT= 53.3			
1T= 53.3 I	***** * FEED *	IP= 739267.71		Ň	
IP= 739267.71	*FEED*<<<* B00ST *<<<<<<<<<	IH= 78.73 I		1/	
14= 78 73 1	***** * HEATER *	/\ B= 2,23		17	

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+ + + + + + + + + + + + + + + + + + + +	/\ /\ /\ /\ iFEED OUT P/H! /\ iT= 53.3 !/\ iP= 739267.71 /\ iH= 78.73 !/\ *** iB= 2.23 ! <<<* iSTATE=SUBCL ! *FEI *PRI *PRI	<pre>iFEED IN P/H i iT= 29.9 i iP= 739267.7i iH= 57.51 i iB= 2.61 i iSTATE=SUBCL i e+++ e<<<<<<<<<<<<<<<<<<<<<<<<<<</pre>	I BOT IN EXP I // IT= 29.9 // IP= 739267.71 // IH= 56.23 // IB= -0.99 // X/LI STATE=SUBCLI // // ### # // ### # // ### # //>>>>>>> ### #	/ / I MIX IN / IT= 30.0 / IP= 360458.5 / IP= 264.50 / IB= 24.62 **** ISTATE=SUPHV *
IP= 739267.71 * * IH= 84.43 1 * * IB= -0.98 I * * ISTATE=SUBCL I * * * * * * * *		I BOT OUT P/HI IT= 29.9 IP= 739267.71 IH= 56.23 1 IB= -0.99 I ISTATE=SUBCL I	// IBOT OUT EXP I // // IT= 30.1 I // // IP= 360458.51 // // IH= 56.23 I // // IB= -1.23 I // // ISTATE=SUBCL I //	<pre>/ I MIX OUT } / IT= 41.4 / IP= 360458.5 / IH= 93.05 / IB= 2.08 / ISTATE=2 PHA re>
I BOT HX IN I * * IT= 105.9 ! * * IP= 739267.71 * * IH= 126.64 ! * * IB= 3.67 ! * * ISTATE=SATDL ! * * * * * * * * * * * * * * *				ABSORBER INI IT= 41.4 L IP= 360458.51 IH= 93.05 I IB= 2.08 I ISTATE=2 PHA I
IT= 105.9 ****** /\ IP= 739267.7! *BOIL* /\ IH= 126.64 ****** /\ IB= 3.67 \/ /\ ISTATE=SATDI \/ /\			/\ ******** /\ T= 30.0 >>> * ABSORE /\ *TSAT= /\ *******	****** * SER * >>> 30.0= ******
	^	· `.		IT= 30.0 I IP= 360458.51
*COLUMN PLATE NUMBERS: TOTAL=11: R *RECTIFYING HX TOTAL= 0.000KW: STR *FEEDPLATE: TEMP= 60.91 DEG C: LIQUI *ABSOLUTE REFLUX FLOW TO DISTILLATE *ACTUAL REFLUX FLOW TO MINIMUM REFLU	ECTIFYING= 5: STRIPPING IPPING HX TOTAL= 9.076H D CONC= 0.289: VAP CONC= FLOW RATIO= 1.65 X FLOW RATIO= 1.19	5= 3 = (W = 0.765= = = = = 8005T=		IT= 30.0 P= 360458.51 H= 57.51 B= 2.36 STATE=SATDL
*COLUMN PLATE NUMBERS: TOTAL=11: R *RECTIFYING HX TOTAL= 0.000KW: STR *FEEDPLATE: TEMP= 60.91 DEG C: LIQUI *ABSOLUTE REFLUX FLOW TO DISTILLATE * I EVAPORATOR I CONDENSER IAB * I EVAPORATOR I CONDENSER IAB	ECTIFYING= 5: STRIPPING IPPING HX TOTAL= 9.076H D CONC= 0.289: VAP CONC= FLOW RATIO= 1.65 X FLOW RATIO= 1.19 SORBER BOILER PUMP 			IT= 30.0 P= 360458.5 H= 57.51 B= 2.36 STATE=SATDL
*COLUMN PLATE NUMBERS: TOTAL=11: R *RECTIFYING HX TOTAL= 0.000KW: STR *FEEDPLATE: TEMP= 60.91 DEG C: LIQUI *ABSOLUTE REFLUX FLOW TO DISTILLATE * ACTUAL REFLUX FLOW TO MININUM REFLU * I EVAPORATOR I CONDENSER IAB * ' I EVAPORATOR I CONDENSER IAB * ' I I I * ' I I I * ' I I I * ' I I I I * ' I I I I * ' I I I I * ' I I I I I * ' I I I I I I * ' I I I I I I I I I I I I I I I I I I	ECTIFYING= 5: STRIPPING IPPING HX TOTAL= 9.076H D CONC= 0.289: VAP CONC= FLOW RATIO= 1.65 X FLOW RATIO= 1.19 SORBER BOILER PUMP .159 16.176 0.069! FEED 	BOOST= 0.765= BOOST= 0.000= CTAL =	/\ \/ /\ \/	IT 30.0 IT 30.0 IP 360458.5 IH 57.51 IB 2.36 ISTATE=SATDL IT 30.0 IT 30.0 IP 360458.5 IH 57.51 IB 2.36 ISTATE=SATDL ISTATE=SATDL IT 29.9

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NO	I TEMP I DEG C	I K I Y/X	HEAT TEL	FEED FLOW KG/S	IA/BY CHI I KJ/S I	PHASE	I MASS	IENTHALPY	IENTROPY	IAVAIL/TY I KJ/KG	IFUG COEF	FUG COEF	FLOW KG/S
1	1 30.0	1 1.01	15.980 	0.0000	-0.086 	LIQ VAP	10.990	1 63.66 1 200.28	1 0.257	1 37.86 1 0.00	0.872	0.001	0.1165
2	 31.1	 1.05 		0.000	 -0.018 	LIQ VAP	10.945 10.990	64.31 200.83	0.266 0.709	34.05 37.91	0.896 0.869	0.135 0.748	0.0449 0.1165
3	1 1 33.6 1	1 1.14	1 0.0001 1 1	0.0000	 -0.024 	LIQ Váp	10.851 10.973	1 65.76 1 202.08	0.282	27.59	0.953	0.134 0.761	0.0430 0.1157
4	38.7	1 1.38	0.000 	0.0000	-0.042 	LIQ VAP	 0.681 0.937	1 68.96 1 204.86	0.303	18.09 34.40	1.096	0.137 0.769	0.0404 0.1138
5	1 1 47.1	 1.83 	1 0.0001 1 1	0.000	-0.049 -1	LIQ VAP	 0.479 0.878	74.88	0.326 0.745	9.04 31.49	1.385 0.887	0.156 0.781	0.0384 0.1112
6	1 1 55.7 1	 2.33 	0.000 0.000	0.000	 -0.028 	LIQ VAP	 0.347 0.810	81.59 213.90	0.344	4.36 28.99	1.722 0.895	0.190 0.792	0.0382
7	1 1 60.9 1	1 1 2.64 1	1 0.0001	0.2858	 -0.122 	LIQ VAP	10.289 10.765	85.89 216.58	0.355	2.65	1.942	0.215 0.798	0.3376
8	66.8	1 2.98	 -3.025 	0.000	 0.157 	LIQ VAP	! 10.238 10.709	90.86 220.11	0.368	1.51 26.33	1 2.198 0.906	0.249 0.804	0.3228 0.1226
9	76.0	1 1 3.50 1	-3.025 	0.0000	 0.108 	LIQ VAP	10.175 10.613	1 98.95 1 225.35	0.387	0.80 24.71	2.624 0.915	0.313 0.812	0.3129 0.1078
10	 89.7 	 4.18 	-3.0251 	0.0000	 0.048 	LIQ VAP	 0.108 0.450	 111.41 233.89	0.416	1.34 23.68	3.292 0.929	0.431 0.824	0.3112
11	 106.0	1 1 4.79 1	 -16.176 	0.0000	2.814 1	LIQ VAP	10.050 10.237	1 126.78 1 245.25	0.450	3.70 25.35	4.125	0.610 0.836	0.2151 0.0961
	1	1	1		1 1		1	a.,	3	1.	•s = _s		1

NOTE; A/BY CH is for streams only.

FIGURE 10 CONTD

KJ/S

PIPEWORK		
CONDENSER OUT TO PRECOOLER IN	0.000	0.000
PRECOOLER LIQ OUT TO EXPANSION IN	0.000	0.000
EVAPORATOR OUT TO PRECOVER IN	0.000	0.000
MINING OUT TO ABSORBED IN	6.000	0.000
ADCODDED OUT TO BUMP IN	0.000	0.000
PUMP OUT TO FFED SPLIT	0.000	0.000
FEED SPLIT TO PREHEATER IN	0.000	0.000
PREHEATER OUT TO FEED JOIN	0.000	0.000
FEED SPLIT TO RECTIFIER HX IN	0.000	0.000
RECTIFIER HX OUT TO FEED JOIN	0.000	0.000
BOTTOM OUT OF COLUMN TO HX IN	0.000	0.000
BOTTOM HX OUT TO PREHEATER IN	0.000	0.000
PREHEATER OUT TO EXPANSION IN	0.000	0.000
DISTULATE EXPANSION	-0.035	1.061
BOTTOMS EXPANSION	-0.052	1.557
	0 071	
PUMP		•
INTERNAL HEAT EXCHANGERS		
PRECOOLER	-0.060	1.824
PREHEATER	-0.110	3.336
EXTERNAL HEAT EXCHANGERS		
EVAPORATOR	0.001	-0.034
EVAP REFRIGERANT STREAMS ONLY -0.842		05 / 87
EVAP CARNOT -0.843		25.487
ABSORBER	1152	
ABSORBER AND MIXING REFRIGERANT STREAMS ONLY	-0.803	24.267
CARNOT FROM AVERAGE TEMP U.UUU		
COLUMN COLUMN		
CARNOT ONLY	3.237	
(STREAMS ONLY = 1.838)	-1 400	42 344
COLUMN ONLY=CARNOT-STREAMS	-1.400	42.510
(MINIMUM SEPERATION WORK- 2.832)		
CONDENSER		
COND CARNOT	0.000	0.000
CONDENSER STREAMS ONLY	-0.008	0.105
	0.000	

FIGURE HIO CONTD

					-	
2					× H. 10	
R 11 -R 22	ABSORPTION REFRIGERA	ION SYSTEM			+ +	
R 11 CHL	RODIFLUOROMETHANE	(CHCLF2) +	I DATA AND UNITS	1	÷ •	
R 22 TRI	OF DISTILATES 0.990	CCCL3F J #	IPEPRESSURE KPA	I I T= 15.0 <<<	+ EVAPORATOR +	<<< T= 24.
RATIO OF FEED	TO DISTILLATE= 10.69	· · · · · · · · · · · · · · · · · · ·	H=ENTHALPY KJ/KG		+ TSAT= 7.7 +	
PROPORTION OF	FEED FLOW THROUGH FEEL	PREHEATER= 1.00+	IB=AVAILABILITY KJ/KG		* *	
FFED SUBCOOL IN	G AT ABSORBER OUT=	1.0 DEG #		1 = 7.0	*************	1 EVAP OUT
DISTILLATE SUE	COOLING AT CONDENSER (UT= 1.0 DEG #		IP= 614929.61	/\ \/	IP= 614929
APPROACH FLUID	OFF EVAP TO EVAP IN	= 8.0 DE6 +		1H= 82.15 I		1H= 254.40
APPROACH CONDEM All concentral	ISER AND ABSORBER= 10.0 Ions are mass concs of	S.U DEG #	EXPANSION IN	ISTATE=2 PHA		ISTATE=SUPH
ENTROPY AND ENT	HALPY DATUM TEMPERATUR	E = -40.00EG C+	IT= 31.3 I		+E+ \/	
VBTY DATUMS: 1	EMP= 38.0DEG C : PRESS	URE= 101.3 KPA+	IP= 1887815.01		+++ \/	
***********	******************	**************	H= 82.15 B= 73 72			
			ISTATE=SUBCL I	IP/COOL L OUTI	X X	IP/COOL V II
				IT= 31.3 I	********	IT= 8.7
	I COND OUT I			IP= 1889815.01	* *	IP= 614929.
	IP≈ 1889815.0			18= 73.72 I	+ PRECOOLER +	18= 54.56
	IH= 105.85 I	14	a 18	ISTATE=SUBCL I	• •	ISTATE=SUPH
					*	
					/\ \/	
	>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>	>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>	>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>	·>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>	>>> \/	
÷	/\ TSAT= 4	9.0				
= 38.0 >>>	+COND+ >>>			IP/COOL L IN I	V V	IP/COOL V OU
	*****			IT= 48.0 I	V/	IT= 43.0
				IP= 1889815.01	N N	IP= 614929.
	* *			1B= 72.26 I	N/	IB= 50.16
	* *			ISTATE=SUBCL 1	V/	ISTATE=SUPHV
S2	* *					
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I COLUMN FEEDI		F	IT= 96.7 1			
	***** * FEED	•	IP= 1889815.01		V	

				IFFED IN P/H 1			V	
			7	IT= 42.7			Ň.	
	1 2		<u>, , , , , , , , , , , , , , , , , , , </u>	IP= 1889815.01			Ň	
					1 901	T TN EVD I	· · · · ·	
		LEEED OUT	T B/UL /		1.7-		Ň	
	- <u>-</u>				10-1	9/ 1	No.	
		11= 70.		ISTATE=SUBCE 1	17= 1		No.	
	+	IP= 1889	815.UI /\		111=	/3./5	N.	IP= 614929.61
	4 A	IH= 124.	46 1 / +++	****	18=	7.66	N/	IH= 278.10 I
	÷ ÷	18= 11.3	20 <<<+	****************		TE=SUBCLI	~ //	IB= 50.16
	+ *	ISTATE=SU	BCL I +FEI	ED +	/\		********	ISTATE≖SUPHV I
I BOT HX OUT I	+ ÷		+PR8	E +	/ \	***	+ +	
IT= 109.0 I	+ *>>>>>>>>>	>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>	>>>>>>+++E/	AT = >>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>	/\>>>>	>+E+>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>	>>>+ MIX +	
IP= 1889815.01	+ (*		+	+ v	Λ	***		
IN= 132 35 I			****		1		********	
	1				<u></u>		\/	
		I BOT	T TN D/H I	I BOT OUT BAN		. IBOT OUT	EYP I V/	I MTY OUT I
ISTATE=SUBLE T					~			
	+ *	11=	109.0	11= 47+7 1	~	11= 48.7		11= 53.3 1
	+ +	19= 1	1889815.01	P= 1889815.0	Λ	18= 6149	29.61 \/	IP= 614929.61
•	+ *	IH= 1	132.35	IH= 73.75 I	\sim	iH= 73.7	'S I \/	IH= 92.86 I
	+ *	1B=	10.98	IB= 7.66 I	/ \	1B= 6.8	1 1 \/	IB= 8.26 I
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I BOT HY TH I					1		\/	
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IP= 1889815.UI	+ +				\sim		N/	
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IB= 14.31 I	+ *				Λ		\/	IT= 53.3 I
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I BOITOM OUT I	***** /\				<u>//</u>		**********	*
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BOTION OUT T= 127.9 P= 1889815.0 H= 153.11	****** /\ *BOIL* /\				/\ /\ /\ T=	38.0 >>>	+ ABSORBER	* * * * * * * * * * * * * * * * * * * *
IT= 127.9 IT= 127.9 IP= 1889815.0 IH= 153.11 IB= 14.31	****** /\ *BOIL* /\ ****** /\				/\ /\ /\ T=	38.0 >>>	**************************************	* * * * * *
I BOTTOM OUT I IT= 127.9 IP= 1889815.0 IH= 153.11 IB= 14.31 ISTATE=SATDL I	****** /\ *BOIL* /\ ****** /\ \/ /\				/\ /\ /\ T= /\	38.0 >>>	* * * ABSORBER *TSAT= 48.0	* * >>> *
I BOTTOM OUT I IT= 127.9 IP= 1889815.01 IH= 153.11 IB= 14.31 ISTATE=SATDL	****** /\ *BOIL* /\ ****** /\ \/ /\ \/ /\				/\ /\ /\ T= /\ /\	38.0 >>>	* * ABSORBER *TSAT= 48.0	* * * * * * * * * * * * * * * * * * *
IT= 127.9 IT= 127.9 IP= 1889815.0 IH= 153.11 IB= 14.31 ISTATE=SATDL	****** /\ *BOIL* /\ ****** /\ \/ /\ \/ /\			**	/\ /\ /\ T= /\ /\	38.0 >>>	* * ABSORBER *TSAT= 48.0	* * >>> * IABSORBER OUTI
I BOTTOM OUT I IT= 127.9 IP= 1889815.0 IH= 153.11 IB= 14.31 ISTATE=SATDL	****** /\ *BOIL* /\ ****** /\ \/ /\ >>>>>>		,	5 . 5	/\ /\ /\ /\ /\ /\	38.0 >>>	• • ABSORBER •TSAT= 48.0	* * * IABSORBER OUTI IT= 43.0 ID= 414929 41
I BOTTOM OUT I IT= 127.9 IP= 1889815.0 IH= 153.11 IB= 14.31 ISTATE=SATDL	****** /\ *BOIL* /\ ****** /\ \/ /\ \/ /\ >>>>>>		a v	525	ハ ハ ハ ハ ハ ハ ハ	38.0 >>>	* ABSORBER *TSAT= 48.0	* * * IABSORBER OUTI IT= 43.0 IP= 614929.61
I BOTION OUT I IT= 127.9 I IP= 1889815.0I IH= 153.11 I IB= 14.31 I ISTATE=SATDL I	*BOIL* /\ *BOIL* /\ // /\ // /\ // /\ >>>>>>				∧ ∧ ∧ ∧ ∧ ∧ ∧ ∧	38.0 >>>	* * * * * * * * * * * * * * * * * * *	* * >>> * ABSORBER OUT T= 43.0 P= 614929.61 H= 71.34
I BOTTOM OUT I IT= 127.9 IP= 1889815.0 IH= 153.11 IB= 14.31 ISTATE=SATDL 	****** /\ *BOIL* /\ ****** /\ \/ /\ \/ /\ >>>>>>	3: RECTIFYING=	7: STRIPPING	****** a 3 +		38.0 >>>	* * ABSORBER *TSAT= 48.0 \/ \/ \/ \/ \/ \/	* * * IABSORBER OUTI IT= 43.0 P= 614929.6 IH= 71.34 IB= 8.63
I BOTTOM OUT I IT= 127.9 IP= 1889815.01 IH= 153.11 IB= 14.31 ISTATE=SATDL 	****** /\ *BOIL* /\ ****** /\ \/ /\ \/ /\ >>>>>> UMBERS: TOTAL=13 TOTAL= 0.000KW	5: RECTIFYING= S: STRIPPING HX T	7: STRIPPING OTAL= 11.678K	****** = 3 + W +		38.0 >>>	* * ABSORBER *TSAT= 48.0 ***** // // // // // // // // // //	* * * * * * * * * * * * * *
I BOTTOM OUT I IT= 127.9 IP= 1889815.0 IH= 153.11 IB= 14.31 ISTATE=SATDL 	****** /\ *BOIL* /\ ****** /\ \/ /\ \/ /\ >>>>>> UMBERS: TOTAL=13 TOTAL= 0.000KW P=103.95 DEG C:	3: RECTIFYING= 1: STRIPPING HX T LIQUID CONC= 0.1	7: STRIPPING OTAL= 11.678K 50: VAP CONC≠			38.0 >>>	*********** * ABSORBER *TSAT= 48.0 ************************************	* * * * * * * * * * * * * *
I BOTTOM OUT I IT= 127.9 IP= 1889815.0 IH= 153.11 ISTATE=SATDL ISTATE=SATDL 	****** /\ *BOIL* /\ ****** /\ \/ /\ \/ /\ \/ /\ >>>>>> UMBERS: TOTAL=13 TOTAL= 0.000KW IP=103.95 DEG C: X; FLOW TO DISTIL	S: RECTIFYING= S: STRIPPING HX T LIQUID CONC= 0.1 LATE FLOW RATIO=	7: STRIPPING TAL= 11.678K 50: VAP CONC= 3.19	•••••• ≖ 3 + ₩ + 0.428+ +		38.0 >>>	* ABSORBER *TSAT= 48.0 \/ \/ \/ \/ \/ \/ \/ \/ \/	* * * ABSORBER OUTI T= 43.0 P= 614929.61 H= 71.34 B= 8.63 STATE=SUBCL
I BOTTOM OUT I IT= 127.9 IP= 1889815.0 IH= 153.11 ISTATE=SATDL 	****** /\ *BOIL* /\ ****** /\ \/ /\ \/ /\ \/ /\ >>>>>> ******* /\ \/ /\ \/ /\ \/ /\ >>>>>> ******* /\ \/ /\ \/ /\ \/ >>>>>> ***************************	5: RECTIFYING= N: STRIPPING HX T Liquid Conc= 0.1 Late flow Ratio= Refiux flow Rati	7: STRIPPING OTAL= 11.678K 50: VAP CONC= 3.19 0= 1.48	= 3 + ₩ + 0.428+ +		38.0 >>>	*	* * * ABSORBER OUTI T= 43.0 P= 614929.6 H= 71.34 B= 8.63 ISTATE=SUBCL
I BOTTOM OUT I IT= 127.9 IP= 1889815.0 IH= 153.11 IB= 14.31 ISTATE=SATDL 	****** /\ *BOIL* /\ ****** /\ \/ /\ \/ /\ \/ /\ >>>>> ******* /\ \/ /\ \/ /\ >>>>> ******* /\ \/ /\ \/ /\ >>>>> *******************************	5: RECTIFYING= 4: STRIPPING HX T LIQUID CONC= 0.1 LATE FLOW RATIO= REFLUX FLOW RATI	7: STRIPPING OTAL= 11.678K 50: VAP CONC= 3.19 0= 1.48	= 3 + ₩ = 0.428+ +		38.0 >>>	*	* * * * * * * * * * * * * *
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I BOTTOM OUT I IT= 127.9 IP= 1889815.0 IH= 153.11 ISTATE=SATDL ISTATE=SATDL *COLUMN PLATE N *RECTIFYING HX *FEEDPLATE: TEM *ABSOLUTE REFLUX * I EVAPOR	HUMBERS: TOTAL=13 TOTAL= 0.000KW P=103.95 DEG C: IX FLOW TO DISTIL FLOW TO MINIMUM	3: RECTIFYING= N: STRIPPING HX T LIQUID CONC= D.1 LATE FLOW RATIO= REFLUX FLOW RATI R IABSORBER BO	7: STRIPPING OTAL= 11.678K 50: VAP CONC= 3.19 0= 1.48 DILER I PUMP I	= 3 + W + 0.428+ B00ST+		38.0 >>>	*	<pre>* * * * * * * * * * * * * * * * * * *</pre>
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I BOTTOM OUT I IT= 127.9 IP= 1889815.0 IH= 153.11 ISTATE=SATDL 	****** /\ *BOIL* /\ *BOIL* /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ <td>S: RECTIFYING= S: STRIPPING HX T LIQUID CONC= 0.1 LATE FLOW RATIO= REFLUX FLOW RATIO (ABSORBER BO (113.360 32.) (113.360 33.) (113.360 /td> <td>7: STRIPPING OTAL= 11.678K 50: VAP CONC= 3.19 0= 1.48 DILER PUMP 801 0.541 FEED COLUMN/HX! T</td> <td>= 3 + W + 0.428+ BOOST+ 0.000+ + 0TAL +</td> <td></td> <td>38.0 >>></td> <td>* ABSORBER TSAT=</td> <td><pre>* * * * * * * * * * * * * * * * * * *</pre></td>	S: RECTIFYING= S: STRIPPING HX T LIQUID CONC= 0.1 LATE FLOW RATIO= REFLUX FLOW RATIO (ABSORBER BO (113.360 32.) (113.360 33.) (113.360 7: STRIPPING OTAL= 11.678K 50: VAP CONC= 3.19 0= 1.48 DILER PUMP 801 0.541 FEED COLUMN/HX! T	= 3 + W + 0.428+ BOOST+ 0.000+ + 0TAL +		38.0 >>>	* ABSORBER TSAT=	<pre>* * * * * * * * * * * * * * * * * * *</pre>	
I BOTTOM OUT I IT= 127.9 IP= 1889815.0 IH= 153.11 IB= 14.31 ISTATE=SATDL 	****** /\ *BOIL* /\ *BOIL* /\ *BOIL* /\ // /\ // /\ // /\ >>>>>> UMBERS: TOTAL=13 TOTAL= 0.000KW /P=103.95 DEG C: X FLOW TO DISTIL FLOW TO DISTIL FLOW TO MINIMUM	S: RECTIFYING= S: STRIPPING HX T LIQUID CONC= 0.1 LATE FLOW RATIO= REFLUX FLOW RATIO I 13.360 I32.	7: STRIPPING 0TAL= 11.478K 50: VAP CONC= 3.19 0= 1.48 01LER I PUMP 801 0.541 FEED COLUMN/HX! T	= 3 + W + 0.428+ = BOOST= 0.000+ ***** 0.000+ ***** 0.000+		38.0 >>>	* ABSORBER TSAT=	<pre>* * * * * * * * * * * * * * * * * * *</pre>
I BOTTOM OUT I IT= 127.9 IP= 1889815.01 IH= 153.11 IB= 14.31 ISTATE=SATDL 	****** /\ *BOIL* /\ *BOIL* /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ <td>3: RECTIFYING= 4: STRIPPING HX T LIQUID CONC= 0.1 LATE FLOW RATIO= REFLUX FLOW RATIO 1 13.360 132. 1 1 1 1 1 1 1 1 1 1 1 1 1</td> <td>7: STRIPPING OTAL= 11.678K 50: VAP CONC= 3.19 0= 1.48 DILER PUMP 801 0.541 FEED COLUMN/HX! T 0.0000 0</td> <td>= 3 + H + 0.428+ + BOOST+ + 0.000+ ++++++ 0TAL + + .6206+</td> <td></td> <td>38.0 >>></td> <td>* ABSORBER TSAT=</td> <td><pre>* * * * * * * * * * * * * * * * * * *</pre></td>	3: RECTIFYING= 4: STRIPPING HX T LIQUID CONC= 0.1 LATE FLOW RATIO= REFLUX FLOW RATIO 1 13.360 132. 1 1 1 1 1 1 1 1 1 1 1 1 1	7: STRIPPING OTAL= 11.678K 50: VAP CONC= 3.19 0= 1.48 DILER PUMP 801 0.541 FEED COLUMN/HX! T 0.0000 0	= 3 + H + 0.428+ + BOOST+ + 0.000+ ++++++ 0TAL + + .6206+		38.0 >>>	* ABSORBER TSAT=	<pre>* * * * * * * * * * * * * * * * * * *</pre>
I BOTTOM OUT I IT= 127.9 IP= 1889815.01 IH= 153.11 IB= 14.31 ISTATE=SATDL *COLUMN PLATE N *RECTIFYING HX *FEEDPLATE: TEM *ABSOLUTE REFLU *ACTUAL REFLUX *	****** /\ *BOIL* /\ *BOIL* /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ /\ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ <td>5: RECTIFYING= 4: STRIPPING HX T LIQUID CONC= 0.1 LATE FLOW RATIO= REFLUX FLOW RATIO 113.360 132. 1 1 1 1 1 1 1 1 1 1 1 1 1</td> <td>7: STRIPPING OTAL= 11.678K 50: VAP CONC= 3.19 0= 1.48 DILER PUMP 801 0.541 FEED COLUMN/HX T 0.0000 D 0.138</td> <td>= 3 + W + 0.428+ + BOOST+ + 0.000+ + 0TAL + 0TAL + + .6206+ +</td> <td></td> <td>38.0 >>></td> <td>*</td> <td><pre>* * * * * * * * * * * * * * * * * * *</pre></td>	5: RECTIFYING= 4: STRIPPING HX T LIQUID CONC= 0.1 LATE FLOW RATIO= REFLUX FLOW RATIO 113.360 132. 1 1 1 1 1 1 1 1 1 1 1 1 1	7: STRIPPING OTAL= 11.678K 50: VAP CONC= 3.19 0= 1.48 DILER PUMP 801 0.541 FEED COLUMN/HX T 0.0000 D 0.138	= 3 + W + 0.428+ + BOOST+ + 0.000+ + 0TAL + 0TAL + + .6206+ +		38.0 >>>	*	<pre>* * * * * * * * * * * * * * * * * * *</pre>
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I BOTTOM OUT I IT= 127.9 IP= 1889815.01 IH= 153.11 ISTATE=SATDL I 	<pre>#BOIL# /\ #BOIL# /\ #BOIL# /\ #BOIL# /\ // /\ // /\ // /\ >>>>>>> #############################</pre>	5: RECTIFYING= 4: STRIPPING HX T LIQUID CONC= 0.1 LATE FLOW RATIO= REFLUX FLOW RATIO= 113.360 132. 1 1 0.0859 15.0. 38.0 & PUMP KU AS *-	7: STRIPPING OTAL= 11.678K 50: VAP CONC= 3.19 O= 1.48 DILER I PUMP I 801 I 0.541I FEED I COLUMN/HXI T I 0.0000 I 0 0.138 127.9 DEG C 1.6	= 3 + W + 0.428+ = BOOST= 0.000+ ***** 0.000+ ***** 0.000+ ***** * 0.000+ ***** * 0.000+ ***** * 0.000+ ***** * *		38.0 >>>	* ABSORBER *TSAT= 48.0 \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/ \/	<pre> * * * * * * * * * * * * * * * * * * *</pre>

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	I DEG C	i y/x	I KW	I KG/S	i KJ/S	I	I CONC	IENTHALP I KJ/KG	YIENTROP	YIAVAIL/T KI KJ/KG	YIFUG COEL ICOMPT 1	FIFUG COE ICOMPT 2	FI FLOW I KG/S
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	1			1	1	I VAP	10.997	1 264.94	1 0.000	1 0.00	1 0.000	1 0.000	1 0.1853
2	49.6	1 1.02	0.000	1		1		!	1	1	1	1	1
	1	1	I	1 0.0000	1 ~0.040	I VAP	10.969	1 107.33	0.421	69.37	0.802	1 0.205	1 0.1282
-	1	1		l	1	1	1	1 204.70	1 0.705	1 //.04	1 0.792	0.632	0.1853
3	1 50.8	1 1.05	0.000	0.0000	1 -0.064	I LIO	10,925	1 107.23	1 0.429	64.10	0.822	L D.204	1 0 1277
	i	1 × 1				I VAP	10.976	1 265.11	1 0.908	1 75.87	1 0.794	1 0.641	1 0.1863
- 4	1 53.6	1 1.13	1 0.0001	0.0000	/ / -0.133	1 10		1 107 44		1	!	1	1
	!	1	1 1		1	I VAP	10.945	1 265.49	1 0.440	1 54.72		1 0.203	0.1265
	1 50 7	1 4 77	1		1	1	1	ì	1	1 /2:01	1 0.777	1 0.046	0.1858
2	1 37.7	1 1.33	1 0.0001	0.0000-	1 -0.277	I LIO	10.663	1 107.53	1 0.452	1 39.91	1 0.988	1 0.208	0.1234
	1	i	i i			I VAP	10.883	266.41	1 0.923	1 66.95	0.811	1 0.657	1 0.1846
6	1 71.5	1 1.77	1 0.0001	0.0000	-0.453	LIO	10.433	1 111 36	1 0 444	1 27 97	1 4 979		!
		1			1	VAP	10.767	1 269.45	1 0.935	58.62	0.831		0.1226
7	I 86.8	2.33	1 0 0001	0.0000			1	1	1	1	i	1 0.077	0.1015
	1 0010	1	1 0.0001	0.0000	-0.388		10.263	1 120.38	1 0.480	15.37	1.694	0.313	0.1273
	I	1	1 1	i	i i	, var	10.012	212.19	1 0.939	50.39	0.857	0.700	0.1806
8	98.2	1 2.69	1 0.0001	0.0000	-0.153	LIQ	10.182	128.65	1 0.494	1 13.00	2 024	0 797	
		1	1 1			VAP	10.491	1 274.89	1 0.933	45.80	0.876	0.363	0.1330
9	103.9	2.84	1 0.000	0 4204 1	-0.700	1.70	F I		1	1	1		1
		1	1 1	0.0208 1	-0.3201	VAP	10.150	133.11	0.502	12.55	2.192	0.423	0.7941
		1	1 1	. 1	i		1	210.00	1 0.720	44.04	0.886	0.722	0.1910
10	107.0	2.92	1 0.0001	0.0000 1	-0.1551	LIQ	10.135	135.51	0.506	12.47	2,280	0.445	0 8057
- 1		1	4 4	. /	1	VAP	10.395	276.81	0.925	43.28	0.892	0.726	0.2315
11 1	112.6	3.04	1 -5.8391	ຄ.ຄດກດ ເ	0.7681	1.10	10 400 I	4/0 00			1	1	
1	1		1 1	1	1	VAP	10.332	-140.08	0.513		2.444 1	0.489	0.7878
42	440.7		! !	1	1		i i			72.10	0.703 1	U./33	0.2431
14	117.3	3.16	-5.8391	0.0000 1	0.7181	LIQ	10.081	145.75	0.523	13.08 I	2.639 i	0.545	0.7747
i	i		i i	1		VAP	10.258	279.30	0.905	41.48	0.917	0.741	0.2252
13	127.8	3.28	1-32-8011	0.0000 i	6.7681	LIO	0.050	153.05	0.534	44 70 1	2 880 1		
1			!!	I		VAP	0.165 1	281.07 1	0.887	41.76	0.936	0.621	0.5625
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FIGURE H11 CONTD

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			Ň	1H= 50.84 I
* KW 1.466 2.426 1.572 2.528 0.011	0.000#		Ň	18= -4.97 I
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+CONCENTRATION 0.990 0.021 0.265		<u></u>	17	IH= 50.84
*		*****	· · · · · ·	18= -4.69
+ MINIMUM REFLUX FLOW (KG/S) = 0.0060	-			ISTATE=SUBCL
***************************************	*******			
*CAPNOT COP= -37.728 BASED ON 23.3, 21.4 & 115.3 DEG C	+			
+COP= 0 577 COPHEAT= 1.575 PUMP KW AS %= 0.4	+			
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NO	I TEMP I DEG C	I I Y	к 7/Х	IHEAT TF	IFEED FLOW 1 KG/S	IA/BY CH	I PHASE	I MASS	IENTHALP	YIENTROPY IKJ/KG K	AVAIL/TY	IFUG COEF	IFUG COEF	FLOW KG/S
1	1 29.7	1	.01	1 2.426	1 0.0000	-0.067	LIQ VAP	10.990	63.36 200.16	0.256 0.000	32.10 0.00	1 0.873 1 0.000	0.001	0.0177
2	 30.8 	11	.05	i 0.000	1 0.0000 1	 -0.003 	LIQ VAP	10.945 10.990	l 64.01 200.71	I I 0.265 I 0.709	28.27 35.92	0.897 0.870	0.134	0.0070
3	 33.4 	11	.15	0.000	1 1 0.0000 1	 -0.004 	LIO VAP	10.847 10.972	! 65.52 202.03	 0.281 0.716	21.51 34.58	I I 0.957 I 0.873	0.133 0.763	0.0067
4	 38.9 	11	.40	1 1 0.000	 0.0000 	i -0.008 	LIQ VAP	10.667 10.934	1 1 68.97 1 205.04	 0.303 0.729	11.36 32.29	 1.112 0.879	0.137 0.771	0.0063
5	 48.0 	 1 	.90	 0.000 	 0.0000 	 -0.009 	LÍO Vap	 0.457 0.869	 75.47 209.88	 0.327 0.748	1.89 29.23	I I 1.432 I 0.889	0.160 0.784	0.0060 0.0168
6	 57.0	1 2	.44	0.000	1 1 0.0000 1	 -0.005 	LIQ VAP	 0.327 0.797	 82.64 214.67	 0.347 0.764	-2.59 26.64	1 1 1.796 1 0.898	0.197 0.795	0.0060 0.0165
7	62.3	 2 	.75	1 1 0.000	 0.0417 	-0.020 -0.020	LIQ VAP	1 10.273 10.750	 87.00 217.35	 0.358 0.772	-3.99 25.31	 2.020 0.903	0.224 0.801	0.0500
8	68.5	 3 	.11	 -0.379	 0.0000 	0.028 	LIQ VAP	1 10.222 10.690	 92.29 221.09	 0.371 0.780	-4.95 23.93	 2.297 0.908	0.262	0.0484 0.0188
9	78.1	3	.64	 -0.379	0.0000	 0.125 	LIQ Vap	 0.161 0.586	 100.79 226.62	 0.391 0.790	-3.13 22.26	 2.748 0.918	0.331 0.816	0.0476 0.0172
10	91.2	4	.27	 -0.379 	 0.0000 	0.127	LIQ VAP	 0.100 0.428	 112.74 234.91	 0.419 0.798	0.10	 3.396 0.931	0.449 0.826	0.0478 0.0164
11	104.8	14	.78	 -0.379 	0.0000	 0.040 	LIQ VAP	 0.052 0.249	 125.61 244.48	 0.447 0.799	4.13 22.57	1 1 4.098 1 0.947	0.600 0.837	0.0486
12	115.3	 5. 	.05	 -2.528 	0.000	 0.573 	LIQ VAP	 0.021 0.107	 135.73 252.56	 0.468 0.791	7.80 30.47	 4.647 0.963	0.737 0.846	0.0312
NOTE	E A/BY	I CH	is	l l for stream	as only.	1		I	1	1 1		1	I I	

FIGURE H12 CONTD

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