THE QUANTIFICATION OF HAZARDS AT LPG INSTALLATIONS

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Master of Philosophy

THE UNIVERSITY OF ASTON IN BIRMINGHAM

October 1987

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SUMMARY

The literature relating to properties, utilisation, storage and transportation of LPG has been surveyed. Potential hazards were reviewed, particularly those related to fireballs arising from unconfined vapour clouds, boiling liquid / expanding vapour and pool fires. These confirmed the need for quantitative techniques for hazard evaluation and a critical analysis of such techniques, their applications and limitations was carried out.

The main parameters innvolved in evaluating thermal radiation hazards were: mass of fuel, equivalent spherical diameter, duration, height and surface temperature of the fireball.

Published empirical correlations for fireball size and radiation were compared with theoretical models and observations and the best fitting models were established, accurate to $\pm 5\%$.

Three new correlations have been proposed to fit published experimental data which relate radiation intensity on human skin to exposure time and to tissue damage, within $\pm 8\%$.

Thermodynamic considerations have been applied to calculate the theoretical flame temperature of a fireball, giving good agreement with previous models.

A proposed correlation for radiation from pool fires also shows reasonable agreement with published results. However, many factors for pool fires remain unpredictable, such as: shape and size of flame envelope, its temperature and emissivity.

Four methods to find acceptable spacing distances between vessels in hazardous areas are discussed. The results show spacing distance related to fuel released and to explosion yield factor for different damage overpressures. The maximum predicted spacing distances are to be preferred because of the relative unpredicability of unconfined vapour cloud explosions.

The quantitative methods discussed enable a reasonable assessment of major LPG hazards. The possibility of a major fire or explosion at an LPG installation may be minimised by recommended safety measures.

The available hazard evaluation methods have been collected, compared and their limitations assessed in a manner which should be useful to engineers and safety and loss prevention personnel.

Key words: Liquefied Petroleum Gas Fire, Explosion, Hazards I dedicate this thesis to the memory of my father, to my mother for her patience and to my husband for his support.

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CHAPTER ONE

INTRODUCTION

Liquefied Petroleum Gas (LPG) is a term used to describe a variety of hydrocarbons which exist as vapours under ambient conditions of temperature and pressure, but can be liquefied by the application of moderate pressure. LPG originates from oil or gas fields, or from refineries where the crude oil is processed. LPG is used mainly for heating or as a feedstock for chemical processes.

In recent years the rapid growth of operations involving the storage, transportation and use of LPG has resulted in problems related to environmental protection, energy conservation, safety and loss prevention. Despite the high standards of care taken in the design, construction and inspection of new and existing process plants, hazardous incidents still occur. The scale and variety of LPG operations have increased to an extent that fire and explosion incidents that occur may put not only the whole process plant and its personnel at risk, but also possibly a large number of inhabitants and buildings in the surrounding area. For example on November 19th, 1984 a disaster involving an LPG installation occured in Mexico City which resulted in the deaths of over 500 people and injury to 7,231, of whom 144 died in hospital.

The main types of hazard associated with LPG plants are:

i. Major fires: pool or flash fires can occur when the three conditions of fuel, air and ignition sources are available. Here radiant heat is the main effect causing burns to exposed persons, and a danger of secondary fires or explosions.

- ii. Explosions: these include;
 - (a) Confined Vapour Cloud Explosions (CVCE): when a fuel/air mixture is ignited within equipment or buildings, allowing the pressure to build up inside the confined volume until rupture of the container or collapse of the building wall occurs.
 - (b) Unconfined Vapour Cloud Explosions (UVCE): when a large fuel/air cloud burns in free space at a sufficiently rapid rate to generate pressure waves which propagate both through the vapour cloud and into the surrounding environment.
 - (c) Boiling Liquid Expanding Vapour Explosions (BLEVE): when a pressure vessel containing flammable liquid is heated so that the metal loses strength and ruptures. The effects of BLEVEs are thermal radiation caused by a fireball, a blast wave and missiles.

Many experimental and theoretical approaches have been published in an attempt to analyse and mitigate the above problems. However, the subject is extremely complicated; it is affected by a number of factors, eg the physical properties and conditions of the gases and liquid involved, the nature and type of storage and transport, the degree of hazard involved, the topography of the region considered, the atmospheric and weather conditions and the density of neighbouring populations.

Therefore, quantitative methods for the identification and assessment of hazards have always been used by the process industries to try to ensure that plants are reasonably safe. The need to quantify hazards in process plants arises because of the need to avoid losses and

costs that could cause business interruption and sometimes complete closure. Safety is an important factor in the appraisal of the acceptability of an industrial activity; it should be considered together with the economic benefits to the community.

The present work includes an evaluation of the hazard ranges for fireball and pool fires, to people, processes and buildings, based on the radiant flux densities which have an effect on human skin, buildings and processes. This study also includes predictions of spacing distances for plants involving UVCEs for which several methods have been proposed for the calculation of acceptable spacing.

The results computed from the various methods used have been compared. The maximum figure is preferred for conservative spacing and the design of mitigatory measures. The purpose of this study is to present an assessment of the main quantitative methods available to assess LPG hazards in the process industries and to draw together sufficient detail to be useful to engineers and safety and loss prevention personnel concerned with LPG handling and use.

CHAPTER TWO

GENERAL CHARACTERISTICS OF LIQUEFIED PETROLEUM GASES

2.1 DEFINITION

The term "liquefied petroleum gas" or LPG, refers to hydrocarbons obtained from petroleum refining operations (ie distillation, reforming, stabilisation), or from compression and/or absorption processing of natural gas and which are gases at normal atmospheric temperature and pressure, but can be readily liquefied with either a moderate increase in pressure or a moderate drop in temperature or both. These hydrocarbons including propane, butane, butylene and propylene and mixtures of these gases are the principal liquefied petroleum gases [1, 2, 4]. The more readily-liquefiable gases of this group are commercial propane and commercial butane, each of which contains several of the other hydrocarbons mentioned in varying amounts. Butadiene is also a liquefied petroleum gas; it is chiefly used in the rubber industry and not as a constituent of LPG fuels.

2.2 SOURCES OF LPG

The main sources of liquefied petroleum gases are natural gas the crude oil/gas mixtures that come out of actively producing oil and natural gas wells, which are removed as condensible products by compression and/or absorption processes, and from the crude oil stabilisation process which is applied in order to provide an appropriate vapour pressure for shipping. These sources are

supplemented by secondary supplies produced within the course of certain refining processes in oil refineries [1, 2].

The composition of liquefied petroleum gases varies according to both the conditions used in the refining processes and the type of crude oil used as feedstock

2.3 GENERAL CHARACTERISTICS OF LPG

The two principle liquefied petroleum gases in general use are defined in BSS 4250 and, from the principle constituents, are termed 'commercial' propane and butane. The contaminants usually comprise varying amounts of, for example, propylene and butylene plus some ethane and possibly some methane. Almost pure propane and butane can be obtained for special purposes.

The general characteristics of LPG are summarised as follows [2, 3, 4, 5]:

- a. LPG at normal temperature and pressure is a gas which is heavier than air. Butane is about twice as dense and propane one and a half times as dense as air at normal temperature and pressure.
- b. LPG is easily-liquefied under moderate pressure at atmospheric temperature. The density of the liquid is approximately half that of water.
- c. Pure LPG is non-corrosive to steel, copper and copper alloys.
- d. LPG is colourless in the liquid and vapour phases. However, any leak is often detected by the presence of frost due to the evaporation of the LPG which causes a local reduction in temperature and freezing of moisture from the surrounding air.
- e. LPG is odourless, so to assist detection a stenching agent (such

as ethyl mercaptan) is added to enable detection by smell at a concentration of one fifth of the lower flammable limit [3], except when the intended use requires a gas free from odour, as in the manufacture of aerosols.

f. LPG is non-toxic but has anaesthetic properties when inhaled in quantities for a long period; inhalation in moderate concentration produces nausea and headache, ie at concentration of 10% the vapour is not noticeably irritating to the eyes, nose or lungs, but will produce slight dizziness in minutes [4]. Table 2.1 lists the properties of chemically pure hydrocarbons [6], and Table 2.2 lists the properties of commercial propane and butane [4, 7].

2.4 HAZARDOUS PROPERTIES OF LPG

Nowadays the preference for gaseous fuel rather than liquid or solid, has produced situations in which the public may be increasingly exposed to potential risk of injury from fires or explosions. The hazardous properties of LPG are summarised as follows:

- a. LPG in the liquid phase has a low relative density, ie half that of water, and this has implications for the control of fire after large accidental spills, since it floats and will spread on water.
- A large quantity of vapour can be obtained from a small quantity of liquid (1 volume of liquid produces over 200 volumes of vapour).
- c. LPG becomes flammable when mixed with air in a concentration within the flammable range by volume, for example the flammability limits of LPG in air (by volume) are 1.5% to 10% [8]. So 4.5 litres of butane when vaporised will produce not

less than 900 litres of gas at atmospheric pressure and 60 °F (15.6 °C); thus at 5% concentration in air, 18.12 m³ of flammable mixture would be formed.

- d. LPG vapour is denser than air, ie propane vapour at ambient temperature having a density of approximately one and a half times that of air, while that of butane is approximately twice the density of air, will sink to the lowest possible level, possibly through drains and sewers, cable conduits or natural fissures and flow downhill. Unless dispersed such accumulation may persist for long periods with the possibility of subsequent ignition at a considerable distance.
- e. LPG has a calorific vlaue of 2 to 3 times that of natural gas, so heat radiation from LPG fires is relatively high.
- f. LPG in the liquid phase evaporates rapidly at ambient temperature causing severe frost "burns" when in contact with skin.

2.5 UTILISATION

The expanding field of utilisation of liquefied gaseous fuels has been largely reflected in the increased number of uses for which these gases have been found acceptable. Because of the large population areas not served by gas mains, ie without convenient supplies of natural gas, its utilisation reaches from the more common domestic uses to specialised commercial applications [3, 7]. The farm, camp or resort operator now has clean, convenient LPG fuel available for the domestic gas range, the gas refrigerator, the gas water heater and the gas fired space heater, to mention the four principal gas fuel applications. More extended uses include that for commercial refrigeration in preserving fruits and vegetables, gas engine power for sawing wood and motor fuel for the operation of trucks and tractors used on the farm, and on construction and road building projects.

Another rapidly-growing classification is the commercial and industrial application in which the portability of the LPG cylinder is important. These include those of the plumber, mechanic, tinsmith, and other tradesmen serving the home and industry [3].

The utilisation field in internal combustion engines is very broad. It includes both the stationary type of gas engine, such as that used for the generation of electric power, for waste pumping and for oil field use.

In the transportation field its use extends from the smallest to the largest trucks and buses and its application in air-conditioning machines for rail cars.

The high octane value of LPG permits its use in all types of gasoline and oil engines after these have been converted with special tanks, heat exchangers and carburettors. Reduced maintenance costs, increased economy and saving in crank case oil consumption are some of the advantages obtained in using propane or butane as fuel. So there are benefits both to the industry in utilising LPG, and to the environment because of the low sulphur content which almost eliminates the production of sulphur dioxide (SO₂) during combustion [3, 7].

Many uses for these gases have been developed in chemicals manufacturing. At the present time iso-butane is made into motor fuel by suitable processes. Special grades of propane, normal butane, isobutane and butadiene are being used in the manufacture of nitroparaffins, plastics, synthetic rubber and synthetic textiles.

In summary, wherever heat, air conditioning, power or refrigeration is required there is scope for the utilisation of LPG.

TABLE 2.1

PHYSICAL PROPERTIES OF LIGHT HYDROCARBONS [6]

	Methane	Ethane	Propane	Isobutane	Butane	Pentane
Molecular volume of gas, cu.ft. @ 60 °F†	378.7	375.8	372.7	366.7	365.40	54 T - 1
Molecular weight of gas	16.04	30.07	44.09	58.12	58.12	72.15
Gal./lbmole at 60 °F	6.4	# 9.64	10.41	12.38	11.94	13.71
Weight:						
% carbon	74.88	79.88	81.72	82.66	82.66	83.33
% hydrogen	25.12	20.12	18.28	17.34	17.34	16.66
Specific gravity:						
Of liquid (water = 1)	0.248	0.377	0.508	0.563	0.584	0.631
Of liquid, °A.P.I	340	1 247	147	120	111	93
Of gas (air = 1)	0.555	1.048	1.550	2.077	2.084	2,490
Weights and volumes:						
Lb/gal. liquid	2.5	† 3.145	4.235	4.494	4.873	5.250
Cu. ft. gas/gal. liquid	59.0	\$ 39.69	36.28	30.65	31.46	27.67
Cu. ft. gas/lb. liquid	24.8	12.5	8.55	6.50		
Ratio, gas volume to liquid volume§	443	293.4	272.7	229.3	237.8	207.0
Initial boiling point (atmospheric pressure)	-259	-128.2	-43.7	10.9	31.1	97
Heat value (gross):						
B.t.u./cu. ft. gas	1,012	1,786	2,522	3,163	3,261	4,025
B.t.u./lb. liquid	23,885	22,323	21,560	20,732	21,180	21,110
B.t.u./gal. liquid		70,210	91,500	103,750	102,600	110,800
Vapour pressure, lb/sq. in. abs						
At -44 °F		88	. 0	-9	-12	-14
At 0 °F		206	38	12	-7	-13
At 33 °F		343	54	17	0	-11
At 70 °F		563	124	45	31	-6
At 90 °F		710	165	62	44	
At 100 °F		-	189	72	52	4
At 130 °F		-	275	110	81	11
At 150 °F			346	138	87	21
Latent heat of vaporisation at boiling point:						
B.t.u./lb.	221	211	185	158	167	153
B.t.u./gal.	553	664	785	742	808	802
Specific heat:						
Of liquid, Cp at 60 °F., B.t.u./(lb.)(°F)		0.780	0.588	0.560	0.549	
Of gas, Cp at 60 °F., B.t.u./(lb.)(°F)	0.526	0.413	0.390	0.406	0.396	0.402
Of gas, C _v at 60 °F., B.t.u./(lb.)(°F)	0.402	0.347	0.346	0.373	0.363	0.376

† Ideal gas = 379.5 cu. ft. @ 60 °F or 15.5 °C
Apparent values for dissolved methane at 60 °F
§ Based on "perfect gas".

TABLE 2.2

TYPICAL PROPERTIES OF COMMERCIAL LPG GRADES [4, 7]

Properties		Commercial butane	Commercial propane
Relative density o at 15.6 °C (60 °	f liquid F)	0.57 to 0.58	0.5 to 0.51
Imperial gallons/to	on at 15.6 °C	385 to 393	439 to 448
Litre/tonne at 15.6	5 °C (60 °F)	1723 to 1760	1965 to 2019
Relative density of with air at 15.6	f gas compared C and		
Volume of cos (it		1.90 to 2.10	1.40 to 1.55
liquid at 15.6 °C	and 1015.9 mbar	406 to 431	537 to 543
Volume of gas (ft ³ 60 °F and 30 in 1) per lb of liquid at Hg	6.5 to 6.9	8.6 to 8.7
Boiling point (°C) pressure	at atmospheric	-2 approx	-45 approx
Vapour pressure (h at their maximum	pars) for products a specified vapour		
pressure. Temp	-40 °C	-	1.38
	-17.8 °C		3.11
	37.8 °C	1.95	5.52
	57.0 C	5.86	15.5
	13 C	0.89	18.6
Latent heat of vapo	risation (kJ/kg)		
at 15.6 °C		372.2	358.2

Cont'd/ ...

TABLE 2.2 - Continued/...

Properties	Commercial butane	Commercial propane
Latent heat of vaporisation (Btu/lb)		
at 15.6 °C	160	154
Specific heat of liquid at 15.6 °C (kJ/kg °C)	2.386	2.512
Specific heat of liquid at 60 °F (Btu/lb °F)	0.57	0.6
Sulphur content, percent weight	Negligible to 0.02	Negligible to 0.02
Calorific values		
Gross: MJ/m ³	121.8	03.1
MJ/kg	49.3	50
Net: MJ/m ³	112.9	86.1
MJ/kg	45.8	46.3
Therm/ton (gross CV)	475	485
Air required for combustion		
(m ³ to burn 1 m ³ of gas)	30	24
Limits of flammability (percentage		
by volume of gas in a gas-air		
mixture to form a combustible		
mixture)		
lower flammable limit	1.8	22
upper flammable limit	9.0	10.0

CHAPTER THREE

STORAGE, TRANSPORTATION AND MODES OF CONTAINER FAILURE

3.1 STORAGE SYSTEMS

Liquefaction of petroleum gases under certain conditions of pressure and temperature permits convenient and econmical storage and transportation of large quantities of these gases.

There are two types of storage:-

3.1.1 Pressurised Storage:

Light gases such as propane and butane liquefy readily when subjected to a pressure equal to their respective equilibrium vapour pressure; this pressure varies according to temperature [2, 3, 4].

		Tempera	ture °C
		15.6 °C (60 °F)	37.8 °C (100 °F)
Equilibrium	Butane	103 kPa (15 psig)	483 kPa (70 psig)
vapour pressures	Propane	518 kPa (75 psig)	1400 kPa (210 psig

In practice the commercial grades of LPG are not pure, being contaminated with small amounts of other low molecular weight hydrocarbons. In the storage of commercial grades of LPG, it is necessary to take into consideration that the equilibrium vapour pressures of commercial grades of propane and butane are somewhat higher than those for the pure gases. This is illustrated in Figure (3.1).

Pressure storage tanks must therefore be designed and constructed to withstand the highest vapour pressure encountered in use. This will depend upon the highest temperature to which the contents will be elevated in use, usually by solar heating.

Static installation tanks are usually constructed in accordance with a relevant standard such as BSS 5500, ACTC Rules and others. The design pressure must not in any case be less than 489 kPa for butane and 1470 kPa for propane.

Pressure storage tanks are fabricated from special steel and are inspected for soundness by a variety of sophisticated techniques for incipient cracking and full weld integrity. It is usual to inspect them at about five yearly intervals in service.

Occasionally pressure storage tanks at fixed installations are buried below ground to eliminate fire impingement and solar heating upon the shell. These are given external protective coatings to resist soil corrosion, and in addition cathodic protection is employed. It is important to ensure that such tanks are securely fixed since ingress of water around even full tanks will produce severe flotation induced stresses which could rupture associated piping. Indeed, any associated piping needs to be of high integrity and well-supported and protected.

These submerged tanks may be more expensive but are a viable proposition in congested areas where spacing requirements cannot be met.

Above ground pressure storage tanks at fixed installations are spaced to provide access for fire fighting [9] and to avoid the spread of fire from tank to tank. The tanks are usually elevated above the



Figure 3.1 Approximate Maximum Vapour Pressures for Commercial Propane and Butane

ground and the ground beneath them is concreted, and sloped away from the tank, to prevent accumulation of spilled LPG below the tank. Such installations are sometimes fitted with water drench systems intended to cool the shell of the tank in the event of a fire below, or of high radiant heating from adjacent fires. Such above ground tanks are often grouped in a fenced compound, to prevent trespassing or tampering. Sabotage is a very real threat.

All large scale pressure storage vessels are fitted with springloaded pressure relief valves set to vent vapour to atmosphere in the event of an undue pressure increase. Ideally the valve capacity should be such that it will cope with the maximum vapour production at the reference temperature plus the vapour production under fire conditions. Installation of a relief valve is not, unfortunately, sufficient to guarantee tank integrity in the event of fire; there are instances on the record of valves sticking shut.

These tanks need a variety of safety devices in addition to relief valves. All openings should be equipped with excess-flow valves that prevent uncontrolled product flow to atmosphere, in the event that piping connected to a tank opening is ruptured or broken-off. Some high pressure tanks are equipped with rupture disks which permit the vapour to escape at a greatly accelerated rate.

A precaution which must be taken into consideration for relief valves and rupture discs is that connections should be equipped with stacks so that discharged vapour will not strike personnel. These stacks should be located so that they will not vent into enclosed spaces or towards air intakes or sources of ignition.

Small scale storage of LPG is always under pressure in containers as diverse as the small disposable cartridges used for

fuelling cigarette lighters, to the 100 kg cylinders, ie "bottles".

Cylinders are tested to the minimum pressures for commercial propane of 2200 kPa and for commercial butane of 750 kPa, and are provided with radically different coupling threads to prevent butanerated equipment from being inadvertently fitted to a propane cylinder, and vice-versa.

Cylinders are subjected to visual examinations to check for any mechanical damage and that they are leak-free prior to filling. However, the accident record is replete with instances of cylinders attached to portable appliances being responsible for massive damage due either to slow leakage and accumulation and subsequent ignition of explosive levels of gas within buildings or by catastrophic failure when subject to fire. To counter this last threat, most cylinders are now being fitted with relief valves to relieve internal pressures caused at elevated temperatures.

The common configuration of storage tanks [10] as illustrated in Figure 3.2 and 3.3 are:

- Cylindrical these pressure tanks range in size from 4.35 kg "bottles" to approximately 165,000 gal "blimps" and in pressure from 200 bar (in small size only) down to 2 bar. The cylindrical type of tank is the most commonly used since it is economical to manufacture, of convenient configuration for movement, and can be mounted on single foundations using conventional erection procedures. It can be designed with good integral structural strength. Rail tank cars, transport trucks and barge tanks are invariably cylindrical.
- Spherical these tanks are usually fabricated at the erection site. They are subject to the same limitations in size as horizon-











tal cylindrical tanks. Their principle advantage is that they represent a minimum expenditure on steel. Segments of spherical tanks can be formed at the factory and stacked for shipment; they are the favoured type for the export trade because of space saving during transportation.

iii) Spheroid - these tanks are generally used in the vapour pressure range below 15 psia. They can be made in very large sizes. Configurations may involve combinations of shapes such as the sphere/torus type and the more oblate spheroid type.

3.1.2 Refrigerated Storage

Butane and propane can easily be liquefied by cooling below their respective boiling points. When chilled as liquid they exert little pressure upon the container which has advantages for engineering purposes. The tank can be of relatively light construction and there are no construction material problems such as the cryogenic-shock associated with liquefied natural gas (methane) which has a boiling point of approximately -160 °C.

Refrigerated tanks are fitted with a pressure relief valve to relieve the increase in internal pressure following failure of the refrigerating system, or fire impingement.

It is usual to insulate such tanks heavily with a variety of materials: expanded polystyrene, rock wool and perlite are common. Any insulation material needs to fulfill a number of criteria, such as [3]:

- a. withstanding impingement of water jets.
- b. resisting the ingress of water vapour
- c. maintaining effectiveness even if mechanically damaged.
d. resisting fire, either intrinsically or behind protective cladding.

Refrigerated storage tanks are constructed with an outer cladding, such as aluminium sheeting, to protect and contain the insulating material, which may be of the order of 900 mm in thickness; the interspace may also contain inert gas. Because the insulation and refrigeration systems add greatly to the complexity of such storage, refrigeration is usually only applied for large static installations of 2000 tonnes or more, or for bulk carriage by sea [2].

The Canvey Report [11] has identified a potential hazard from large refrigerated butane storage tanks that is of interest. Referring to British Gas refrigerated tanks of 5000 and 10,000 tonnes capacity, it points out that these are used near atmospheric pressure and were designed primarily to withstand the head of liquid. The mode of failure of such a tank when containing butane is diametrically opposed to that of an identical tank containing oil. If such a tank, containing oil, were to develop a horizontal split at a welded seam, then the weight of the upper part of the tank would tend to keep the split closed. Relief valves for refrigerated storage are set to quite low values, a few centimetres of water only; however this slight positive pressure in the same type of tank, but containing butane, would cause the top of the tank to lift and the split to gape further open. Thus the outflow of the tank contents would increase and possibly engulf the surroundings with consequent dangers.

Although bunding is not recommended around pressurised storage tanks or semi-refrigerated storage tanks, creating as this would the potential for a large pocket of explosive vapours, they are spaced at appropriate separation distances [12]. In general bunds are

provided for atmospheric storage tanks and for fully refrigerated storage tanks to contain spilled refrigerated LPG. This has the effect of limiting heat input to the spilled liquid, reducing the rate of vapour generation and subsequent formation of a large explosive vapour cloud It is also argued that such containment of a large spill is [12]. desirable to facilitate 'mopping up' operations, where the liquid could be transferred to road or rail tankers and thus be contained under pressure conditions as the LPG liquid subsequently warmed to ambient temperature. So it is possible to envisage circumstances under which such bunding would be a positive disadvantage in the event of tank leakage. Should such a spill ignite there, a contained and possibly sustained fire would ensue at the tank wall, and the light cladding used to protect the insulation would probably fail. Heat input into the tank contents could then generate vapour in excess of relief valve capacity and perhaps lead to tank failure.

Furthermore the presence of a surrounding pool of cold butane would preclude the use of water to cool the tank wall, as this water would perforce drain into the bund. Experiments have shown that under these circumstances water freezes to form an ice slurry which, being denser than the butane, sinks to the bottom of the pool. In large quantities this would cause displacement of the butane onto the surrounding site, over the top of the bund. This would increase the rate of evaporation and the incident would escalate rapidly in both size and seriousness [3].

Some practical aspects of the design of bunds for flammable liquids have been discussed by Hearfield (12). One fact which should be noted is that a leak in the side of a tank may form a horizontal jet which may jump the bund if the latter is too close to the tank. Therefore the bund wall should be far enough from the side of the tank

to prevent a jet jumping over or, alternatively, the bund should be surrounded by an impervious surface sloped inward to the bund drain area.

The corners of the bund should be rounded and not at a right angle, since it is difficult to extinguish a fire in a 90° angle corner because of the air compression effect.

The bund should be adequately drained, with a valve on the inlet end of the drain but outside the bund wall. Walls for full bunding should not be so high as to hinder fire fighting. The ICI LFG Code suggests a maximum height of 2 m with an allowance of 0.15 m freeboard above the predicted liquid level. This leaves room for a foam blanket. There should be a minimum of two access points on opposite sides of the bund to allow safe access in all wind directions. Access may be provided by steps over the bund.

Recommendations for pressure storage vessels vary. Hughes recommends that LPG vessels smaller than 30,000 UK gal (136 m³) should be surrounded by a layer of gravel only, while vessels larger than this should stand on a concrete apron sloping down to a catchment area again provided with gravel to assist vaporisation. A low protective wall 0.4 m high may be provided if considered necessary.

The HSE recommendations for LPG are that there should be low walls not exceeding 0.61 m around the vessels [12]. These should be built sufficiently far from the vessels to allow free circulation of air, the minimum distance being 1.5 m even for small installations. The walls should not be designed to retain any specific quantity of liquid.

An unusual hazard from refrigerated LPG storage may occur when a large refrigerated tank is used to store high molecular weight

hydrocarbon such as pentane in a warm climate [3]. In this case, refrigeration is necessary to keep the vapour pressure at a low value although the ambient temperature is high. However, in a reported incident when the refrigerant line started to leak the low molecular weight hydrocarbon refrigerant into the pentane this led to an increase in the vapour pressure to a value which was sufficient unduly to stress the tank, due to the fact that the relief valve was "stuck shut". In this case the tank attempted to become spherical and eventually the top blew off. In the description of this incident a cloud of vapour rose 45 to 60 metres in the air and then, being denser than air, as are propane and butane, settled back and spread over the surrounding area. After thirty seconds, or so, the cloud found an ignition source and a large fire storm ensued.

This incident may point to the need to ensure that all refrigerated storage of LPG utilises as refrigerant the same material as that to be stored.

3.2 TYPE OF CONTAINERS

3.2.1 Cartridges

Disposable cartridges are used in large numbers, largely in the leisure field, in content of weight of up to approximately 0.45 kg [7]. The major disadvantage of this type of cartridge is that it cannot be removed from the appliance until it is exhausted, otherwise the gas in its liquid phase may flash vaporise and may result in cold burns, or fire if a source of ignition is present.

3.2.2 Cylinders

Refillable cylinders are used mainly for domestic and light

industrial applications and range in content weight from approximately 5 kg to 46 kg capacity. They are designed specifically for either butane or propane due to the considerable gauge pressure differences:

Butane: 2 bar (28 psig)

Propane : 7 bar (100 psig)

The fittings on these cylinders are not interchangable. These cylinders are pressure vessels and therefore have to be tested periodically before and during usage. The first retest is required 10 years after the date of manufacture with subsequent retest required at 5 year intervals. These retests may be accomplished by visual inspection procedures as prescribed by the ICC [13]. Each cylinder should bear a manufacturer's serial number, year of manufacture, the specification to which is was built; the date of test, the test pressure, and the minimum hydraulic capacity of the vessel. The cylinders are visually examined prior to filling and the valve checked for leaks with soap solution.

3.2.3 Bulk Tanks

Storage at user's premises normally consists of individual tanks ranging from under one tonne to over 100 tonnes capacity. At a small number of premises with large consumption and at refineries much larger tanks may be found. Most LPG tanks are cylinders with dished ends, although some small tanks and most very large ones are spherical in shape.

All the cylinders and bulk vessels referred to above must be physically adequate for the stresses involved. They must be gas tight, be provided with safety relief valves where necessary, and be marked for ready-identification. Their design, characteristics, testing, inspection, retesting, handling, and siting when stationary are all

covered by established codes of practice, such as those issued by the LPGITA. Standards, or regulations, some of which are mandatory.

Before embarking on any project involving the installation of LPG facilities, it is important to ensure that any statutory requirement in respect of such installations will be compiled with. A check should be made that the proposed installation will comply with the requirements of all appropriate bodies, such as in the UK, the Factory Inspectorate, who administer the Highly Flammable Liquids and Liquefied Petroleum Gases Regulations 1972 (BI 917) made under the Factories Act 1961, and the Local Fire Authority.

In addition to conforming to any statutory requirements, it is advisable to be guided by appropriate Codes of Practice. In this connection a number of codes have been prepared by the LPGITA (UK) covering both LPG installation and the utilisation of LPG [4].

3.3 TRANSPORTATION

Producers and consumers are, in most instances, separated by considerable distances, since the production plants are located principally in the oil and gas producing regions of most countries, while the potential markets extend to sections far removed from these points. Hence large quantities of propane and butane must be transported over long distances by safe and economical methods.

For several years the only facilities for transporting these liquid gases were small metal cylinders, commonly called "bottled gas" cylinders, and only those customers who were reasonably close to the source of supply could economically use this type of fuel.

Liquefied petroleum gases are transported in much the same manner as other petroleum products, except that the metal containers

must be of sufficient strength to withstand the vapour pressure resulting from certain maximum atmospheric temperatures that will be encountered. The containers must also be equipped with safeguards as called for by local regulations.

The pressure within the container increases with the temperature of the liquid, which in turn is governed by heat transfer from the outside temperature. This heat transfer is relatively slow, so tank temperatures, rarely approach maximum atmospheric temperatures, and are usually lower by -12 °C (10 deg F) or more [13]. LPG is transported in varying quantities by road, rail, ships and pipelines.

Road tankers are used for transporting bulk loads between 1 and 20 tonnes, and railcars carry individual bulk loads of 10 to 90 tonnes. Tanks of more than 5 m³ capacity are fitted with internal baffles to minimise internal surging of the contents. The maximum capacity of the tankers for use on the road in the UK is determined by the current Motor Vehicle (Construction and Use) Regulations, and by ullage (air space) needed.

Liquefied petroleum gas has a relatively high volumetric coefficient of expansion and, therefore, any container should be filled to a limit which permits liquid expansion due to normal rises in temperature without danger of overstressing the container by hydraulic pressure. So it is necessary to ensure that the maximum amount of LPG in any road tanker should be such that the tank will be not more than 97% full with the contents at the highest temperature likely in service, this is termed the Reference Temperature for Filling Ratios and is currently in the UK 42.5 °C for tanks of 5 m³ water capacity and below, and 38 °C for tanks over this [14]. The maximum permitted LPG filling by weight is given by [14]:

$$W_1(kg) = 0.97 \text{ x } V_1 \text{ x } \rho_{(ref)}$$

.... (3.1)

where: V_1 = water capacity of vessel in litres $\rho_{(ref)}$ = the relative density of LPG at the reference temperature for filling ratio (RTFR)

In any case, the tank should not be filled so that a liquid full condition can arise at a temperature 5 °C in excess of the RTFR [14].

At low ambient temperature there may be a fall in the LPG vapour pressure to a value below that of the atmosphere. Therefore the tank must be designed to withstand this unaccustomed external stress, or alternatively be equipped with some system to relieve the vacuum. It is usual to assume for this purpose a design temperature of -10 °C, when the vapour pressure of n-butane is about 35 kPa below atmospheric.

Tanks for road transport are frequently used to carry a variety of materials. If a tank is so used, it is essential to ensure that it is suitable for all the substances handled.

Some important features of road tankers which should be considered are:

- i design basis,
- ii material of construction,
- iii reference temperature,
- iv minimum wall thickness,
- v safety valves, and
- vi attachment to vehicle

LPG is transported by pipelines which are usually buried about a metre below ground. The older pipelines had no specific external protection, then the wrapping of lines was developed, culminating in the present practice of using a coal tar/glass fibre wrap [12]. But corrosion still occurred at points where the wrapping was badly done or was damaged or where there were line joints. This problem has been largely overcome by the use of cathodic protection. Minimum external corrosion rates are obtained with the new method of external protection which utilizes a combination of wrapping and cathodic protection.

Coastal and ocean going ships with multiple pressure storage tanks of up to 3000 tonnes total LPG capacity are in service, and refrigerated low pressure storage tankers of 3000 to 4000 tonnes LPG capacity have recently been introduced.

Hazards presented by the transport of hazardous material are mainly fire and explosion. Fire is more likely than explosion where there is a loss of containment of flammable material from a rail, barge or ship tank or from a pipeline. But Unconfined Vapour Cloud Explosions and Boiling Liquid Expanding Vapour Explosions may occur as a result of transport accidents.

Road accidents leading to the release of the flammable material are due in most cases to a tanker crashing or overturning.

Rail accidents are mostly due to train crash or derailment. Mechanical failures of rolling stock and faults in the rail track are frequent causes of these. Another hazard of railways is a fire along the track caused by engine or brake shoe sparks.

The main cause of failure of pipelines, besides corrosion, is interference. In particular damage occurs to pipeline from earthmoving and excavating equipment.

Finally the hazards associated with ships are fire or explosion

on the ship itself, or spillage of flammable material from the ship.

Spillage at sea is likely to be the result of collision or stranding. Factors which increase the probability of collision are congestion of shipping lanes, fast steaming and difficult weather conditions.

The effects are limited to the ship itself or occasionally to other ships nearby if a spillage of flammable liquid occurs well out to sea. However, if it takes place near the coast, particularly in port, people on land may be endangered.

3.4 MODES OF CONTAINER FAILURE

Failures of LPG storage systems and the subsequent release of quantities of energy by fires and explosion have lead to massive losses of life and property. The actual potential of such fires and explosions are examined and quantified in Chapter 4; however, as the initiating failure of a storage system is of paramount importance in determining the ultimate scale of destruction, these failures are considered here. Assessment of the historical record allows identification of several modes of failure.

It is also possible to identify other failure modes that are feasible but have yet to be realised.

3.4.1 Spontaneous Structural Failure

In any large fabricated steel structure, there are inevitably small defects consisting usually of cracks up to 2 cm long by 0.5 cm deep [11]. These may be due to poor weld fusion, or may be due to a high cooling rate leading to embrittlement and crack generation by residual stresses in the so-called heat affected zone. These cracks grow slowly while the tank is in service, especially with a pressure tank, where

there are many changes of stress as the tank adapts to varying load patterns. The longer the crack becomes, the faster it grows; if it is not repaired it can either lead to leakage or convert suddenly to a fast fracture mode allowing the tank to rupture suddenly with the generation of fragments.

3.4.2 Corrosion

The overall chemistry of the rusting process of iron and steel is complex, with the eventual formation of a complex of iron oxides and peroxides on to the surface of the corroding metal in the presence of water and a dilute acid. However, the first stage of the rusting process is the formation of ferrous ion according to the equation

$$Fe (s) + 2H^{+} (aq) = Fe^{2+} (aq) + H_{2} (g) \qquad \dots (3.2)$$

to rust via
other reaction ---->

This can be further simplified as:

$$Fe(s) = Fe^{2+}(aq) + 2 electrons$$
 (3.3)

Rusting tends to occur in that part of an iron or steel object which is under some stress, producing an 'anodic' site relative to the remainder of the system.

LPG storage vessels, especially those buried below ground, are extensively protected against corrosion by both protective coating and by the use of cathodic protection. Examination of expression (3.3) allows an explanation of the mechanism by which cathodic protection operates. If the production of ferrous ions can be inhibited by supplying more electrons thus

$$Fe^{2+}(aq) + 2e^{-} = Fe (metal)$$
 (3.4)

then corrosion will not take place whilst the supply of electrons lasts. In practice, this electron source is usually provided by the use of a metal such as magnesium or zinc, in close contact with the vessel to be protected, and which is a better reducing agent than iron or steel. The overall chemistry is

$$Mg(s) + Fe^{2+}(aq) = Mg^{2+}(aq) + Fe(s)$$
 (3.5)

and in electrochemical terms, iron and magnesium are considered to form an electrochemical cell, with water as electrolyte. Within this notional cell, magnesium acts as the anode and iron as cathode (hence 'cathodic' protection) and the magnesium ions are lost in the water. So long as the iron continues to act as the cathode, reaction (3.3) cannot proceed [3]. The overall process is summarised in Figure 3.4.

3.4.3 Mechanical Damage

A variety of interactions have been identified which may cause mechanical damage to LPG storage vessels severe enough to cause loss of containment [11]. In the overwhelming majority of cases on record it is the damage to associated piping, caused for instance by vehicle impact, that causes this category of damage. Once containment is lost below the liquid level then massive pressure - driven discharge of the contents will ensue.



Stage 1 : Iron atom donates two electrons to hydrogen ions in solution in electrolyte. These are replaced by electrical conduction from a magnesium atom.



Stage 2: The magnesium ion is now soluble in the electrolyte and is lost into solution. The two reduced hydrogen atoms combine to form a hydrogen molecule, which is lost as gas from the electrolyte. There is no loss of iron; therefore corrosion has been inhibited.

Figure 3.4 Mechanism of cathodic protection against corrosion

The course of events for mechanical damage to LPG storage is summarised below:



Damage to refrigerated storage tanks is more likely to occur due to their lighter construction. However, the lower rate of vaporisation for refrigerated LPG tends to limit the consequences of such a spill, providing that the spill is contained within a relatively small area to limit the input of heat. However, if the secondary containment were lost, then spread of liquid onto the surrounding site would be most serious, as unlimited heat would effectively be available to produce vaporisation [3].

3.4.4 Flame Impingement

When flames contact the vessel below the liquid level, the liquid serves to absorb the heat and tank metal below this level therefore remains at a safe temperature. But with continued relief valve operation, the liquid level will drop exposing a greater area of metal to the effects of heating. Flames contacting the tank above the liquid level can create temperatures within the metal high enough to weaken it seriously. When this happens the pressure within the tank can cause the metal to thin and eventually tear.

Therefore, if flames impinge upon LPG storage tanks, two consequences may arise:

- there is an increase in internal pressure as the contents increase in temperature.
- the strength of the steel container decreases at elevated temperatures.

The heating is due in most cases to leakage of hydrocarbons under the tank, either from the same tank or from a neighbour. In theory the relief valve should vent the tank as the temperature rises, but in practice there have been cases of them sticking closed. In any event, the venting capacity may be inadequate and the pressure will be certain to increase. When tanks explode under these conditions, they are said to BLEVE. This is derived from the expression Boiling Liquid Expanding Vapour Explosion and is most simply described as the sudden rupture of a pressure vessel containing a flammable liquid above its normal boiling point, due to a fire surrounding the vessel. Such fires also lead to progressive weakening to the point of rupture. The main consequences of a BLEVE which are discussed in detail in Chapter 5 are:

- 1. intense transient radiation from the fire ball following a ground flash.
- limited blast as has been reported as severe blast wave damage from the BLEVE at Feyzin refinery [15].
- 3. missile generation, possibly involving large fragments of the

tank, being projected considerable distances. It has been reported in the Mexico disaster [15] that 15 of 48 cylindrical tanks each weighing at least 20 tones were trasnformed into missiles and flew over 100 m. Those travelling further landed 1000 or 1200 m away from the scene and destroyed everything in their path, including several houses.

There are many BLEVE incidents [15] in the accident record as shown in Table 3.1 concerned with rail and road tankers. Mississauga is an example of a rail incident which is described briefly in Table 3.2.

3.4.5 Blast Damage

One danger arises from the 'domino effect' where an intact LPG tank in close proximity to some initiating explosion is itself damaged, releasing additional fragments, LPG, or light hydrocarbons into the existing fire situation. In practice, LPG tanks are usually either spherical or cylindrical and are highly resistant to catastrophic failure from blast waves. However, there is a very great threat in the fact that blast wave energies are sufficient to actually move a tank on its mountings, thereby rupturing associated piping [3].

3.4.6 Missile Interactions

Missile interactions can perhaps be considered as a special case of mechanical failure, and an extension of the previous failure mode. Explosions in process vessels, storage vessels or buildings can be expected to produce a range of missiles, with different masses, travelling at various velocities. Upon impact with an LPG storage vessel there is a wide range of possible outcomes, ranging from elastic collision with little or no damage, to complete penetration of the vessel

TABLE 3.1

BLEVE INCIDENTS

Date	Country	Accident
Jan 4, 1966	Feyzin, France	Explosion of tanks at refinery. 18 fatalitites. 40 injuries
Jul 5, 1973	Kingman, Arizona, USA	BLEVE of tank car caused by explosion and fire of LPG distribution plant. 13 fatalities. 96 injuries.
Jun 21, 1970	Crescent City, Illinois, USA	Derailment and explosion of tank cars. 66 injuries.
Feb 9, 1972	Tewkesbury, USA	Collision tank vehicle with pipelines causing BLEVE of storage tank, 2 fatalities. 21 injuries.
Mar 3, 1980	Los Angeles, California, USA	Overturn and explosion tank vehicle loaded with gasoline. 2 fatalities. 2 injuries.
Jul 23, 1984	Illinois, USA	Explosion and fire at refinery. 15 fatalities. 22 injuries.
Nov 19, 1984	San Juanico, Mexico City	Series of explosions and fires. 500 fatalities. 7000 injuries.

TABLE 3.2

EXAMPLE OF A RAIL INCIDENT INVOLVING LPG (MISSISSAUGA DISASTER)

SCENARIO 11.53

SATURDAY, NOV 10, 1979

Train 54 of CP Rail suffered a derailment in Mississauga.First car derailed was a toluene tank carTwenty-three cars went with it (21 tank cars)

Involved 11 propane

- 4 NaOH
 - 3 toluene
 - 2 fiber glass insulation
 - 3 styrene
 - 1 chlorine

Fire spread throughout most of derailed cars.

- 12.15/12.20 8th, 12th and 13th propane tank cars BLEVE'd. 7th car containing $Cl_2 - 2^{1/2}$ feet hole in shell.
- 1.30 Readable copy of manifest obtained.
- EVACUATION Almost 250,000 people for periods up to 5 days.
- CAUSE The toluene car suffered a 'hot-box' in its right-rear journal box. The rear-axle and its two wheels separated from the truck and the car 1 mile before derailment.
- DAMAGE 3 cars BLEVE'd within 30 minutes with 3 great fireballs.

Large parts of bodies projected:

8-145 feet east 12-440 feet southeast 13-2222 feet northeast (other parts in all directions)

75,000 remained evacuated until Friday, 16th November.

wall and consequent release of the contents. It has been reported [16] that penetration by fragments of other tanks was responsible for the subsequent BLEVE's in Mexico's disaster.

3.4.7 Sabotage

The location of large amounts of stored energy in a relatively small space allows massive destruction by initiating fires or explosions by acts of sabotage.

The tanks should be enclosed to protect them against damage, a strong wire fence should be put up instead of a solid wall, which could contain any explosive mixture [1], and this security fence should be at distance of 15 m or more from the tank [2]. More stringent security measures may be appropriate if sabotage is a possibility.

The GAO Report recommended [3] that LPG sites should be situated in remote areas, and such siting is prudent not only on the grounds of public safety, but also on the basis of reduction of the propaganda value to terrorist organisations from destruction of such sites.

CHAPTER FOUR

CONSEQUENCES OF SYSTEM FAILURES AND THE HAZARDS INVOLVED

In the previous chapter, the systems of LPG storage and transport were reviewed together with various failure modes to which they are liable, some of which are probable and some remote. In each case of failure the end result was envisaged as a loss of containment. This would result in either a slumping vapour cloud or a jet depending on the mode of failure as discussed in 3.1 and 3.3.

It is now pertinent to consider the further consequences of these losses. The leak or cloud may of course disperse harmlessly (i.e without ignition or contact with personnel) depending on the circumstances, for example being well mixed by wind which may result in a gas concentration below the lower flammable limit so that the mixture is 'lean' with insufficient gas to support combustion or the ignition source might be well away from the plant in a harmless area. The consequences of releases of LPG are summarised in Figure 4.1. The diagram identifies a number of consequences which must be considered as serious hazards.

4.1. HAZARDS TO PERSONNEL

4.1.1 Cold "Burns"

When any liquid vaporises the heat required to effect the evaporation is drawn from the surroundings, which are cooled as a result to a lower temperature. Where the rate of evaporation is rapid,



FIGURE 4.1 FLOW CHART OF CONSEQUENCES OF LPG FAILURES

as when a liquid is above its boiling point at ambient temperature (the case with a release of LPG), severe cold "burns" may result from skin contact.

4.1.2 Asphyxiation

Propane and butane are not 'toxic', in the normal sense of the word but possess anaesthetic properties, as do all petroleum vapours, and should not be inhaled in quantity. Occupation exposure limits are recommended by the Health and Safety Executive on advice from HSC's Advisory Committee on Toxic substances (ACTS). These limits are considered to represent good practice and realistic criteria for the control of exposure, plant design, engineering controls and if necessary, the selection and use of personal protective equipment.

It is important to control exposure so as to avoid both short term and long term effects. The long term exposure limit is concerned with the total intake over long periods and is therefore appropriate for protecting against the effects of long-term exposure or reducing the risks to an insignificant level. The short term exposure limit is aimed primarily at avoiding acute effects, or at least reducing the risk of their occurence. Both limits are expressed as time weighted average (TWA) concentrations which are simply airborne concentrations averaged over a specified period of time. The period for the long term limit is normally eight hours, while the short-term exposure limits are normally expressed as ten minute TWA's. The occupational exposure limits (threshold limit) for butane is 600 ppm at long term exposure limit and 750 ppm at short term exposure limit [17]; suprisingly no limit is stated for propane.

A large concentration of vapour in a confined space will displace air to an extent that there could be a deficiency of oxygen for normal respiration. At least 18 % oxygen is required in air, e.g in a confined space, for workers not wearing breathing apparatus. Concentrations liable to cause serious oxygen deficiency would always be in (or above)the flammable range and constitute a serious fire and explosion risk.

4.1.3 Thermal Radiation Effects

The effects of fires on human beings increase with the heat flux and exposure time. There is however, a threshold flux which

can be sustained for a long period without causing serious injury. This flux was suggested to be 5 kW/m² [11].Further study [18] suggests that this value should be reduced slightly to 4 kW/m². For shorter exposure times (as might be experienced near a fireball),the 'blistering of skin' criteria have been adopted by the Safety and Reliability Directorate (UKAEA). Much lower intensities can cause problems with the eyes by the focussing of radiation on to the retina.

TABLE 4.1

RELATION BETWEEN HEAT FLUX AND EXPOSURE TIME AND THE EFFECTS ON HUMANS; AND THE LIMITING FLUX FOR SECONDARY FIRES

$Q kW/m^2$	t	Effect	
0.2	long	Limiting safe flux for eyes	
4	long (>60 sec)	Limiting safe flux for humans	
12.6	long	Limiting flux for secondary fires	
6.5	-20 sec	Blistering of skin	
11	-10 sec	Blistering of skin	
20	-5 sec	Blistering of skin	

Table 4.1 indicates that with long periods of exposure (> 30 seconds), the following criteria may be adopted [19]:

- Secondary building fires are possible and people, whether indoors or outdoors, are at serious risk if the flux is greater than 12.6 kW/m².
- Only people outdoors are at serious risk if the flux is greater than 4 kW/m².

The latter figure represents the flux which would cause skin blistering after 30 seconds this time being taken as representative of the time it may take to seek shelter or to escape.

For duration <30 seconds the fluxes necessary to cause skin blistering were adopted as the criteria of injury to people outdoors and secondary fires respectively [19].

In conclusion therefore all people inside the secondary fire radius are assumed to become casualties. Only those outdoors will be at risk between this radius and the radius for hazard range to people.

4.1.4 Explosion Effect

Explosion damage constitutes the effect of thermal radiation and the effect of pressure waves generated by the blast. The effects of the former were covered in the previous section. Humans are relatively resilient to overpressure generated in an explosion as shown in the Table 4.2 [20].

TABLE 4.2 EFFECT OF OVERPRESSURE ON HUMANS

Peak Overpressure		Human effects
(psi)	(kPa)	
5	34	Threshold of eardrum damage
10	69	Threshold of lung damage
40	276	Threshold of mortality

An explosion causes casualties primarily as a result of flying fragments, e.g of glass, and falling masonry as well as direct blast injuries. Damage caused by blast is shown in Table 4-3 [19, 21].

TABLE 4.3 STRUCTURAL DAMAGE CAUSED BY BLAST

Peak	Overpressure	Structural Damage
psi	<u>kPa</u>	
<1	<7	Window breakage
1-3	7-21	Walls collapse
3-5	21-34	Reinforced structures distort,
		Atmospheric storage tanks fail
5-7	34-48	Wagons and plant items overturned
>7	>48	Extensive overall damage

4.2 FIRE HAZARDS

The greatest potential hazard associated with the bulk storage of LPG is that of fire and explosion. In the process industries, fires cause more serious accidents in total than explosions and toxic releases, although the accidents in which the greatest loss of life and damage occur are generally caused by explosion [22].

Fire or combustion, is a chemical reaction in which a substance reacts with oxygen from the air and heat is released. Thus fire occurs in the gaseous state, solids and liquids either vaporise, or yield combustible vapours, which ignite and burn. There are three conditions essential for a fire:

- A fuel; gas or vapour within certain concentration limits.
- An oxidant; usually air containing a minimum concentration of oxygen.

Heat; an ignition source of minimum temperature and energy.

These three conditions are often represented as the fire triangle shown in Figure 4.2. If one of these conditions is missing fire will not generally occur (exceptions arise, e.g with oxidising agents, pyrophoric chemicals, etc).

> FIGURE 4.2 FIRE TRIANGLE

4.2.1 Flammability Limits

A flammable gas burns in air only within a limited range of compositions. Below a certain concentration of the flammable gas, the lower flammability limit, the mixture is too 'lean', whilst above a certain concentration, the upper flammability limit, it is too 'rich'. The concentration between these limits constitutes the flammable range. The lower and upper flammability limits are generally expressed in per cent by volume of vapour in air as shown for selected substances in Table 4.4 [6].

TABLE 4.4

VALVES OF LOWER AND UPPER FLAMMABILITY LIMITS FOR DIFFERENT COMPONENTS

Component	Flamability Limits		
*	Lower ($\% v/v$)	Upper (% v/v)	
Methane	5	15	
Ethane	3	12.4	
Ethylene	2.7	36	
Propane	2.1	9.5	
Propylene	2.4	11	
n-Butane	1.8	8.4	
Cyclohexane	1.3	7.8	
Hydrogen	4	75	
Ammonia	15	28	
Acetylene	2.5	100	
Carbon Monoxide	12.5	74	

All gas compositions between the lower and upper limits are flammable. The energy produced by the combustion reaction is a minimum at the limits and proceeds with increasing magnitude to the maximum energy at the stoichiometric composition at which each reactant fully reacts. The rate at which a flame propagates through a gas is dependent on several factors, such as the pressure, temperature, composition and the type of event that initiated the reaction [23].

Flammability limits, are affected by pressure, temperature, and addition of inert gas [22, 23].

Normal variations of atmospheric pressure do not have any appreciable effect on flammability limits. The effect of larger pressure changes is specific to each mixture. A decrease in pressure below atmospheric can narrow the flammable range by raising the lower flammability limit and reducing the upper limit until the two limits coincide and the mixture becomes non-flammable. This effect is illustrated for methane in Figure 4-3 [22]. Conversely, an increase in pressure above atmospheric can widen the flammable range. This effect is shown in Figure 4.4 for natural gas [22]. It may be noted that the effect is more marked on the upper than on the lower flammability limit.

Flammability limits are also affected by temperature. This effect is illustrated for methane in air in Figure 4.5 [22]. An increase in temperature tends to widen the flammable range. The effect of both pressure and temperature for methane-air mixtures is illustrated in Figure 4.6 [23].

The flammability limits are also affected by the addition of inert gas such as nitrogen, carbon dioxide or steam. This effect is shown for methane in Figure 4.7 [22].

The oxygen concentration also affects the flammability limits. In general, increasing oxygen content alters the lower flammability limit only slightly, but it's affect in raising the upper flammability is marked.

In general, carbon dioxide causes a greater narrowing of the flammable range than does nitrogen. For many flammable gas-air systems, the mixtures can be rendered non-flammable by the addition of approximately 30% of carbon dioxide or 40% of nitrogen [22].

4.2.2 Ignition Sources

When an accidental release of combustible substance has occured there is the possibility of a succeeding fire and explosion.



Figure 4.3 Effect of pressure on flammability limits of methane in air.



Figure 4.4 Effect of pressure on flammability limits of natural gas in air.



Figure 4.5 Effect of temperature on flammability limits of methane in air.



Figure 4.6 Influence of pressure & temperature on flammability of methane in air.



Figure 4.7 Effect of inert gases on flammability limits of methane in air.



There are normally a number of potential ignition sources within and in the neighbourhood of a chemical plant. Sources of ignition include the following [22]:

Flames, direct heat and hot surfaces,

Welding and cutting,

Mechanical sparks,

Vehicles: e.g, spark-ignition engines or the exhaust pipes on diesel engines,

Self heating e.g, as in lagging fires,

Static electricity e.g, discharges from hydrocarbon liquid fuels during pumping, splash-filling, rapid (air) venting.

4.2.2.1 Flames:

The flames of burners in fired heaters and furnaces, including boiler houses, may be sources of ignition on process plants. The source of ignition for the explosion at Flixborough may well have been burner flames on the hydrogen plant [22]. The flame at a flare stack may be another source of ignition.

Hot soot from chimneys of flare stacks [22] may be a source of ignition. This is a particularly important potential source of ignition on ships.

Cigarettes are another possible source of ignition. A cigarette itself is not always hot enough to ignite a flammable gas, but any match used for lighting is an effective ignition source. So smoking should be prohibited in the area of flammable hazards.

In welding operations it is necessary to ensure that no smouldering material such as oil-soaked rags have been left, e.g due to the spray of sparks, because smouldering material may act as a source of ignition. The welding torch itself is also a potent ignition source.

Machinery in distress, such as a pump with a faulty bearing, may run hot and this hot spot is a potential source of ignition. Small process fires of various kinds such as pump or flange fires, may consitute a source of ignition for a larger fire.

4.2.2.2 Welding and Cutting:

Acetylene welding, and oxy-acetylene cutting operations, are among the most common sources of ignition. Either the flame itself or the white hot shower of metal particles may reach and ignite flammables. The explosion hazard with 'empty' drums and tanks is well known; without adequate cleaning and purging such containers will retain flammable gas or vapours.

4.2.2.3 Mechanical Sparks:

Potential sources of mechanical sparks include metal tools, falling objects and cigarette lighter flints. Metal tools can, in principal, cause sparks. Evidently upon striking metal the sparks need to be above a certain minimum energy level. Non-sparking tools are available made of soft non-ferrous metals such as beryliumcopper alloy; these are used extensively in the chemical industry but are not necessarily safer if there is a chance of picking up grit. In the presence of more easily ignited gases such as hydrogen however the hazard arising from the impact of steel on steel may be appreciably reduced if non-sparking tools are used but the greater risk due to impact of metal on rock would still remain. Smears of light alloy material on steel tools, particularly if the tools are rusty, significantly increase the ignition hazard and must be avoided if such tools are used.

Falling objects like tank washing machines or cathodic protection anodes in ships cargo tanks can cause sparks which may cause more serious hazards in a space which already contains a flammable atmosphere.

Finally cigarette lighter flints can also cause a mechanical spark if the lighter is dropped.

4.2.2.4 Vehicles:

A chemical works may contain at any given time considerable numbers of vehicles. These vehicles are potential sources of ignition. The ignition of the flammable vapour cloud in the Feyzin disaster may well have been caused by a car passing on a nearby road [22].

It is necessary therefore to exclude ordinary vehicles from the area of flammable hazards. Compression ignition (diesel) vehicles are usually considered safe because they have no high tension ignition system but spark arresters need to be fitted to their exhausts.

4.2.2.5 Self Heating:

Self heating, or spontaneous combustion, of a solid material is a process of slow oxidation. Material undergoing self-heating may act as a source of ignition or may give rise to a fire or an explosion.

Self-heating can occur in oil rags left on steam pipes, oil dispersed in cotton waste, in lagging soaked with oil and coal stored in heaps. The rate of heat loss is restricted by insulation in such cases so that the temperature, and hence the reaction rate, can

escalate. Self heating requires that heat is generated in a mass of material more rapidly than it can be removed. The rate of reaction typically increases exponentially with temperature curve, H in Figure 4.8 [24]. In this Figure different rates of heat loss are presented as lines A, B, C and D. With a mass initially at temperature T_A , the temperature of the surroundings and the cooling rate following line A then clearly since the generation rate exceeds the removal rate the temperature of the mass increases. A stable situation will however occur at 'X'. If the initial temperature was T_B and the cooling rate followed line B the cooling is inadequate and a thermal runaway occurs which can lead to a fire. In this situation T_C represents the critical temperature of the surroundings.

The rate of cooling commencing at T_A might fall to follow to line D, e.g, due to insulation. Line D then represents a critical minimum rate to avoid a thermal runaway.

Mitigation of accidents due to self-heating can be achieved by good housekeeping, using appropriate lagging thicknesses and protection and by cooling hot materials before they are sent to storage, or it may be necessary to purge systems with inert gas in some cases, for instance where flammable solids of high specific surface are stored.

4.2.2.6 Static Electricity:

The hazard of static electricity usually occurs in the process industry in fluid handling operations such as pipeline flow, truck filling, in sprays and mists such as in steam cleaning and steam leaks.

Pumping of relatively pure organic materials can result in an accumulation of charge on the liquid surface in a receiving tank; if



Figure 4.8 Heat Balance Relationship H = rate of heat generation A,B,C,D = rates of heat loss.
the liquid is insufficiently conductive the charge may build up over a period, then bleed off rapidly resulting in a spark. Slow pumping, subsurface discharge, earthing and bonding or the addition of anti-static additives may be used to avoid static electric discharge [22].

The Fire Protection Association have analysed 206 fires in the process industries over the period 1963-1974. The results of the analyses are shown in Table 4.5, giving the source of ignition for the ensuing fire [25]. It is impossible to eliminate all sources of ignition for hydrocarbon-air mixtures due to their very low ignition energies. The design philosophy for hydrocarbons should therefore be to keep them entirely separated from air, or to depress the limits of flammmability by inerting.

Typical sources of ignition in LPG facilities include naked lights, incendiary sparks, hot surfaces and electrical equipment [2].

TABLE 4.5

THE SOURCE OF IGNITION FOR FIRES WHICH THE FIRE PROTECTION ASSOCIATION ANALYSED IN THE PROCESS INDUSTRIES OVER THE PERIOD 1963-1974.

Source of Ignition	No. of fires	70
Electricity	30	14.6
Hot surfaces	30	14.6
Spontaneous ignition	20	9.7
Frictional heat and sparks	15	7.3
Gas burners and flames	12	5.8
Cutting and welding	12	5.8
Static	7	3.4
Other known	17	8.2
Unknown	63	30.6
	206	

4.3 EXPLOSION HAZARDS

An Explosion involves a rapid release of energy which causes a shock wave moving away from the source [26]. This energy may originally have been stored in the system in a variety of forms; these include nuclear, chemical, electrical, mechanical or pressure energy. However, the energy release is not usually considered to be explosive unless it is sufficiently rapid and concentrated to produce a pressure wave that can be heard.

Many types of processes may lead to explosions in the atmosphere. Table 4.6 gives a list of the more significant types of explosion including theoretical models, natural explosions, intentional explosions and accidental explosions.

TABLE 4.6

TYPES OF PROCESSES WHICH MAY LEAD TO EXPLOSION

Theoretical Models	Natural	Intentional	Accidental
Ideal gas	Lighting	Nuclear	Confined vapour
			in Enclosures
Real gas	Volcanoes	High Explosives	Unconfined vapour clouds
Constant velocity Piston	Meteors	Electical spark	Unstable materials during manufacture transport, storage and use
Bursting sphere	Fires	Bomb calorimeter	Nuclear reactor Runaway

One of the most important properties which determines the behaviour of any explosion process is the energy distribution in the system and the manner in which it changes with time as the pressure wave propagates away from the source. All the energy is initially stored in the source in the form of potential energy. At the instant when the explosion starts, this potential energy is redistributed to produce kinetic and potential energy in different parts of the system.

Another view of the source of the explosion reveals that the characteristics of an explosion depends upon the material involved.

High Explosives: are considered to be substances which detonate, (produce a shock wave faster than the speed of sound in the medium) releasing energy very rapidly and creating very high pressures.(e.g TNT)

Conversely low explosives: which are considered to be substances which deflagrate or burn rather than detonate, and therefore create lower pressures (e.g hydrocarbon components).

An explosion can be produced by the detonation by a high explosive, the rapid burning of a flammable gas cloud, or a sudden failure of a pressure vessel containing high pressure gas. When used to describe the rapid burning of a flammable gas cloud, the term explosion refers to events where the flame speed is >170m/sec [19]. In order to achieve this the flame must propagate through a cloud premixed with air. The resulting over-pressures are sufficient to cause damage inside and outside the cloud as the shock wave propagates into the atmosphere beyond the cloud. This type of explosion may also be referred to as a deflagration [19].

4.3.1 Occurrence of Explosive Vapour Clouds

Spillage of a flammable material due to a loss of containment or major release does not necessarily give rise to an explosion

hazard, although it is always a potential source of fire. For a vapour cloud explosion to occur a substantial quantity of the flammable vapour must be released into the air as a cloud which must then be ignited by a source of sufficient energy.

The flammable materials that can form a potential vapour cloud are divided into the following categories [22, 26]:

- 1 Liquid below its atmospheric boiling temperature
- 2 Gas under pressure
- 3 Liquid boiling below atmospheric temperature
- 4 Liquid above atmospheric boiling temperature

In category 1, loss of containment due to the rupture of the pressure shell will cause a quantity of the liquid to flash off to form vapour. The amount flashed off depends upon such factors as the original pressure, the vapour pressure at storage temperature, and the ratio between specific and latent heats (27). This limited quantity of material may be increased by frothing and entrainment of air. Thus when liquid ammonia is lost from a vessel effective gas-liquid separation occurs only if the pressure is released slowly. If the vessel is suddenly de-pressurized gas bubbles will form throughtout the liquid and a mixture of gas and fine liquid aerosol will be ejected with explosive force. Even if the liquid contributes little to the explosive potential of the mixture the entrained fraction is available to feed subsequent fires and explosions and to vaporise on any warm plant in the vicinity of the leak and so add to the size of the cloud [28].

In the case of a gas emission (category 2), the amount released

is usually limited because although gases are normally stored under high pressure their density is relatively low. Although large-scale, low pressure vessels exist, they rarely fail catastroph-ically. Failures of a non-catastrophic nature (usually referred to as 'leakbefore-break') [21] occur more frequently but, because of the low pressure, do no generate large volumes of explosive mixtures.

Category 3, includes cryogenic liquids (e.g. LNG) which are intrinsically more dangerous if stored under high pressure than at low temperature. Refrigerated storage may pose other problems due to steel embrittlement at low temperatures.

In this case, loss of containment will not result in such rapid flashing of liquid and a vapour cloud can only be formed by evaporation from the cold liquid.

In category 4, liquids are the most hazardous. They contain pressure energy for atomisation and thermal energy to sustain evaporation. Furthermore, their density is likely to be more than that of category 3 flammable LNG and LPG, so the mass released is likely to be more than with category 3.

4.3.2 Vapour Cloud Dispersion

It is frequently asserted than any flammable vapour cloud which disperses near ground level within a process plant will find an ignition source. However, there are several documented incidents in which clouds have dispersed without ignition [29].

The main factors that determine the probability of ignition include the nature of the material, the size of the cloud, and location of the source. Small leaks have a high probability of harmless dispersion because concentration falls to well below the lower

flammable limit before a source of igntion is encountered. For example leakage from a gasoline can during normal filling extends only of the order of 2 m downwind. This may also occur with large leaks in remote locations, such as storage areas. For large clouds in process areas, ignition is much more likely, and generally fairly immediate. Kletz [30] estimated that for small leaks about one in 30 ignites. It has also been suggested [30] that for releases greater than 10 tonnes of flammable vapour the probability of ignition can be as high as one in two.

The movement of the vapour cloud and the extent of its flammable region, depend upon the mixing process during the dispersion of the fuel. In general, by simple diffusional mixing, the pure fuel at the spill will be diluted to below the upper flammability limit somewhere in the inner region of the cloud and to below the lower flammability limit somewhere in the outer region of the cloud.

A variety of factors affects dispersion. Models should include the effect of the vapour density but common atmospheric dispersion models, like the Gaussian model [31] are only applicable to gases with a density approximately equal to that of air. Many gases have considerably higher or lower densities and a different dispersion behaviour may be expected, especially when large amounts are released. The direction and velocity of discharge, as well as the nature and size of spill, are other important factors affecting the dispersion model. There are two general classes of spill to consider: the massive single spill produced by the catastrophic or rapid rupture of a storage tank, and the continuous leak produced by either a relatively small hole in a tank or a break in a pipeline. Dispersion models for each type have been derived and may be applied to calculate the location of the upper and lower flammability limits for

flammable materials. For a massive single spill the severity of the danger rises at first and then decreases as more and more of the spill is diluted to below the lower flammability limits of the fuel. In the case of a continuous spill, as long as the spill is occuring, the danger increases to an asymptotic limit as the delay to ignition increases, because the extent of the flammable region must eventually be stabilised by dispersion in the atmosphere [32].

The dispersion of an unconfined vapour cloud is dependent upon a variety of factors including direction and velocity of discharge, air temperature and wind conditions on site, effects of buildings and plant, vapour and ground temperatures, and density of the vapour [33]. Normal air pollution dispersion models have been used to predict dispersion of such vapour clouds, but for heavier than air vapours the clouds are likely to be more stable than predicted from Pasquill theories [34]. Experimentation in wind tunnels [35] showed that a rapid release of LPG vapour tended to produce plumes which were:

- Initially influenced by gravitational effects (slumping) giving rise to a rapid spread upwind as well as downwind of release,
- Greatly influenced by slope effects, i.e. topography, and would travel considerable distances downhill from the point of release,
- Relatively stable, i.e. turbulence within the plume due to ground roughness was suppressed, and
- Considerably affected by wind speed, i.e. high wind speeds greatly reduced the stability.

4.3.3 Ignition and Combustion

4.3.3.1 Ignition

A vapour cloud will continue to disperse into the atmosphere unless and until it is ignited upon reaching a suitable ignition source. From reported incidents only about 1 cloud in 6 has drifted more than 100 m before ignition occurred [29].

Immediate ignition of an instantaneous pressurised burst always gives rise to a fireball. An immediate ignition of a continuous pressurised release may be assumed to give rise to a torch. Immediate ignition of a refrigerated spill will give rise to a pool fire.

With both a pressurised release and a refrigerated spill a delay in ignition will give rise to an explosion [19].

Delays as long as 15 minutes have been reported with a large leak and proper wind conditions [29]; an enormous combustible volume can be formed in this time. The consequences of releases of pressurised and refrigerated liquefied flammable gases are shown in Table 4.7 [21].

TABLE 4.7

THE CONSEQUENCES OF RELEASES OF PRESSURISED AND REFRIGERATED LIQUEFIED FLAMMABLE GASES

Time of ignition	Refrigerated release	Pressurised release
Immediate (within seconds)	Pool fire	Fireball
Slightly delay (within 30 sec)	Diffusion flash fire	Premixed flash fire/explosion
Delayed (minutes)	Premixed flash fire/ explosion	Premixed flash fire/ explosion

4.3.3.2 Combustion

The way in which LPG releases burn is dependent upon a number of factors including [19]:

- the conditions under which the material is released, e.g. temperature, pressure, rate of release, quality of release,
- the nature of the flammable cloud at the point of ignition, e.g.
 its size, shape, composition, degree of confinement,
- the nature of the ignition source.

The modes of combustion can be described in terms of whether the flames are [19]:

- premixed or diffusion, and
- stationary or propagating.

Pool fires and torches are examples of stationary diffusion flames. Both represent combustion of the release at source. A necessary prerequisite for a pool fire is that the release gives rise to the formation of a liquid pool. Pool fires are, therefore, more likely for the release of liquids stored below their boiling point or for pressurised releases with a low flash fraction. Torches describe the combustion of gas or liquid sprays from pipework or a small hole. They will be most likely with pressurised releases.

Fireballs and diffusion flash fires are examples of propagating diffusion flames. They occur following the ignition of a cloud the bulk of which is above the UFL. The edges of the cloud, however, will be within the flammable range due to diffusion and mixing with the surrounding air. Ignition is possible at such points and may be followed by rapid flame propagation through the flammable region so

that the central rich core of the cloud is enveloped in flame. Subsequent burning is controlled by the rate of entrainment of air and its mixing with fuel and will be a relatively slow process. The cloud burns as a fireball when the whole of the cloud will appear to be on fire, this taking place at eddy boundaries where the fuel-air composition is in the flammable range. Fireballs are more likely following ignition of a hemispherical or spherical cloud such as will exist shortly after a pressurised release. Diffusion flash fires are more likely following ignition of clouds with heights much less than their maximum width or length, e.g. those formed by slumped hemispherical clouds, by evaporation from liquid pools or plumes from continuous releases [19].

Propagating premixed flames occur when a cloud of gas mixture inside the flammable range is ignited. The flame propagates outwards from the ignition source consuming the mixture. For very low propagation speeds the rate at which hot combustion products are generated is low and so expansion can take place easily without any significant overpressure being generated. Under these circumstances the cloud will burn as a flash fire.

As flame speeds increase combustion products are generated more quickly so that pressure is built up both ahead and behind the flame front. At flame speeds >170-200 m/sec the overpressures are sufficient to cause extensive damage both within the cloud and outside it as the pressure waves propagate and decay in the atmosphere beyond the burning cloud. The event that produces damaging overpressure is referred to as an explosion [19].

4.3.4 Types of Explosion

4.3.4.1 Confined Vapour Cloud Explosion (CVCE)

Confined vapour cloud explosions occur when fuel-air mixtures are ignited in buildings or vessels and confinement allows the pressure to build up until the containing walls rupture due to the rise in the internal pressure [22, 36]. Whether or not this rise is sufficient to cause rupture depends upon many parameters such as:

- the design of the enclosure,
- the amount of hydrocarbon present,
- the degree of mixing with air,
- the extent and speed at which the enclosure or building vents the combustion products and so lowers the effective internal pressure.

In the case of a pressure vessel, where the bursting pressure should be known, the energy resulting from ignition of the mixture would be given by the isentropic expansion energy

$$\Delta E = \frac{P_0 V_0}{\gamma - 1} \left[1 - \frac{P_0}{P_f} \right]^{1 - \gamma} \dots 4.1$$

where ΔE = Isentropic expansion energy, J

 P_o , P_f = Initial and final absolute pressure, N/m²

 $V_0 = Original volume, m^3$

 γ = ratio of specific heats

This is the same equation as for the bursting of a pressure vessel and can be calculated in the same manner.

Normally Pf is unknown but is low in the case of confinement within ordinary buildings or structures. Following ignition of a massive release of materials, CVCE's will add substantially to the local severity within a UVCE and probably result in much of the major damage found within the confines of the cloud. However, confined vapour cloud explosions are generally insufficiently large to do more than local damage to the plant. This type of explosion is common where LPG cylinders develop a slow undetected leak which results in internal concentrations of air-vapour mixtures between the UFL and LFL. When an ignition source is suddenly introduced into the building, a violent explosion can result. There is probably a good case for always isolating portable LPG appliances at the cylinder outlet valve when not in use; this would greatly reduce the risk of confined vapour explosions. CVCE's will normally also be a hazard in situations where unconfined vapour cloud explosions are possible. The spacing policy relating to this type of explosion is related to the production of missiles from building or plant.

4.3.4.2 Unconfined Vapour Cloud Explosion (UVCE)

An unconfined vapour cloud explosion takes place when a large fuel-air cloud burns in a relatively unconfined area with sufficient speed to generate pressure waves which propagate through both the vapour cloud and the surrounding atmosphere [26, 37]. The pressure waves generated can be thought of as the physical mechanism by which chemical energy released during combustion of fuel-air cloud is transported to distant points where it may cause damage through the action of incident pressure waves on structures. Events of this type usually involve large releases of 5 tonnes or more

of common hydrocarbons [19]. However, Briscoe [37] suggests that this should be over 10 tonnes.

Explosions are caused by local ignition (ie at a single point) followed by flame propagation throughout the rest of the vapour cloud. Hence the rate of combustion is directly related to the speed with which the flame propagates.

Two forms of propagation have been distinguished, deflagration when the flame advances at subsonic speed, and detonation when the flame advances at supersonic speed.

Flames in a deflagration may travel at a range of speeds; higher flame speeds produce higher pressure levels and gas velocities. Generally pressure levels are relatively low, ie less than 20 m/s flame speeds give negligible pressure levels and greater than 250 m/s give a level of 0.1 bar [37]. In the event of detonation, flame speeds of 2000 m/sec could produce pressure levels of approximately 16 bars, but these are unlikely in an UVCE. When the shock waves reach the end of the vapour cloud they begin to decay since they are no longer sustained by chemical energy. It is this decaying wave which causes most of the damage outside the factory site.

4.3.4.3 Boiling Liquid Expanding Vapour Explosion (BLEVE)

All liquefied gases are stored in containers at temperatures above their normal boiling points and remain under pressure only so long as the container remains closed to the atmosphere. This pressure ranges from less than 0.06 bar (1 psi) for some liquid natural gas containers to 15 bar (200 psi) for liquefied gas containers at normal storage temperatures. If the pressure is suddenly reduced

to atmospheric, as would be the case if the container developed a crack, the sensible heat which is stored in the liquid at the high pressure causes the liquid to evaporate very rapidly with consequent rapid cooling of the liquid. Failure of the container is most often due to the weakening of the container due to flame impingement and consequently the metal temperature increasing. Most steels used for pressure vessels storing or transporting LPG will steadily lose tensile strength at temperatures above 400 °F. Thus, container steels exposed to an external fire above the liquid level within the container, above the vapour space, will increase in temperature shell below the liquid level will remain very near to the boiling point of the pressurised stored liquid so long as the internal pressure does not decrease. The liquid to gas expansion ratio may be from 200:1 up to 600:1.

The expansion of the liquid into vapour due to a sudden decrease in internal container pressure is the very process that tends to increase the length of the small, initial crack at a rate close to the velocity of sound in metal.

There are three separate damage producing effects [22, 38]:

A blast wave due to relief of internal pressure,

- Thermal radiation due to the effects of the fireball resulting from massive burning of the contents of the vessel in the air,

Projection of large fragments, scattered for considerable distances, by the violent rupture of the tank.

The main effect arises from thermal radiation and people at distances of 300 metres or more have been cited as having been injured. The secondary, and more specific, hazard due to this type

of occurrence is that of projectiles, extending to distances of 300 metres for large fragments and 500-600 metres for smaller fragments.

CHAPTER FIVE

BLEVE'S INVOLVING PRESSURISED STORAGE TANKS

5.1 LPG TANK FAILURE MECHANISMS

Most of the LPG storage tanks in service have been designed in accordance with the requirements stated in Codes of Practice [39]. Some Codes state that bulk pressure vessels must be designed and constructed to a recognised standard and give examples such as ASME, BS 1515, BS 5500, etc., and that they must be designed for the highest pressure to which they will be subjected in service. Normally these LPG tanks are designed for a burst pressure of 1000 to 1250 psig. Safety pressure relief valves are installed to prevent the internal pressure from rising more than about 10 percent above the maximum design pressure. The flow capacity of the valves is designed to maintain safe tank internal pressure levels with the tank and its contents exposed to excessive heat gains, such as from a fire. It is generally accepted that the maximum allowable heat gain is about 1.43 x 10^9 kJ.sec/m² (35,000 Btu.h/ft²) from a spill fire [39].

It is assumed that adequate welding procedures and inspections have been carried out during the fabrication of an LPG storage tank, in accordance with the appropriate pressure vessel Code, depending on whether or not the storage is refrigerated, e.g. BS 5500. Therefore, the most probable source of tank failure is explosive failure due to excessive heat gain by the tank during exposure to severe fire conditions. As is well documented by others [39], sudden release of the stored energy within a high temperature and high pressure LPG tank can produce hazards ranging from sudden total explosive

disintegration of the tank, with resulting high velocity large metal sections, to projectile like motion of a large section of the tank.

An examination of the fire record relating to tank fires shows that four types of fire condition frequently occur. These types are as follows [39]:

Pressure relief valve vapour fires: This type of fire condition is caused by moderately high pressure vapours discharging through spring-loaded pressure relief valves which have to be located on all LPG storage tanks of an industrial size (30,000 gallon capacity or more). Ignition is usually produced by static discharge. The resultant fire can be extremely dangerous due to the high rate of heat input into the tank vapour space. Extinguishing such pressure relief flames is recommended to minimize such heat input conditions, providing an explosive situation will not be produced by the release of the LPG vapours through the relief valves.

Radiant heating effects from a remote fire: An LPG tank will be partially exposed to incident heat fluxes from fires at other locations such as adjacent tankage, spills at unloading stations, or other types of flammable hydrocarbon fires (e.g., liquid spills). If such fires are not controlled or extinguished, an unprotected LPG tank will eventually develop a pressure relief type valve fire as described above.

Direct contact heat transfer from spill fire flames: Many LPG tanks and tank farms utilize local impounding dikes to prevent the spread of flammable product and to limit the extent of flammable vapour dispersion. In many cases if a fire occurs this will result in direct

flame contact on the bottom and lower sides of the LPG tank and radiation on to the upper surfaces.

High pressure impingement fire condition: This is the most dangerous type of fire due to the greatly increased heat fluxes, especially if the flame impingement is on the vapour space or on an empty LPG tank which does not have the heat capacity protection provided by the liquid contents of a full LPG tank. Such fires are experienced from the failure of loading lines, for instance, when a tank car turns over and both liquid and vapour are released through the relief valve.

The failure of an LPG tank exposed to a fire condition is related to its steel strength characteristics as a function of temperature. The strength of LPG tank steels increases initially as the temperature increases, up to a maximum somewhere between 350 and 450 °C beyond which the strength starts to decrease. At a temperature of about 540 °C the strength of a typical LPG tank will be reduced from a design pressure of about 7000 kPa to some 2000 kPa, and further heating to 600 °C will lower the strength still further to approximately 1300 kPa [39].

In heating LPG tanks by exposure to fire, some of the absorbed heat flux goes into raising the stored liquid temperature and some goes into raising the steel temperature. The heat absorbed by the liquid causes liquid to vaporise, with a resultant increase in tank vapour pressure. When this pressure reaches the relief setting of the safety relief valves, bursting discs, etc., this excess pressure is allowed to vent to the atmosphere. During severe fire conditions, such as in direct exposure to spill fires or impinging pressure fires, the heat input is sufficient to cause a continual increase in the tank internal

pressure. Meanwhile the heat input into the tank steel not in contact with liquid product, causes an increase in the steel temperature. In time a point is reached where the steel temperature in a localised area is reduced to the local bursting strength below that being maintained by the tank pressure relief valve system. At that time the vessel will rupture and a boiling-liquid expanding vapour explosion (BLEVE) will result. A typical sequence is illustrated in Figure 5.1 [38].

5.2 DAMAGE PRODUCING EFFECTS OF BLEVE'S

There are three separate damage producing effects [22, 38]:

- i A blast wave due to relief of internal pressure,
- ii Thermal radiation due to the effects of the fireball resulting from the massive burning of the contents of the vessel in the air, which will be discussed in Chapter 6,
- iii projection of large fragments, scattered for considerable distances by the violent rupture of the tank.

5.2.1 Blast wave effect

Damage from failure of a vessel under pressure is partly due to the shock wave produced in the surrounding air by the sudden release of energy.

The work of Strehlow and Ricker [40] showed a simple relationship between Ps, the dimensionless shock overpressure and Rs the dimensionless shock radius for all sphere bursts. Their work presents a graphical method of predicting the shock overpressure at any radial distance from a spherical vessel which has burst. All pressures are reduced to dimensionless overpressures by means of the ambient atmospheric pressure Po. Thus the dimensionless over-



Figure 5.1 The development of a BLEVE

pressure in the sphere which is initially at absolute pressure P_2 is given by

$$\overline{P}_2 = \frac{P_2 \cdot P_0}{P_0} \qquad \dots 5.1$$

similarly the shock overpressure

$$\overline{P_{s}} = \frac{P_{s} - P_{o}}{P_{o}} \qquad \dots 5.2$$

The radial distances are made dimensionless by dividing by the radius R_0 to which the gas in the sphere would have expanded adiabatically, if reduced from P_2 to P_0 . The scaled radial distance of the shock front is given as,

$$\overline{R}_{S} = \frac{R_{S}}{R_{O}} \qquad \dots 5.3$$

For a gas with a specific heat ratio of γ_1 initially stored at P₂ in a sphere of radius r₀.

$$R_0 = \left[\frac{E_1}{P_0}\right]^{1/3}$$
 5.4

Applying Brode's formula [40] which has been suggested for calculating the effective energy stored in a pressurised sphere of ideal gas, as shown in equation 5.5

$$E_{1} = \begin{bmatrix} P_{2} - P_{0} \\ \hline \\ (\gamma_{1} - 1) \end{bmatrix} V_{0} \qquad \dots 5.5$$

with $V_0 = \begin{bmatrix} 4\pi \\ 3 \end{bmatrix} r_0^3$, then equation 5.5 becomes

 $E_1 = \begin{bmatrix} 4\pi \\ -3 \end{bmatrix} \begin{bmatrix} P_2 - P_0 \\ -\gamma_1 - 1 \end{bmatrix} r_0^3 \qquad \dots 5.6$

then substituting equation 5.6 in equation 5.4 gives:

$$R_{o} = \left[\frac{4\pi (P_{2} - P_{o})}{3(\gamma_{1} - 1) P_{o}}\right]^{\frac{1}{3}} r_{o} \qquad \dots 5.7$$

At the instant of failure the scaled shock front radius \overline{R}_{so} is given by equation 5.8

$$\overline{R}_{so} = \frac{r_o}{R_o} = \left[\frac{3(\gamma_1 - 1)}{4\pi P_2}\right]^{\frac{1}{3}} \dots 5.8$$

The contact surface pressure for the particular sphere gas/ambient air conditions, determines the initial reduced shock front overpressure \overline{P}_{so} at \overline{R}_{so} . This value may be found from equation 5.9 by an iterative technique [40].

$$1 + \overline{P}_{so} = (1 + \overline{P})_{2} \begin{bmatrix} 1 - \frac{(\gamma_{1} - 1) \overline{P}_{so} (a_{o} / a_{2})}{(2\gamma_{o} [2\gamma_{o} + (\gamma_{o} + 1) \overline{P}_{so}])^{\frac{1}{2}}} \end{bmatrix}^{\frac{2\gamma_{1}}{\gamma_{1} - 1}} \dots 5.9$$

where

 γ_1 , γ_0 = the specific heat ratio of the stored gas and air respectively

 a_0 , a_2 = the sonic velocity in air and in the sphere, respectively, at the instant of rupture

The initial values of $\overline{R}_{s} = \overline{R}_{so}$ and $\overline{P}_{s} = \overline{P}_{so}$ are plotted at the point 'x' in Figure 5.2, to determine the appropriate curve which predicts \overline{P}_{s} for \overline{R}_{s} beyond the sphere radius ro. Note from equations 5.8 and 5.9 that the initial point on Figure 5.2 [41] is independent of the size of the spherical vessel, but from equation 5.7, it is clear that the range Rs, which is obtained from equation 5.3 is directly related to the vessel radius r_o.

The fact that all the overpressure versus scaled distance curves shown in Figure 5.2 are roughly parallel allows construction of a graphical technique for determining the actual overpressure-distance relationship for any specific spherical vessel burst. Thus the same method is used to find the overpressure distribution for different storage pressures as shown in Figure 5.3, and for different sphere diameters as shown in Figure 5.4.





Example: Catastrophic Failure of Propane Tank Mass of propane = 30 tonnes Storage pressure = 20 bar gauge Ambient temperature = 300 K Diameter of storage sphere = 5 m

Calculation of blast damage from vessel rupture can be carried out by making an assumption such as, taking atmospheric pressure as 1 bar approximately, the reduced overpressure in the vessel initially is equal to the gauge pressure (ie $\overline{P}_2 = 20$) and the specific heat ratios for each of LPG and air are 1.2, 1.4 respectively.

So from equation 5.8

$$\overline{R}_{SO} = (3 \text{ x} - \frac{0.2}{80\pi})^{1/3} = 0.134$$

$$\log R_{so} = -0.874$$

since

$$\frac{a_0}{a_2} = \left(\frac{\gamma_0}{\gamma_1}\right)^{1/2}$$

then

$$\frac{a_0}{a_2} = \left(\frac{1.4}{1.2}\right)^{1/2} = 1.08$$

Substituting the values of \overline{P}_2 , a_0/a_2 , γ_1 and γ_0 in equation 5.9 gives,



Figure 5.3 Overpressure vs Radial Distance following rupture of a 2.5 m Spherical Storage Diameter, assuming Different Storage Pressures.



Figure 5.4 Overpressure vs. Radial Distance following rupture of Spheres at 20 Bar Storage Pressure, assuming Different diameters.

$$1 + \overline{P}_{so} = 21 \left[1 - \frac{0.2 \times 1.08 \overline{P}_{so}}{\left\{ 2.8 (2.8 + 2.4 \overline{P}_{so}) \right\}^{\frac{1}{2}}} \right]^{12}$$

Solving this equation gives $\overline{P}_{so} = 3.1$ log $\overline{P}_{so} = 0.49$

The conditions at the moment of rupture are located at the point 'x' on Figure 5.2, the overpressure distribution is found as follows from the broken line on Figure 5.2.

From equation 5.8

$$R_0 = \frac{r_0}{\overline{R}_{SO}} = \frac{2.5}{0.134} = 18.7 \text{ m}$$

From equation 5.3

 $R_{s} = R_{o} \overline{R}_{s} = 18.7 \overline{R}_{s}$

The radii quoted in Table 5.1 were obtained knowing the value of \overline{R}_{s} which corresponds to a known value of overpressure from Figure 5.2. A computer program was written to enable rapid calculations using this method; this is given in Appendix A and the result of typical calculations are given in Table 5.1. Typical damage at each radius can be estimated from Table 5.2 [42].

5.2.2 Projectiles

The sudden release of energy during an explosion may project

from the epicentre a wide variety of missiles. In the case of BLEVE type incidents, the projectiles can range in size from small splinters of steel, to large storage tank sections of up to three quarters of the tank in size. Incidents are on record of such large fragments being projected for up to a mile and the danger these pose is apparent [3].

TABLE 5.1 OVERPRESSURE vs RADIAL DISTANCE FOLLOWING CATASTROPHIC FAILURE OF PROPANE TANK AT 20 BAR GAUGE STORAGE PRESSURE

Overpressure	Radial Distance
(kN/m^2)	(m)
309	2.5
281.5	2.95
189.2	4.69
111.68	7.7
69.1	11.79
36.8	18.79
22.4	30.32
17.13	36.3

The consequences from this type of event are very hard to predict because the trajectories of any fragments depend upon such variables as shape, size and angle of projection as well as the effect of wind. The wind direction (and its likely effect) in any particular incident is largely unpredictable and its effect on missiles is likely to be small.

Upon rupture tanks containing LPG usually fragment into a number of parts, eg 10 to 20. With cylindrical tanks the number is considerably less. Energy released upon rupture of LPG tanks is partly converted into kinetic energy of the fragments. As a result, the fragments may be scattered great distances. Figure 5.5 [16] shows the

TABLE 5.2

DAMAGE CAUSED BY VARIOUS AMPLITUDES OF SHOCK WAVES

PRESSURE kN/m ²	DAMAGE
0.2	Occasional breaking of large glass windows already under strain.
0.7	Breakage of windows, small, under strain.
1.0	Typical pressure of glass failure.
2.0	"Safe Distance" (probability 0.05 of no serious damage beyond
2.0	Missile limit
2.0	Some damage to house ceilings 10% window glass broken
2.8	Limited minor structural damage.
3.5-6.9	Large and small windows usually shattered; occasional damage to window frames
4.8	Minor damage to house structure
5.2	Breakage of small windows, not under strain
6.9	Partial demolition of houses, made uninhabitable.
6.9-13.8	Corrugated asbestos shattered. Corrugated steel or aluminium
	panels, fastenings fail, followed by buckling wood panels
0.0	(standard housing) fastenings fail, panels blown in.
9.0	Steel frame of clad building slightly distorted.
13.8 20.7	Partial collapse of walls and roofs of houses.
15.0-20.7	Concrete or cinder block walls, not reinforced, shattered.
17.3	Lower limit of serious structural damage.
20.7	Heavy machines (ut 1350 kg) in industrial hail it is
20.7	damage
20.7	Frameless self-framing steel papel building demolished.
	frame building distorted and pulled away from foundations
20.7-27.6	Frameless, self-framing steel panel building demolished Rupture
	of oil storage tanks.
27.6	Cladding of light industrial building ruptured.
34.5	Wooden utilities poles (telegraph etc) snapped. Tall hydraulic
	press (18000 kg) in building slightly damaged.
34.5-48.3	Nearly complete destruction of houses.
48.3	Loaded train wagons overturned.
48.3-55.2	Brick panels 0.2-0.3 m not reinforced, failed by shearing or flexure.
62.1	Loaded train/box-cars completely demolished.
69.0	Probable total destruction buildings. Heavy (3200 kg) machine
	tools moved and badly damaged. Very heavy (5500 kg) machine
	tools survived.
2000	Limit of crater lip.



position of the larger fragments of LPG tank in the San Juan Ixhuatepec incident up to distances of about 400 m. At least four large fragments came down at distances of 600, 750, 1100 and 1200 m [16]. It has been considered [3] that a safe evacuation radius from pressurised storage tanks of LPG should be a least 600 m.

There is, however, one area of consideration where the geometry of the projectiles, together with weight, is accurately known, that of gas bottles in storage at LPG facilities. The probability of a gas bottle striking LPG storage tanks and pipes, leading to large scale spills, is extensively covered in the Canvey Report [11]. The conclusion reached was that a 14.5 kg bottle with an initial velocity of 100 m/s will upon impact:

- i bounce off storage spheres,
- ii possibly penetrate cylindrical tanks (bullets),
- iii certainly rupture piping.

Gas bottles are present at filling facilities in quite large numbers, and in the past a number of fires at these installations have resulted in projectile damage at distances up to some hundreds of metres. As a result of these experiences, there has been a move to fit relief valves to bottles to relieve any excessive pressure build-up in a fire situation. This measure substantially reduces the chance of failure of the bottle and subsequent BLEVE. However, the plume of burning gas from a relieving bottle will extend for some distance, endangering fire-fighters and spreading the fire. Furthermore, if the vapour vented from a fire exposed cylinder is not ignited immediately, due perhaps to a local oxygen deficiency, then delayed ignition might well lead to a semi-confined explosion with missiles ejected a considerable distance. To prevent this occurrence, bottles should not be stacked on pallets more than five high.

Finally, injury due to missiles has not been evident, in major incidents and, therefore, historically, missile production would appear to be a low risk primary hazard. (This effect is, of course a function of the distance of personnel from the epicentre of the BLEVE, so that in 'minor' incidents involving small cylinders being fitted/adjusted the incidence of injury from fragments would be expected to be greater.) These projectiles might, however, give rise to a secondary hazard on site due to rupture of storage tanks etc. Theoretically this low risk would appear to be a product of the small size of people in relation to the area in which the missiles fall and the fact that in urban environments the majority would be under cover giving an unquantifiable level of protection from a missile.

5.3 PROTECTION OF LPG TANKS

LPG tanks exposed to severe fire conditions suffer an increase in internal pressure while simultaneously undergoing a decrease in tank burst strength. Although the required safety valves can hold the internal pressure level to about 20% to 25% of the design burst strength, the heat absorption from exposure to a fire will rapidly lower the steel burst strength to a point where the steel is not capable of withstanding even this safe relief pressure and explosion results. Tests have been conducted in the USA [39] using a standard 30,000 gallon LPG blimp with a standard paint finish and filled with LPG which was exposed to direct flame contact from an impounded JP-4 fuel spill fire beneath the blimp. The tank vented after about 12 minutes of fire exposure and exploded after about 25 minutes, 60% of the contents still being liquid. This points to the need to keep the tem-



Figure 5.6 Methods of protecting a pressure vessel against fire (Support legs are also normally protected by insulation and water sprays) perature of LPG storage tanks low, especially the parts of the tank that are not wetted (and thus cooled) by the internal contents.

There are several methods currently in use for thermal protection as shown in Figure 5.6 [43] and these will be discussed in turn.

5.3.1 Sloping the ground

The base on which the pressure vessel stands should be impervious and sloped so that any spillage is collected. A slope of 1 in 40 is recommended [44]. Spillage of flammable liquid cannot then accumulate under the vessels, as it did at Feyzin [43]. The collecting area should be such a distance away from the tanks, and of such a surface area, that the flame from a fire in the collecting pit will not impinge on the tank (considering that flames may be at angle of 45° to the vertical and that the flame length may be twice the pool diameter).

In the case of process vessels the principle of sloping should be applied so that the gradient is not less than 1 in 60. It should lead to a safe controlled drainage or collecting point. If the ground is level, or worse, slopes towards the centre of the plant, spillage will remain in the plant area. Also, hydrocarbons will float on top of the water used for cooling the vessels and could be a hazard to other plants nearby.

5.3.2 Water Cooling

Water application by either direct contact water spray fixed piped system, monitor nozzles, and hose lines are classic methods for cooling LPG tanks exposed to fire conditions. The theory is that, providing the water film is maintained over the tank, then the temperature cannot exceed the boiling point of water, and tank shell strength remains adequate. However, if the exposure to fire is very

strength remains adequate. However, if the exposure to fire is very severe, the water may well be vaporised before contacting the vessel, or high velocity vapour and flame effects may blow aside the water film and permit direct flame impingement on the steel shell. There are recorded data which substantiate the inadequacy of water systems for LPG tank protection; their complexity makes them unreliable and very costly, both from an initial installation and from a maintenance viewpoint. Further, there is a lack of evidence to show the effectiveness of fixed water spray systems for the prevention of LPG tank explosions during tank exposure to flame contact from spill fires and impinging vapour/liquid pressure fires.

The rate of water application usually recommended where vessels are liable to be exposed to direct flame impingement is 10 litres/m²/min [44]. Such a rate is hardly practicable to supply if a number of large vessels have to be cooled. For example, 450 m³/hr are required for a 2000 m³ sphere. This reinforces the case for sloping the ground. Where radiant heat alone has to be considered, much lower rates may be sufficient.

Water from fixed equipment may be applied to the top of the vessel in cases where it will be evenly distributed over the whole surface. In other cases, it should be applied directly on to all parts of the vessel as shown in Figure 5.7 [45]. The former is much cheaper but care must be taken to see that obstructions on the vessel do not impede the flow.

Fixed equipment should preferably be permanently drained, periodically tested and serviced to ensure that it does not freeze or rust.

Finally, care must be taken that there is adequate water in the


Figure 5.7 Spray System for Spherical Tank.

fire system or all times to operate the fixed water drenching installation. The quantity of water applied by a fixed system must leave an adequate balance in the total water supply system for use by mobile equipment. Provision must be made for controlled drainage of the applied water. It is not economically practicable in some cases to design and construct underground drains capable of handling the required flowrate.

5.3.3 Insulation

Insulation, particularly of vessels containing high risk material such as liquefied flammable gases, is recommended in some codes such as the ICI LFG Code but not in others [43]. Tests and fire experience have shown that insulation should withstand the jets from fire hoses.

Insulation is on the vessel as an immediate barrier to heat input before the water drench is applied. The advantage of it is when the water supply lines have been damaged by fire or any associated explosion. Nevertheless, insulation should be looked on as additional to water drenching, not as an alternative. Insulation slows down the rate of heat input to the vessel, typically by a factor of about ten [43], but it does not ultimately prevent the vessel getting too hot. With insulation there is more time to commission water drenching and augment fixed systems with portable monitor sprays. Figures 5.8 (a and b) show the application of insulation [45]. Methods which have been suggested as a means of providing fire insulation may be summarised as follows.







(a) Structural Fire Protection, Elevation

(b) Concrete Fire Protection.

5.3.3.1 Mounding or burying of the LPG Tank:

Below ground burial is the easiest method of thermal protection to understand and accomplish, but it is the least desirable method due to the high potential for corrosion and safety problems associated with undetected LPG leaks. The seriousness of underground leakage of LPG should not be underestimated; Andrews [3] reports on a test to determine the problems attendant upon the sealing of a leak in a 8" diameter LPG pipeline buried 4ft below ground in sandy soil. The pipeline was deliberately drilled with a 3/32 inch hole prior to burial, and upon pressurising the pipeline with LPG, the escaping vapour was ignited as it issued through the soil. After some hours a mechanical excavator was moved into position, protected by water sprayed from hoses, and proceeded to uncover the pipeline. However, at a depth of two feet the excavator met with rock-hard ground, which was in fact frozen sand and water and LPG due to the rapid evaporation at the point of leakage and a consequent drop in temperature. Two water jets were needed, over a considerable time, to soften the sand so that it could be removed. Even after removal it continued to burn. This is indicative of the type of difficulty that might well face the emergency services in the event of an accident with a buried LPG tank.

Above ground burial, or mounding, offers protection from flame impingement, but has inherent with it the problems of tank and piping system redesign or requalification if the tank to be mounded was initially designed for above ground service. All piping must usually be by way of top-side manhole connections. It has been recommended [43] that burying a tank never be used.

5.3.3.2 Standard Insulation Systems

Conventional or standard insulation systems such as cork, glass-wool or aggregates such as vermiculite, perlite or calcite require an outer metallic sheath for environmental protection and to provide mechanical attachment of the insulation. The lack of firm bonding between the insulation and the LPG tank shell permits moisture entrapment and subsequent corrosion. During severe flame impingement fires, the relatively thin metallic sheathing deforms with loss of the previously protected thermal insulation.

Tests have shown that vermiculite-concrete exposed to a fire, stands up well to fire and water jets without damage [43].

Low temperature vessels should be provided with fire protection on the outside of the heat insulation because the insulating materials available for low temperature duty do not have good fire resistance, for example, cork or foam glass covered with a vermiculite concrete layer 10-20 mm thick. One disadvantage with an insulating cover is that it is difficult to inspect the vessel or supports for corrosion. On one occasion a vessel collapsed due to corrosion of the supports. Fortunately this occured when it was filled with water for test. Insulation of a vessel should be designed so that sections can be removed for inspection of the vessel and supports.

5.3.3.3 Refractory protection systems

Refractory materials are normally used to give thermal protection to the inside of furnaces, kilns and high temperature process line [39]. When applied to the fire protection of steels where a hydrocarbon spill fire can be expected to reach maximum intensity within a few seconds, such materials will crack under the condition of

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large thermal stresses, leaving large areas of the tank shell exposed to direct flame. Basically most refractory materials are designed to be brought very slowly up to their operating temperature, as well as being cooled down quite slowly.

5.3.3.4 Cement Compounds

These compounds including ordinary concrete and gunite provide good fire protection insulation systems for steel structures [39]. They are, however, quite heavy, expensive, in most cases corrosive, and exhibit poor mechanical bonding properties.

5.3.3.5 Ablative Coating

These types of material provide good protection for structural steels. The fundamental principle is to apply a coating that gradually erodes due to the heat input from the impinging fire. The solid material changes into a gas, utilising heat energy that would otherwise be used for heating of the unprotected LPG tank. The temperature rise in the protected vessel is in direct proportion to the thickness of the coating and its thermal properties. Some ablative coatings result in a simultaneous generation of a tough microporous char layer which provides additional insulating properties. The main problem with such materials is the complexity in the application, which makes it quite costly on an installed basis [39].

5.3.4 Depressurisation

Reducing the pressure in a vessel exposed to fire leads to a reduction of the stress on the metal and removes the danger of the vessel bursting. Also, this method may be used to prevent damage to items of equipment which are expensive to replace and may take a long time to obtain.

The designer needs to decide how quickly the pressure should be reduced and by what means.

Klaassen [44] suggests that means for vapour depressurising should be provided on all vessels, handling flammable liquids which operates above 3 bar and that the depressurising valve should be sized so that the pressure falls to half of design pressure in ten minutes. The American Petroleum Institute [46] recommended that the pressure should be reduced to 8 bar or 50 percent of design, whichever is lower, in 15 minutes.

The stress to rupture depends on the temperature, falling rapidly for carbon steels at temperatures above about 500 °C. It is therefore important to know the temperature that can be attained by an unwetted vessel exposed to fire. The API [47] assumes 690 °C, but in theory, much higher temperatures can be reached [43]. Assuming a temperature, and using Figure A3 in "Design and Installation of Pressure Relieving Systems in Refineries", can provide guidance on the percent of depressurising. As an example with an assumed temperature of 750 °C and using the above mentioned Figure for carbon steel vessels designed to BS 1515 shows a need to depressurise to about 20 percent of design pressure.

As the steel subject to higher stresses can fail more quickly at the temperature assumed, the aim should be to achieve this pressure within ten minutes. This is the shortest time in which a vessel has been known to burst when exposed to a fire [43].

The allowable depressurising time is affected by insulation and sloping the ground. If the vessel is insulated the rate of heat input is reduced and this time can be extended to an extent which depends on the quality of the insulation. Therefore, an insulated vessel should be depressurised within thirty minutes. Also the heat input into the vessel will be reduced when the ground under the vessel is sloped. This will lead to an increase in the depressurising time to twenty minutes.

If a vessel is insulated and the ground is sloped, then the depressurising time becomes sixty minutes. However, although this may be adequate to protect the vessel, another purpose of de-pressurising is to reduce the size of any leak which occurs in the vessel or its connecting pipework.

Vapour depressurising can be achieved by de-pressurising the vessel through existing lines installed for process purposes. If these are not present, then a relief valve by-pass may be installed. The depressurising valves, whether part of the process or specially-installed, must be operable in a fire situation. This usually means that they have to be remotely operated by electric or pneumatic power. The operating button must be well away from the vessel or it may not be accessible when a fire occurs.

Instead of a relief valve by-pass, a relief valve can be obtained which can be lifted by a remotely-operated pneumatic device. When this device does not operate, the valve functions as a normal relief valve.

It is not necessary to provide depressurising facilities on every vessel. If a group of vessels are normally connected together they can be depressurised as a whole.

The material discharged through a vapour depressurising valve, like the material discharged through a relief valve, must be disposed of in a safe way. It may be discharged to a flare system or it may be

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discharged through a vent or tail-pipe at such velocity that good mixing occurs. In the case of fire on the tail-pipe of the vapour depressurising valve due to formation of a cloud of flammable vapour, steam dispersion may be installed.

Finally the tail-pipe should be positioned so that any flame on it cannot impinge on other equipment.

CHAPTER SIX

EVALUATION OF THERMAL RADIATION HAZARD RANGES

A number of accidents have occurred leading to a fireball. These have involved the release of the contents from pressurised installations, or storage, or rail and road tankers containing LPG fuels. A list of incidents for which quantitative data are available is shown in Table 6.1 [48], together with the source of that information.

In evaluation of the thermal radiation hazard represented by a fireball, there are four primary parameters which are relevant [48]:-

- i the mass of fuel involved
- ii the fireball diameter
- iii the duration time, and
- iv the surface temperature

The consequences of an LPG fireball which must be considered as serious hazards are:

- i a Major fire
- ii an Unconfined Vapour Cloud Explosion (UVCE)
- iii a Boiling Liquid Expanding Vapour Explosion (BLEVE)

6.1 FIREBALL CHARACTERISTICS

The rapid combustion of flammable vapour in a spherical envelope is generally termed a fireball. The strong buoyancy forces of the hot, burning gases result in turbulent mixing and the formation of a mushroom-shaped cloud, often with a 'stem' of flame emanating

	and the second second					
Reference	Date	Place	Fuel container	Fuel	Quantity involved (tonnes)	Reported fireball diameter (m)
49	21.6.70	Crescent City Illinois, USA	Rail tank car	Propane	75	150-200
50	9.3.72	Lynchburg Virginia, USA	Tank truck	Propane	6	120
51	5.7.73	Kingman, Virginia, USA	Tank car	Propane	45	300
52	11.1.74	West St Paul, Minnesota, USA	Tank	DATI	10	100
53	17.1.74	Aberdeen, Scotland, UK	Road tanker	Butane	2	70
54	26.11.76	Belt, Montana USA	Rail tank car	DAT	80	300
55	28.12.77	Goldanna, Indiana, USA	Rail tank car	DAT	70	300-350
38	11.7.1978	San Carlos, Spain	Road tanker	Propylene	23.5	125
38	29.4.79	Eagle Pass, Texas, USA	Storage tank	DdT	25	340
16	19.11.84	San Juanico, Mexico City	Storage tanks	DdT	2500-3000	300

DATA ON SOME INCIDENTS RESULTING IN FIREBALLS

TABLE 6.1

from the source of the fuel. It is the high degree of turbulent mixing and rapid air entrainment which allows large quantities of fuel to be consumed in a short period [48].

Large quantities of heat are radiated from the hot mass of burning gases and the high radiant flux levels represent the major hazard from fireballs, extending well beyond the region that the flame envelopes. The main parameters relevant to a fireball are presented in the following sections.

6.1.1 Fireball Duration

The duration of the fireball is related to the mass of combustible fuel by the relationship;

$$t = A m^a \qquad \dots 6.1$$

where A and a are constants and m is the mass of fuel (kg). Table 6.2 and 6.3 [56] summarise the constants from reported data. Whilst there is some agreement between the various researchers this is not evidence for the adequate measurement of fireball duration since it can be partly attributed to the way in which the duration is defined by different researchers [48].

Hasegawa and Sato [57] defined an effective fireball duration as the time for which the fireball area exceeded half the maximum value whereas Fay's data [58] for 'complete combustion' of the fuel, and the data presented by Hardee *et al* [59], are for the combustion time until the fireball starts to lift off the ground. He suggests that the total combustion time will be about twice the lift-off time and this is consistent with the total duration data presented by Fay & Lewis [48].

TABLE 6.2

Flammable	Scale	A	a	В	b	Authors
Methane	0.1-10 kg	2.57	1/6	6.36	0.325	Hardee et al
Propane	0.037-0.37 g	2.53	1/6	6.28	1/3	Fay and Lewis
Pentane	0.3-6.2 kg	1.10	0.1	5.28	0.277	Hasegawa
Propene	0.124-452 kg	0.32	1/3	3.51	1/3	Maurer et al

VALUES OF THE PARAMETERS IN EQUATION 6.1 AND 6.2 FOUND EXPERIMENTALLY

TABLE 6.3

VALUES OF PARAMETERS IN EQUATION 6.1 AND 6.2 PROPOSED IN VARIOUS REVIEW PAPERS

Authors	Α	а	В	b
Gayle	0.245	0.356	3.68	0.326
Brasie	0.30	1/3	3.8	1/3
Marshall	0.38	1/3	5.5	1/3
Roberts	0.45	1/3	5.8	1/3

The difference between the exponents obtained may well be explained by the scale of the experiments. Small scale studies on quiescent fireballs by Fay [58], Hardee [59] and pressurised fuels by Hasegawa [57] all show low values while Maures [60] represents a higher exponent value. This implies that very small scale laboratory fireballs do not provide an accurate guide to the duration of larger scale events.

A computer program was written to enable rapid calculation of fireball duration using different authors' parameters; this is given in Appendix B. Typical results are shown in Table 6.4.

6.1.2 Fireball Diameter

Experimental studies of fireballs have been carried out with small quatities of quiescent gaseous fuels contained in soap bubbles by Fay and Lewis [58], Lihou and Maund [56], and in balloons and plastic bags as done by Hardee *et al* [59].

Studies have also been carried out by Hasegawa and Sato [57], Maurer *et al* [60], in which the fuel was rapidly released from a vessel after super-heating to a selected pressure.

The relationship obtained by different authors, relating maximum diameter with fuel mass is illustrated in equation 6.2. The parameters quoted in Table 6.2 and 6.3 are for a mass m (kg) of flammable producing a fireball of diameter Ds (m).

$$D_{\rm S} = {\rm Bm}^{\rm b} \qquad \dots \quad 6.2$$

There is considerable disagreement between various authors concerning the correlation which should be used to predict the maxi-

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TABLE 6.4

Mass (kg)	Gayle	Brasie	Marshall	Robert	Hardee	Fay
5	0.4	0.5	0.6	0.8	3.4	3.3
50	1.0	1.1	1.4	1.7	4.9	4.8
100	1.3	1.4	1.8	2.1	5.5	5.4
500	2.2	2.4	3.0	3.6	7.2	7.1
1,000	2.9	3.0	3.8	4.5	8.1	8.0
10,000	6.5	6.4	8.2	9.7	11.9	11.7
50,000	11.5	11.0	13.9	16.5	15.5	15.2
100,000	14.8	13.9	17.6	20.8	17.4	17.1
500,000	26.2	23.7	30.0	35.6	22.7	22.3

TYPICAL RESULTS OF FIREBALL DURATION (SEC) USING DIFFERENT AUTHORS' PARAMETERS

mum diameter of the fireball as shown in Tables 6.2 and 6.3.

A computer program was written to enable rapid calculation of fireball diameter using different authors' parameters; this is given in Appendix B. Typical results are shown in Table 6.5.

6.2 HAZARDS FROM FIREBALLS

Fireballs are hazardous because they are very hot, may be very large, and move in the atmosphere on currents of air. They are, therefore, a source of ignition and may ignite any combustible material in their path. More serious is their potential effect on people who can suffer severe burns from thermal radiation, even at some distance away from the fireball.

For example, the disaster on the Spanish Campsite in 1978 in which many people were killed and injured occurred when a cloud of propene escaped from a tanker and became a fireball when it found a source of ignition [56]. People were severely burnt by thermal radiation even some distance from the fireball because they were scantily clad. From the work of Stoll and Chianta [61] on the thermal radiation damage to pig skin, the following relationship has been derived. It enables the steady incident flux density q_2 (kW/m²) to be related to the duration te (seconds) which will cause severe blistering

$$q_2 = 50/te^{0.71}$$
6.3

Stoll's equation has been used [56] in the estimation of spacing distances in previous work. A new correlation on thermal radiation damage to human skin has been obtained and this will be described in Section 6.3.

TABLE 6.5

Mass (kg)	Gayle	Brasie	Marshall	Robert	Hardee	Fay
5	6.2	6.5	9.4	9.9	10.7	10.7
50	13.2	14.0	20.2	21.3	22.7	23.1
100	16.5	17.6	25.5	26.9	28.4	29.1
500	27.9	30.1	43.6	45.9	47.9	49.7
1,000	35.0	37.9	54.9	57.9	60.0	62.7
10,000	74.1	81.6	118.1	124.6	126.9	134.9
50,000	125.2	139.5	201.9	212.9	214.1	230.5
100,000	157.0	175.7	254.3	268.2	268.2	290.4
500,000	265.3	300.3	434.6	458.3	452.5	496.3

TYPICAL RESULTS OF FIREBALL DIAMETER (m) USING DIFFERENT AUTHORS' PARAMETERS

With a steady radiant flux, secondary fires in buildings can occur when the flux density is 12.6 kW/m² or more. Process plants and flammable storage tanks suffer severe damage at 37.8 kW/m² [62]. As a general rule of thumb, one may assume that under the steady flux densities quoted above, building and process equipment will ignite after 1000 seconds (about 17 minutes) [56]. Thus the radiation dosage for secondary fires may be set out as follows;

For buildings:

 $12.6 \text{ kW/m}^2 \text{ x } 1000 = 12.6 \text{ MJ/m}^2 \qquad \dots \quad 6.4$

For process equipment

 $37.8 \text{ kW/m}^2 \text{ x } 1000 = 37.8 \text{ MJ/m}^2 \dots 6.5$

6.3 THERMAL RADIATION DAMAGE CRITERIA

In estimating the hazard due to thermal radiation from LPG flames, calculation of the radiant intensity variation with distance from the flame is an important criterion. However, to quantify the hazards or to evaluate the risk to population and property, the damage criteria have to be considered.

In general, damage to skin exposed to thermal radiation depends on the intensity level and the duration of exposure, the protection offered by clothes and the natural cooling by perspiration and increased blood flow.

From the work of Stoll and Green [63] which includes exposure of human forearms to different intensities of radiation, the times required to produce threshold pain, threshold blister and full blister were determined.

Their data have been correlated in the present work using a computer curve fitting technique, as shown in Appendix C, to give three relationships which enable the thermal radiation incident on human skin to be related to the time of exposure as shown below.

Pain Threshold:	$q = 28.2 (t)^{-0.73}$	 6.6
Threshold Blister:	$q = 60.5 (t)^{-0.75}$	 6.7
Full Blister:	$q = 61.8 (t)^{-0.73}$	 6.8

where q is the radiation intensity (kW/m^2) and t is the time of exposure (sec).

Furthermore the data from their work which represents tissue damage for the threshold blister and full blister have been correlated using the same technique to give two relationships between the damage to tissue and the thermal radiation incident on human skin as shown below.

Threshold Blister:	$Z = 2.61 (q)^{1.09}$	 6.9
Full Blister:	$Z = 3.65 (q)^{1.31}$	 6.10

where Z represents the percent of tissue damage.

6.4 PREDICTION OF THERMAL RADIATION HAZARDS

Heat transfer from the fireball to the target will be primarily by thermal radiation. The radiation flux at a target q_2 (kW/m²) is given by the equation:

$$q_2 = F_{21} \tau \alpha q_1 \qquad \dots \ 6.11$$

where $q_1 (kW/m^2)$ is the emitted flux from the surface of the fireball, τ is the atmospheric transmissivity, α the absorptivity of the target and F_{21} the view factor.

Transmissivity can be estimated from the following equation [56]

$$\tau = \exp(-7 \times 10^{-4} \text{ L})$$
 6.12

where L is the distance from the edge of the fireball to the target area (m). The attenuation constant for the atmosphere can vary in value from 0.4 km^{-1} to 1.0 km^{-1} between a clear and a hazy day respectively [56]. An average value of 0.7 km^{-1} has been assumed in equation 6.12 and L can be found from the following equation.

$$L = \left[\left\{ \left(\frac{H}{D_s} + 0.5 \right)^2 + \left(\frac{H}{D_s} \right)^2 \right\}^{1/2} - 0.5 \right] D_s \qquad \dots 6.13$$

where Ds is the spherical fireball diameter, H the height of the fireball (m) and R the radial distance (m).

For a spherical fireball of diameter Ds (m) and which has cleared the ground by a height of H (m), the value of F_{21} for a target at a radial distance R (m) from a point directly below the centre of the fireball is given by equation 6.14 which was used to prepare Table 6.6 [56]. TABLE 6.6

DIRECT EXCHANGE FACTORS FOR SPHERICAL FIREBALLS OF DIAMETER Ds TO TARGETS AT

RANGE R, WHEN THE FIREBALL HAS CLEARED THE GROUND BY H

0044 0052 0900 0067 0073 6100. 0084 0089 6000 9600. 1600 6600 .0104 .0106 0106 .0107 0105 0103 0100 7000 3.0 .0100 .0106 .0115 .0119 0057 0067 .0113 2.75 0077 0085 0093 0111 .0122 .0126 .0127 .0125 0122 0118 0114 0109 .0127 0075 0088 0100 0120 0128 0135 .0143 0145 0149 .0154 .0153 .0147 0111 0141 0151 0135 0151 0141 0129 0123 2.5 2.25 .010 .012 .015 .016 .018 019 013 .017 .018 018 019 019 019 019 018 .016 .016 .017 .015 014 014 .016 018 .020 022 023 .024 .024 .024 024 024 .023 023 019 021 .022 020 018 015 2.0 .017 019 029 029 022 024 026 028 .030 .030 030 030 029 .028 026 018 1.8 025 017 023 020 021 .030 .033 035 036 .037 .038 F21: Incident Flux Density/Emitted Flux Density .038 036 036 1.6 .033 029 018 027 .037 037 031 026 024 022 020 048 049 049 049 1.4 038 042 046 .048 .047 .047 .045 .043 .040 .036 .033 029 .027 024 022 020 066 059 1.2 062 065 600 6007 064 .056 .053 .047 042 029 057 061 090 .037 .033 026 023 021 610. 1.0 095 960 095 092 .088 .084 076 .074 690. 048 089 .064 .055 .042 .036 .032 026 028 022 .115 .119 .115 118 .103 080 .109 060 .086 .082 .076 070. 090. .023 .051 .044 .038 .033 029 026 6 .149 .150 .146 138 129 109 076 121 100 960 084 091 064 054 046 040 035 030 027 024 œ. .180 .166 152 .137 .112 196 191 124 106 .083 .068 .057 048 .036 .027 101. 091 041 031 .024 2 246 .233 200 262 .178 .158 .140 .117 660 124 080. .073 090. .050 .043 111. .037 .032 028 .025 9. 315 275 238 156 128 354 206 179 137 120 106 095 690 063 052 025 944 038 033 028 Ś .716 546 340 276 228 .192 426 .163 .140 .122 .107 .056 .151 .084 790. 046 039 .026 .25 .034 029 8 694 .510 309 .250 207 .174 .160 .148 .128 391 690. III 087 .057 047 040 034 030 026 0 R/Ds H/Ds 0.75 0.6 0.7 0.2 0.3 0.4 0.5 0.8 6.0 1.0 1.2 1.4 1.6 1.8 2.0 2.2 2.6 0.1 2.4 0

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$$F_{21} = \frac{(H/D_s + 0.5)}{4\{(H/D_s + 0.5)^2 + (R/D_s)^2\}^{3/2}} \dots 6.14$$

In this work τ and α have been taken for simplicity as unity although for long hazard ranges τ can have a value significantly less than one [19].

The emitted flux density q_1 from the surface of a fireball is dependent upon the fourth power of its absolute temperature $Ts(^{\circ}K)$ as given by the following equation

$$q_1 = \varepsilon \theta T_s^4 \qquad \dots \quad 6.15$$

where ε is the emissivity and θ the Stefan-Boltzmann constant.

Recently surface temperatures were measured by two-colour pyrometry from colour cine films of fireballs which were made by igniting elevated hemispherical detergent bubbles filled with butane and with natural gas in the range 100 to 800 ml [56]. Empirical correlations with mass of flammable were found for maximum spherical diameter, maximum elevation and duration of the visible fireballs.

One way of assessing large-scale phenomena on a small scale is by modelling. Two models based upon material and heat balances have been developed by Lihou [56], which fit the observed diameter and elapsed time relationships. Both models are based upon the experimental observation that the diameters of fireballs increased at a constant rate. Both models will be described in the following section.

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6.5 MATHEMATICAL MODELS FOR HEAT RADIATION6.5.1 Hot Shell-Cold Core Model

In this model the temperature at the core of the fireball is assumed to be ambient temperature, ie 290 °K. Prior to neutral buoyancy, for a butane fireball, the core is assumed to contain no air. But when a fireball begins to rise air enters the core without being heated. Air required for combustion is consumed at the hot shell [56].

6.5.1.1 Pre-lift off Combustion of Butane

A heat balance for a butane cloud initially at 290 °K can be written as follows:

combustion rate = sensible heat to air + radiative heat loss

$$m h \frac{df}{dt} = mrC (T_s-290) \frac{df}{dt} + \pi D^2 \theta \epsilon T_s^4 \qquad \dots 6.16$$

where m: mass of flammable originally in the cloud,

f: fraction of fuel burnt in time t

- h: net calorific value
- r: mass ratio of combustion products from the stoichiometric combustion of unit mass of the flammable gas

C: mean specific heat of combustion product

D: equivalent spherical diameter

Ts: temperature of the hot shell

ε: emissivity

θ: Stefan-Boltzmann constant.

It has been noted that throughout combustion, the equivalent spherical diameter increases at a constant rate $(dD/dt)_c$. Rearranging equation 6.16 will give:

mh - mrC (T_s-290) df =
$$\pi D^2$$
 dt $\theta \epsilon$ T_s⁴

Cancelling dt by making the substitution:

$$D^{2}dt = D^{2} dD (dt/dD)_{c}$$
 6.17

integration can be carried out, with the boundary conditions f = owhen D = Do where

$$Do = 3.57 (m/M)^{1/3}$$
 6.18

and M is the molecular weight of the flammable gas, then

$$[\text{mh-mrC} (T_s - 290)]f = \pi/3 \theta \epsilon T_s^4 (D^3 - D_0^3) (dt/dD)_c$$

therefore

1

$$f = \frac{\pi \,\theta \epsilon \,T_s^4 \,(D^3 - D_0^3)}{3m\{h - rC \,(T_s - 290)\} \,(dD/dt)_c} \qquad \dots 6.19$$

At neutral buoyancy the fireball density equals that of the surrounding air. At 290 °K with a mean molecular weight of 29 this has a density of [273/290 x 29/22.4 = 1.22] kg/m³. Let $f = f_1$ and $D = D_1$ at liftoff, the fireball mass is $m(1+rf_1)$ and its volume is $\pi/6 D_1^3$. Equating its density to 273/224. density = mass/volume

$$1.22 = \frac{m(1+rf_1)}{\pi/6 D_1^3} \qquad \dots \quad 6.20$$

rearranging equation 6.20 gives,

$$D_1^3 = 1.567 (1+rf_1)m$$
 6.21

Substituting f_1 and D_1^3 in equation 6.19 and substituting $\theta = 56.7 \times 10^{-12} \text{ kW/m}^2 \text{.k}^4$ gives the following general equation for denser than air gases;

$$f_1 = \frac{93.05 \varepsilon (T_s/1000)^4 (1 + rf_1 - 29/M)}{\{h - rC (T_s - 290)\} (dD/dt)_c} \qquad \dots 6.22$$

To obtain the relationships for butane the values of r, C, M, h, ε and Ts should be known, for example for butane [56] r = 17.14, C = 1.14 kJ/kg °K, M = 58, h = 45920 kJ/kg and from the experimental work Ts = 1300 K, ε = 1 and $(dD/dt)_c$ = 1 m/sec. Substituting these values in equations 6.22 gives f_1 = 0.00614. From equations 6.18 and 6.21 and putting (dD/dt)c = 1 m/sec give the following relationships for butane

$D_0 = 0.92 m^{1/3}$		5.23
$D_1 = 1.20 m^{1/3}$		6.24
$t_1 = 0.28 m^{1/3}$		6.25

6.5.1.2 Dilution and Combustion

During the completion of combustion of butane clouds, and throughout the combustion of methane clouds, air at 290K enters the core at an average radial velocity $\overline{U}(m/sec)$ through the hot shell. \overline{M} is the average molecular weight of a stoichiometric mixture of combustion products. The rate of increase of volume can be expressed as follows:

$$\frac{\pi D^2}{2} \left(\frac{dD}{dt}\right)_c = \frac{mr}{\overline{M}} \times \frac{22.4}{273} (T_s - 290) \frac{df}{dt} + \pi D^2 \overline{U} \qquad \dots 6.26$$

the heat balance shown as equation 6.16 still applies, because the excess air enters the core without being heated. Combining equations 6.16 and 6.26 gives the following heat and mass balance relationship:

$$mh\frac{df}{dt} = \pi D^{2} \left[\theta \varepsilon T_{s}^{4} + \frac{273\overline{M}C}{22.4} \left\{ 0.5 \left(\frac{dD}{dt}\right)_{c} - \overline{U} \right\} \right] \qquad \dots 6.27$$

the boundary conditions are $f=f_1$ at $D=D_1$; which is given by equation 6.21. This equation can be used to eliminate m, having made the substitution shown as equation 6.17 and integrating

$$f = f_1 + \frac{448 (1 + rf_1)}{273h \left(\frac{dD}{dt}\right)_c} \left[\left(\frac{D}{D_1}\right)^3 - 1 \right] x Q \qquad \dots 6.28$$

where Q = heat flux density function

$$Q = \theta \varepsilon T_s^4 + \frac{273\overline{M}C}{22.4} \left[0.5 \left(\frac{dD}{dt}\right)_c - \overline{U} \right] \qquad \dots \quad 6.29$$

Let D=Dc at the completion of combustion when f = 1.

The experimental data for butane is best fitted by putting $\overline{U} = 0.5$ (dD/dt)c, which eliminates the second term in equation 6.29. Additional constants for butane are $\overline{M} = 28.4$, $f_1 = 0.00614$ and $D_1=1.2 \text{ m}^{1/3}$. Substitution in equation 6.28 gives

$$D_c = 6.45 \text{ m}^{1/3}$$
 6.30

since $tc = \frac{Dc-Do}{(dD/dt)_c}$ and $\left(\frac{dD}{dt}\right)_c = 1$ m/sec

So the following expression can be obtained for the duration of the combustion phase

$$tc = 5.53 m^{1/3}$$
 6.31

For methane, the boundary conditions for equation 6.27 are:

$$f = 0$$
 at $D = D_0$

in which D_0 is found from equation 6.18 with M = 16.

Other constants are [56] r=19, C=1.22, \overline{M} =27.6, h=50213 kJ/kg and from the experiments the following mean values were found: Ts=1500K, ε = 0.5, (dD/dt)_c = 1 m/sec. These parameters used in equation 6.32 with f=1 at D=Dc and t=t_c, with Q given by equation 6.29 when $\overline{U}=0.5 (dD/dt)_c$ will produce equations 6.33 and 6.34.

$$f = \frac{44.8 \times 290}{273 Mh(dD/dt)_c} \left[\left(\frac{D}{Do} \right)^3 - 1 \right] \times Q \qquad \dots 6.32$$

$$D_{o} = 1.42 \text{ m}^{1/3} \qquad \dots \quad 6.33$$
$$D_{c} = 6.95 \text{ m}^{1/3} \qquad \dots \quad 6.34$$
and $t_{c} = Dc - Do/(dD/dt)_{c} = Dc - Do \qquad \dots \quad 6.35$ since $(dD/dt)c = 1$

The computer program presented in Appendix G was used to evaluate the fireball dimensions and hazard ranges by using this model. The predicted results are presented in the same Appendix. These results show that this model cannot be used for large quantities of hydrocarbon. However Lihou [56] suggested that this model gave a satisfactory result for small-scale experiments, ie in the range of 200-800 ml (0.134-0.536 gm).

6.5.2 Isothermal Model

In this model, the assumptions are [56]:

- i The whole fireball burns at isothermal temperature (Tc)
- ii Combustion is controlled by the supply of air and ceases after a time (tc) which is correlated empirically with the mass (m) of the flammable gas initially in the cloud.
- iii The equivalent spherical diameter starts at Do and increases at a constant rate (dD/dt)_c up to a diameter Dc at which the fuel has all disappeared.
- iv A fraction $(1-f_c)$ of the fuel is assumed to be cracked to produce

soot, the volume of which is neglected.

The final diameter (Dc) is found for a stoichiometric mixture of combustion products from the burnt fraction (fc) of the fuel plus a fraction of residual air, all at temperature of Tc.

Since the volume of a sphere is

$$\overline{V} = (\pi/6)D^3$$
6.36

differentiating the above equation to obtain the equation which presents the rate of increase of volume, as follows

$$\frac{\pi}{6} D^3 \left(\frac{dD}{dt}\right)_c = \frac{\pi}{2} D^2 \left(\frac{dD}{dt}\right) = \frac{VT_c}{290} \left(1 + \frac{nf_c}{v}\right) \qquad \dots \quad 6.37$$

where v	=	the stoichiometric molar ratio of air to flammable gas
n	=	the increase in the total number of moles per mole of flammable gas burnt stoichiometrically
v	=	volumetric rate at which air flows into the fireball (m^{3}/sec)
D	=	equivalent spherical diameter
fc	=	fraction of fuel burnt at the end of combustion

Rearranging the above equation gives

$$V = \frac{290 \pi D^2 (dD/dt)_c}{2T_c (1 + nf_c/v)} \dots 6.38$$

A heat balance equation can be written as follows:

combustion rate = sensible heat to air + radiative heat loss

combustion rate =
$$\frac{273 \text{ VMh } f_c}{290 \text{ x } 22.4 \text{ x } \upsilon} \qquad \dots \quad 6.39$$

substitution for V from equation 6.38 gives

combustion rate =
$$\frac{273 \text{ Mh } f_c \left[\frac{290 \pi D^2 (dD/dt)_c}{2T_c (1 + n f_c/v)} \right]}{290 \times 22.4 \times v}$$

sensible heat =
$$\frac{273 \times 29 \times V C_a (T_c-290)}{290 \times 22.4} \qquad \dots \quad 6.40$$

substitution for V from equation 6.38 gives

sensible heat =
$$\frac{273 \times 29 \times C_a (T_c - 290) \times \left[\frac{290 \pi D^2 (dD/dt)_c}{2 T_c (1 + nfc/v)}\right]}{290 \times 22.4}$$

$$= \frac{176.71 \text{ C}_{a} (\text{T}_{c} - 290) \pi \text{D}^{2} (\text{dD/dt})_{c}}{\text{Tc} (1 + \text{nfc/y})}$$

radiative heat loss = $\pi D^2 \epsilon \theta T_c^4$

substituting $\theta = 56.7 \text{ x } 10^{-12} \text{ kW/m}^2 \text{k}^4$ and Cair = 1.09 kJ/kg °K, rewriting the heat balance equation:

.... 6.41

6.09 Mhf_c πD^2 (dD/dt)_c 176.71x1.09 (T_c.290) x πD^2 (dD/dt)_c

$$T_{c} (\upsilon + nf_{c}) = -----+ 56.7 \times 10^{-12} \varepsilon \pi D^{2} T_{c}^{4}$$
$$T_{c} (\upsilon + nf_{c}) = -------+ 56.7 \times 10^{-12} \varepsilon \pi D^{2} T_{c}^{4}$$
$$\dots 6.42$$

cancelling πD^2 from above equation gives:

 $\frac{6.09 \text{ Mhf}_{c}}{T_{c} (\upsilon + nf_{c})} \frac{dD}{dt} - \frac{192.6 (Tc-290)}{T_{c} (\upsilon + nf_{c})} \frac{dD}{dt} - \frac{192.6 (Tc-290)}{(--)_{c}} \frac{dD}{(--)_{c}} = 56.7 \times 10^{-12} \varepsilon T_{c}^{4}$

rearranging gives

 $\frac{6.09 \text{ Mhf}_{c}}{T_{c} (\upsilon + nf_{c})} \frac{dD}{dt} - \frac{192.6 \upsilon (Tc-290)}{T_{c} (\upsilon + nf_{c})} \frac{dD}{dt} - \frac{192.6 \upsilon (Tc-290)}{(--)_{c}} \frac{dD}{(--)_{c}} = 56.7 \times 10^{-12} \varepsilon T_{c}^{4}$

$$\frac{6.09 \text{ Mhf}_{c} - 192.6 \text{ v} (\text{T}_{c} - 290)}{\text{T}_{c} (\text{v} + \text{nf}_{c})} \left[\frac{\text{dD}}{\text{dt}} \right]_{c} = 56.7 \text{ x} 10^{-12} \varepsilon \text{T}_{c}^{4}$$

$$\left[\frac{6.09 \text{ Mhf}_{c} - 31.6 \upsilon (T_{c} - 290)}{T_{c} (\upsilon + nf_{c})}\right] \left[\frac{dD}{dt}\right]_{c} = 9.30 \varepsilon \left(\frac{T_{c}}{1000}\right)^{4}$$

.... 6.43

for a gas cloud of mass m, molecular weight M, initially at 290°K, the initial diameter is given as follows:

volume of sphere
$$(\overline{V}) = \frac{\pi}{6} D^3 = \frac{\text{mass (m)}}{\text{density (p)}} \dots 6.44$$

so D =
$$\left(\frac{6V}{\pi}\right)^{1/3}$$

= $\left(\frac{6m}{\pi\rho_T}\right)^{1/3}$

where

$$\rho_T$$
 = density at given temperature

since
$$\rho_{\rm T} = \rho_0 x \frac{298}{T + 273}$$

and taking T = 17 °C, $\rho_0 = M/22.4$

so
$$\rho_{\rm T} = \frac{M}{22.4} \times \frac{298}{290}$$

then D_o =
$$\left[\frac{6 \text{ m}}{\pi \text{ x} \frac{\text{M}}{22.4} \text{ x} \frac{298}{290}}\right]^{\frac{1}{3}}$$

ie
$$Do = 0.539 \left[\frac{290 \text{ m}}{M}\right]^{1/3}$$
 6.45

The final diameter is given as shown below

$$Dc = 0.539 \{ v + (n+1) f_c \}^{1/3} \{ \frac{Tc m}{M} \}^{1/3} \dots 6.46$$

The constant rate of increase of diameter is given by:

$$\left(\frac{dD}{dt}\right)_{c} = \frac{D_{c} - D_{o}}{t_{c}} = \frac{0.539 (m/M)^{\frac{1}{3}} (T_{c} - 290)^{\frac{1}{3}} \{\upsilon + (n+1)f_{c}\}^{\frac{1}{3}}}{t_{c}} \dots 6.47$$

1

substituting for (dD/dt)c from equation 6.47 into the heat and material balance equation, gives

9.30
$$\varepsilon t_c \left(\frac{T_c}{1000}\right)^4 = \left[\frac{Mhfc - 31.6 \upsilon (T_c - 290)}{T_c (\upsilon + nfc)}\right] \left[\frac{0.539m (T_c - 290) \{\upsilon + (n+1)fc\}}{M}\right]^{\frac{1}{3}}$$

since $tc = Am^{1/3}$ from equation 6.1,

so the heat and material balance equation becomes as follows:

9.30
$$\varepsilon Am^{\frac{1}{3}} \left(\frac{T_c}{1000}\right)^4 = \left[\frac{Mhfc-31.61\upsilon(T_c-290)}{T_c(\upsilon+nfc)}\right] \left[0.539 \text{ m}\right]^{\frac{1}{3}} \left[\frac{(T_c-290)\{\upsilon+(n+1)fc\}}{M}\right]^{\frac{1}{3}}$$

Cancelling $m^{l/3}$ and rearranging the above equation gives the final heat and mass balance equation from which the fireball temperature is related to A and f_c .

$$17.26 \text{ A}\epsilon \left(\frac{T_c}{1000}\right)^4 = \left[\frac{\text{Mhfc}-31.61\upsilon(T_c-290)}{T_c(\upsilon+nfc)}\right] \left[\frac{\{\upsilon+(n+1)fc\}\{T_c-290\}\}}{M}\right]^{\frac{1}{3}}$$
.... 6.48

A computer program was written to enable the rapid calculation

using this method and different authors' parameters to evaluate the fireball surface temperature. This program is given in Appendix D.

Insertion of the parameters proposed by several authors into the isothermal model gave values of B which were reasonably close to the values which they recommended as shown on Table 6.3. This is illustrated by the following example.

Taking the value of ε , h, M, υ , n and fc from Table 6.7 and substituting these in equation 6.48 and taking the value of A recommended by Roberts [64] in Table 6.3, gives the value of Tc.

For example, for butane $\epsilon = 1$, h = 45806 kJ/kg, M = 58, $\upsilon = 30.95$, n = 1.5, $f_c = 0.95$ and taking A = 0.45, gives Tc = 2016 °K.

Taking $f_c = 0.95$ and Tc = 2016 °K, equation 6.46 gives the following expression for the final diameter of a butane fireball, which is in good agreement with the value of B recommended by Roberts [64] in Table 6.3.

$$Dc = 5.66 \text{ m}^{1/3}$$
 6.49

.... 6.50

and
$$t = 0.45 \text{ m}^{1/3}$$

The initial diameter Do is given by equation 6.45. The duration of combustion is given by equation 6.50 from which the rate of growth of diameter for large fireballs would appear to be about 10.5 m/sec.

Similarly the value of A recommended by other authors, Gayle [65], Brasie [66] and Marshall [67] can be substituted in equation 6.48 to give the corresponding value of Tc.

The final diameter of a butane fireball can be obtained from equation 6.46 which is in good agreement with the value of B recomm-

TABLE 6.7

BASIC SET OF PARAMETERS USED IN EVALUATION OF FIREBALL SURFACE TEMPERATURE USING THE ISOTHERMAL MODEL

Component	r	fc	n	υ	М	h(kJ/kg)
CH ₄	18.16	0.906	0	9.52	16	50,145
C ₃ H ₈	16.6	0.949	1	23.8	44	46,450
C_4H_{10}	16.39	0.95	1.5	30.95	58	45,806
C ₅ H ₁₂	16.25	0.955	2	38.09	72	45,446

ended by Marshall in Table 6.3, but not with the value of B recommended by both Gayle and Brasie as shown below.

Taking $f_c = 0.95$, A = 0.38, the value recommended by Marshall in Table 6.3, equation 6.48 gives Tc = 2064 °K, substitution of Tc in equation 6.46 gives the final diameter of a butane fireball as follows.

$$Dc = 5.7 m^{1/3}$$
 6.51

Taking $f_c = 0.95$ and A = 0.3, the value recommended by Brasie in Table 6.3, equation 6.48 gives Tc = 2132 °K, substitute Tc in equation 6.46 gives the following expression:

$$Dc = 5.76 \text{ m}^{1/3}$$
 6.52

Taking fc = 0.95 and A = 0.245, the value recommended by Gayle, use of equation 6.48 produces Tc = 2188 °K. Substitution of Tc into equation 6.46 gives the following expression:

$$Dc = 5.8 m^{1/3}$$
 6.53

Finally the rate of growth of fireball diameter for a large fireball using other authors' parameters is presented in Table 6.8. it appears that only Roberts and Marshall's parameters give close values to those achieved by Lihou's model.
TABLE 6.8 PREDICTED RATES OF GROWTH OF FIREBALL DIAMETER USING VARIOUS AUTHORS' PARAMETERS

Author	Rate of growth (dD/dt)c	
	(m/sec)	
Gayle	19.9	
Brasie	16	
Marshall	12	
Roberts	10.5	

6.6 BOILING LIQUID EXPANDING VAPOUR EXPLOSIONS (BLEVE)

6.6.1 Size of BLEVE

Several correlations have been proposed for calculating the dimensions of a fireball originating from a BLEVE of flammable material as shown below. Some of these correlations are derived from experiments:

$$D = Bm^b$$
 6.54

where D the diameter of the fireball (m)

m mass of fuel (kg)

Other correlations show that B depends not only on the mass of fuel, but also upon gas density as given in Lihou's [56] and Crawley's [36] correlations as shown below

$$D = B [m/M]^b$$
 6.55

Parameters for various authors which apply for both equations 6.54 and 6.55 are given in Table 6.9. In some equations the mass of fuel is in tonnes as used in Crawley's correlation [36].

TABLE 6.9

VALUES OF PARAMETERS PROPOSED BY DIFFERENT AUTHORS FOR USE IN EQUATIONS 6.54 AND 6.55

Author	В	b	
Maurer	3.51	1/3	
TNO	6.48	0.325	
Crawley	105	1/3	
Lihou	3.44	1/3	

A computer program was written to enable rapid calculation of BLEVE fireball diameters and duration using different authors' parameters, this is given in Appendix H. Typical results are shown in Figure 6.1.

A comparison can be made between the values of diameter obtained by applying the various correlations mentioned above and the values obtained in real incidents as described below:

Incident: 1. Oneonta [36], charge 80 te of LPG, diameter observed 122 m.



Figure 6.1 Variation of Fireball Diameter with Mass of Fuel based on Crawley, Maurer, Lihou and TNO correlations.

Mexico disaster [16], 1600 m³ sphere filled to 90% capacity, at density of 512 kg/m³, this would have amounted to 819,200 kg. Fireball dimensions, as shown in photographs and video, are estimated to have ranged from 200 to 300 m.

Comparison between predicted and observed results are shown in Table 6.10. These results show that the Crawley correlation gives a close result to those observed from the real BLEVE incidents mentioned above.

TABLE 6.10 COMPARISON BETWEEN CALCULATED AND OBSERVED FIREBALL DIAMETER

Mass of fuel	Diameter				
Tonne	Crawley	Maurer	TNO	Lihou	Observed
80	128	151	254	149	122
819	278	328	541	323	200-300

6.6.2 BLEVE Model

Based upon the work of Maurer *et al* [60], Lihou derived the following relationships for the duration and the diameter of a BLEVE fireball:

$$t_c = 0.31 \left[\frac{44.8 \text{ m}}{\text{M}} \right]^{\frac{1}{3}} \dots 6.56$$

$$D_{\rm c} = 3.44 \left[\frac{44.8 \text{ m}}{\text{M}} \right]^{\frac{1}{3}} \dots 6.57$$

The work of Maurer *et al* [60] indicates that the maximum amount of LPG which will burn in the time available for mixing with air is 70%. Only about 40% of the flammable mass (that is 28% of the original contents of the vessel) can be sufficiently well-mixed with air to undergo percussive deflagration. Thus 42% of the flammable, originally in the tank, produces a fireball.

Using equations 6.56 and 6.57 in a heat balance, based upon 42% of m undergoing isothermal stoichiometric combustion at Tc, with a constant diameter Ds, the following equation can be used to find Tc.

$$\frac{69700}{M} \left[\frac{T_c}{1000} \right]^4 = h - 1000 \text{ rC} \left[\frac{T_c}{1000} - 0.29 \right] \dots 6.58$$

where Tc isothermal temperature of the fireball (°K)

- h net calorific value of fuel (kJ/kg)
- C specific heat of combustion products (kJ/kg.°K)
- r mass ratio of combustion products per unit mass of flammable gas, from stoichiometric combustion.

A computer program was written to enable rapid calculation

using this method to evaluate the fireball surface temperature; this is given in Appendix E.

6.7 FIREBALL SURFACE TEMPERATURE

The fireball surface temperature can be predicted from the isothermal model, the BLEVE model as done by Lihou [56] and also by a new proposal from thermodynamic calculations. The temperature calculated from thermodynamic considerations is termed the 'theoretical flame temperature'.

Since the final temperature of the combustion products is unknown and must be calculated, the process is assumed to be adiabatic. With the added assumption that kinetic and potential energy terms are negligible, the overall energy balance for this process reduces to

It is assumed that the combustion takes place at 298 °K (25 °C) and that the heat passes into the reaction products adiabatically, heating them to the flame temperature which is assumed at first

since
$$\Delta H^{\circ}_{298} + \Delta H^{\circ}p = \Delta H = 0$$
 6.60
But $\Delta H^{\circ}_{298} = \Delta H^{\circ}c$ 6.61
and $\Delta Hp^{\circ} = \sum_{\text{products}} (nCp_m) (t-25)$ 6.62

Thus
$$\sum_{\text{products}} (nCp_m) (t-25) = \Delta H^{\circ}c$$
 6.63

where $\Delta H^{\circ}c =$ standard heat of combustion $Cp_{m} =$ mean specific heat of combustion product n = number of moles

Since the combustion products are carbon dioxide and water, in addition to the nitrogen, excess oxygen and unburned hydrocarbon then

$$\Delta H_{c}^{\circ} = [(n_{CO_{2}}Cp_{CO_{2}}) + (n_{H_{2}O}Cp_{H_{2}O}) + (n_{N_{2}}Cp_{N_{2}}) + (n_{O_{2}}Cp_{O_{2}}) + (n_{H.C}Cp_{H.C})] [t - 25]$$
.... 6.64

A computer program was written to enable rapid calculations for fireball surface temperature using this method, this is given in Appendix F.

6.8 ELEVATION

The experimental results [56] with elevated fireballs showed that the velocity of rise was very nearly $(dD/dt)_c$. For butane equations 6.45, 6.46 and 6.50 indicate that for large scale fireballs $(dD/dt)_c =$ 10.5 m/sec. Similarly using the same equations shows that for large scale methane fireballs $(dD/dt)_c = 9.9$ m/sec.

Taking a rate of rise of 10 m/sec as suggested by Lihou [56], the simple relationship for all large-scale fireballs is as follows:

H = 10 t

.... 6.65

In the BLEVE model, Lihou suggests [56] the following equation should be used in calculation of the hazard range to people from a BLEVE

$$H = 0.5 Ds$$
 6.66

The final elevation given by equation 6.59, using Robert's parameter, is as follows:

$$H = 4.5 m^{1/3}$$
 6.67

The above equation agrees quite well with the correlation by Fay and Lewis [58] for propane, which is shown below

 $H = 4.1 m^{1/3}$ 6.68

6.9 HAZARD RANGES

Hazard range can be predicted by using the appropriate correlation, then the duration of combustion can be used in equations 6.3 and 6.6 to obtain the limiting values of q_2 . Fireball surface temperature (Tc) can be obtained from the isothermal model, BLEVE model and thermodynamic calculation. Substitution of (Ts) in equation 6.15 evaluates q_1 . A computer program was written to enable rapid calculation for view factors and the ratio of height to fireball diameter, this is given in Appendix G. Table 6.6 is used to find the value of R/Ds for which F_{21} is approximately equal to the limiting values of q_2/q_1 . Hazard ranges are found by multiplying the predicted fireball diameter Ds(m) by the limiting value of R/Ds. Typically, allowance for attenuation will reduce the hazard ranges by up to 20%. Typical results for hazard ranges for 10 tonnes of butane are given in Table 6.11.

TABLE 6.11

TYPICAL RESULTS FOR HAZARD RANGES (m) FOR 10 TONNES BUTANE FIREBALL USING DIFFERENT AUTHORS' PARAMETERS

Author	human range	building range	process range
Gayle	218.61	207.49	133.39
Brasie	265.26	220.37	138.75
Marshall	378.02	307.14	189.01
Robert	404.87	323.89	193.09

6.10 POOL FIRES

A leak of flammable liquid ie oil, oil products or refrigerated LPG contained within a bund will produce a pool which, upon ignition will result in a pool fire.

The application of engineering calculations to separation distances for storage tanks has been treated by Hearfield [12].

Calculations of separation distances related to heat effects are usually based on direct flame impingement and on heat radiation.

It is commonly assumed that in a storage tank fire, the tank is effectively a pool of liquid with a flame burning on its surface and that on a tank of diameter D, the height of the flame in still air is 2D [62, 68].

If there is a wind, however, the flame is distorted, At a wind speed of 2m/s it has been observed that the flame is deformed to an angle of 45° to the horizontal and that on the downwind side of the fire the flame hugs the ground for a distance of about 0.5 D [62].

For the heat radiated from the flame it is usually assumed for simplicity that still-air conditions prevail [62]. The heat radiated is calculated and compared with limiting values.

An equation given by Hearfield [12] for the heat radiated from the envelope of a flame burning on a liquid pool in a storage tank is:

 $Q_1 = K A r \rho C$ 6.69

whe	ere A	the surface area of pool, m ²
	С	the calorific value of liquid, kJ/kg
	Q1	the quantity of heat radiated from the envelope, kW
	r	the liquid burning rate, m/s
	ρ	the density of liquid. kg/m ³
	K	constant (the value generally used by the author is 0.3)

The rate of burning can be predicted from the derivation of Moorhouse [48] which is based on a heat balance on a simple vertical cylindrical flame from a pool fire as shown below.

heat radiated to surroundings = flame surface area x surface emissive power

 $= [(\pi/4)D^2 + \pi DL].E \qquad \dots 6.70$

The above equation can be equated to the fraction of the total heat

liberated by combustion, as shown below

$$[(\pi/4)D^2 + \pi DL].E = f. \ddot{m}. Hc. \pi D^2/4 \dots 6.71$$

Rearranging the above equation, gives

D

$$\ddot{m} = \frac{L}{f Hc} (1 + 4 L/D)$$
 6.72

 E = flame surface emissive power, kW/m² Hc = heat of combustion, kJ/kg f = fraction of heat of combustion of fuel white radiated (the value suggested by the author range of 0.2-0.4) L/D = ratio of flame length to diameter 	vhere i	m	=	fuel burning rate, kg/m ² .sec
 Hc = heat of combustion, kJ/kg f = fraction of heat of combustion of fuel white radiated (the value suggested by the author range of 0.2-0.4) L/D = ratio of flame length to diameter 	E	E	=	flame surface emissive power, kW/m ²
 f = fraction of heat of combustion of fuel white radiated (the value suggested by the author range of 0.2-0.4) L/D = ratio of flame length to diameter 	F	Hc	=	heat of combustion, kJ/kg
I/D - ratio of flame length to diameter	f	f	=	fraction of heat of combustion of fuel which is radiated (the value suggested by the author in the range of $0.2-0.4$)
<u><u><u></u></u><u></u><u></u><u></u><u></u><u></u><u></u><u></u><u></u><u></u><u></u><u></u><u></u><u></u><u></u><u></u><u></u><u></u></u>	L	L/D	=	ratio of flame length to diameter

Taking the value of K = 0.3, f = 0.3, r = \dot{m}/ρ and substituting the value of \dot{m} from equation 6.72 in equation 6.69, gives

$$Q_1 = A. E (1 + 4 L/D)$$
 6.73

Since

$$\mathbf{E} = \varepsilon \boldsymbol{\Theta} \mathbf{T}_{\mathrm{f}}^{4} \qquad \dots \quad 6.74$$

where ϵ	=	emmissivity (for simplicity assumed unity)
θ	=	Stefan-Boltzmann constant (56.7 x 10 ⁻¹² kW/m ² .k ⁴)
Tf	=	flame surface temperature







Receiving Tank

Burning Tank .

Figure 6-2 Radiation from Pool Fires.

The application of equation 6.69 is described by Robertson [62] as follows. It is assumed that the tank roof has completely blow off presenting a pool fire where the diameter of the pool is the diameter of the tank, which is assumed to be cylindrical. If still air conditions are assumed the flame envelope is a cylinder of diameter D with a height of 2D as shown in Figure 6.2 [62]. It is further assumed that the heat radiated from the envelope is in turn equivalent to radiation from a radiating area which is a vertical rectangular plane through the diameter of the flame envelope of width D and height 2D. The flame temperature which is used in equation 6.74 is the flame temperature obtained for propane is 1840 °C (2113° K).

The heat flux received by an object varies roughly according to inverse square of the distance between the radiator and the receiver [62], which can be described for a point source as follows:

$$Q_2 = \frac{Q_1}{4\pi d^2}$$
 6.75

where Q_1 = the quantity of heat radiated, (kW) Q_2 = radiant flux intensity, kW/m² d = distance from tank to the receiver, m

There are three important radiation flux bands to be considered:

i

A flux of 37.8 kW/m² is the approximate maximum radiation any tank should receive from an adjacent tank fire.

ii

The second band of flux is 12.6 kW/m². This flux is the threshold limit over which the resins in wood, building roofing felt, oily rugs and paper will start to emit flammable vapours

which are easily ignited by flying sparks.

iii A flux of 4.7 kW/m² is the approximate threshold of pain on any exposed part of the human body. At this blistering of the skin takes place.

A computer program was written to assist rapid calculation for the distances from a burning tank to the flux band stated above; this is shown in Appendix I.

6.11 WATER REQUIREMENTS

In certain circumstances when a tank farm is constructed, it is not possible to leave a safe distance between each tank due to economic reasons and/or lack of space. Under these circumstances, it is necessary to install water sprinklers near the tanks to spray water onto adjacent tanks to keep them cool should a tank catch fire.

Only those parts of the adjacent tanks 'seen' from the burning tank need to be kept cool using the water, eg the shaded part shown in Figure 6.3. As the tanks are cylindrical, this accounts for half the tank surface which is exposed to the heat [62].

It is possible to calculate the heat flux received by the adjacent tank using equation 6.73. As only half the adjacent tank will receive the heat from the fire [62], the total amount of heat absorbed is by this half of the tank. Therefore, the effective area of the tank receiving the heat is:

Area =
$$\frac{\pi \text{ x diameter x height}}{2}$$
 6.76



Figure (6.3) Effect of radiant heat on adjacent tank.

Therefore the heat flux reaching the tank divided by the area calculated in equation 6.76 gives the amount of heat received by the adjacent tank per square metre.

If water is sprayed onto the adjacent tank, a heat balance over the tank enables the amount of water required to remove the heat from the surface of the tank to be calculated. The heat balance is given by

 $Q = m_1 Cp \Delta T \qquad \dots 6.77$

where	Q	=	heat received on the surface of the tank
	m_1	=	mass of water required to remove the heat
	Ср	=	specific heat capacity of the water
	ΔT	=	temperature rise of the water

But to apply the above equation, the following assumptions had to be made.

1. The water used is initially at 20 °C and rises to 90 °C when sprayed onto the adjacent tank wall. Therefore ΔT in equation 6.77 will be 70.

2. The specific heat capacity of the water varies between 4.183 kJ/kg K at 20 °C to 4.208 kJ/kg K at 90 °C but it is assumed to be constant at 4.191 kJ./kg K. Since 1 kg of water = 1 litre of water, the amount of water needed to keep the surface of the tank cool can be evaluated in litres/m²/sec or litres/m²/min as required.

A computer program including the preliminary estimation of the amount of water required was written to enable rapid calculation using the above method, this is shown in Appendix I. Clearly this can be adopted for different water temperatures. An allowance must also be made for any inefficiencies in water usage.

6.12 ANALYSIS OF THERMAL RADIATION RESULTS

The hazard ranges to people, buildings, and process equipment and the diameter, duration and height of a fireball which is growing and rising steadily with time, have been evaluated.

There is considerable disagreement between various authors regarding the appropriate values of numerical parameters to be used for predicting duration, equivalent spherical diameter, height, and hazard ranges from a fireball as shown in Figures 6.4 to 6.7. These Figures show the relationship between duration, diameter, height, and human hazard range with the mass of fuel. These graphs also show different values at each given mass of fuel for different author's parameters.

The results, as presented in Appendix G, are evaluated in terms of diameter, duration, height of fireball, and human, building and process equipment hazard ranges related to the flammable mass of hydrocarbon gases released. Calculations have been performed using various authors' parameters and Hot Shell-Cold Core, Isothermal and BLEVE models, as presented in Appendix G, to enable a quantitative assessment and comparison of all the methods. It must be accepted that the comparison is limited by the considerable number of assumptions mentioned earlier in Sections 6.4, 6.5 and 6.6. These assumptions may not always be true in real incidents. However, the mathematical models can be applied to compare various layout arrangements and the associated hazard ranges for humans, buildings and equipment can be predicted for different quantities of flammable fuels.

The results illustrated in Figures 6.8 to 6.15 according to different models show that the prediction of diameter and hazard

ranges according to the isothermal and BLEVE models have gross deviations due to several assumptions and empirical equations which may affect their accuracy. In the meantime the hot shell-cold core model gives unreasonable results for large scale scenarios, as presented in Appendix G. However, Lihou [56] suggested that this model had been used satisfactorily for small scale predictions.

The lack of accuracy of the above models may be due to a combination of the following:

- i It has been assumed that the diameter of a fireball is equivalent to the spherical diameter. However, the fireball shape changes with rise. At ignition, it is hemispherical then spherical and after lift off it takes several shapes [69].
- ii A fraction of fuel is assumed to be cracked to produce soot, the volume of which is neglected. However in real incidents luminous flames have high heat radiation rates and these represent the major hazard from fireballs and may cause fatal burns and secondary fires.
- iii Accurate calculation of the thermal radiation incident upon a receiving surface requires knowledge of the degree of attenuation of the radiation by the atmosphere between the flame and the receiver. This is a requirement common to both fireballs and pool fires [48].
- iv All the models are based upon small-scale experimental work in which the equivalent spherical diameter of the fireball increased at a constant rate. Therefore, resolution of the lack of accuracy of the above methods must await large scale experiments.
- v In the case of pool fires, the speed and extent of the spread across any surface will depend upon local topography and wind speed and direction.

The results for fireballs, which are presented in Figures 6.4 and 6.5 show that the four authors' correlations for fireball duration and diameter all give different results, and the deviations between them increase in proportion to the flammable quantity of fuel involved.

For flammable quantities less than 1 tonne, the four authors' correlations give essentially similar hazard ranges. On the contrary, when the flammable quantity is bigger than 1 tonne, the four correlations can clearly be distinguished from each other. In general the difference between the results are not too large, in particular the results for Marshall's and Robert's correlations. However, they are significant and cannot be neglected. In Figure 6.7, it is shown that the Roberts correlation generally provides the biggest human hazard range and the Gayle correlation usually gives the shortest values of human hazard range

In general, as would be expected, all four correlations provide the biggest ranges for human hazard and usually give the shortest values of process equipment ranges, as shown in Figure 6.8.

Also fireball diameter increases linearly with the duration of combustion in both isothermal and BLEVE models, as shown in Figures 6.9 and 6.10, because the assumption is that the fireball size in each model increases at a constant rate throughout combustion.

The results for hazard ranges based on the fireball surface temperature predicted using the thermodynamic equation give slightly smaller hazard ranges than those based on temperature predicted using the isothermal equation, as illustrated in Figure 6.11.

These differences between isothermal and thermodynamic temperature results are not significant, and can be neglected for large scale fireballs. Hence either method could be use to predict the



Figure 6.4 Variation of Fireball Duration with Mass of Fuel based on Different Authors' parameters.



Figure 6.5 Variation of Fireball Diameter with Mass of Fuel based on different Authors parameters.



Figure 6.6 Variation of Fireball Height with Mass of Fuel based on different Authors parameters.

surface temperature of a fireball theoretically. But in the case of the BLEVE model the results for hazard ranges predicted by thermodynamic calculation have larger deviations from the hazard ranges than those based on temperature predicted using the BLEVE equation, as illustrated in Figure 6.12.

In general, results from the two models increase in proportion to the increase of the mass of fuel. The difference between the results of the two models are significant and cannot be neglected, as shown in Figure 6.11 to 6.15. Figure 6.13 shows that the hazard ranges provided by the BLEVE model increase with decreasing number of carbon atoms (ie methane hazard range is the biggest while pentane hazard range is the shortest).

The difference between the four authors' correlations, Hot Shell-Cold Core, Isothermal and BLEVE models appear to be due to the different theoretical bases used for each method, although some common assumptions have been made in all the correlations and models, such as the assumption that the diameter of the fireball is the equivalent spherical diameter. Each method also contains several other assumptions that might cause considerable errors in the calculations. Moreover many factors involved in fireball combustion and growth remain unpredictable (eg wind velocity and direction, weather conditions, degree of confinement).

Therefore, the validity of these results remains unproven so average figures would be expected to give a more realistic answer. However, experience obtained from investigation of real incidents leads to the conclusion that maximum figures are to be preferred wherever possible.

The results for pool fires, which are presented in Appendix I



Figure 6.7 Variation of Human Hazard Range with Mass of Fuel based on different Authors parameters.



Figure 6.8 Variation of the Hazard Range with Mass of Fuel using Robert Correlation.



Figure 6-9 Relationship between Fireball Diameter and Fireball Duration according to Isothermal Model.



Figure 6.10 Relationship between Fireball Diameter and Fireball Duration according to BLEVE Model



Figure 6.11 Variation of Human Hazard Range with Mass of Fuel using Isothermal Model



Figure 6.12 Variation of Human Hazard Range with Mass of Fuel using BLEVE Model.





Variation of Fireball Diameter with Mass of Fuel using BLEVE Model for different Hydrocarbon gases.



Figure 6.14 Variation of Hazard Range with Mass of Fuel using Isothermal Model.



Figure 6.15 Variation of Hazard Range with Mass of Fuel using BLEVE Model.

and Figures 6.16 to 6.19, show that the method which is proposed in this study provides reasonable hazard ranges for humans, buildings and process equipment and the quantity of water rate required (litre/min.m²) to prevent a secondary fire.

Whilst it is not possible to deduce from these calculations the validity of the results obtained, a comparison with those published in the literature shows that they are in good agreement. However, many factors involved in a pool fire remain unpredictable (eg shape and size of the flame envelope, temperature and thermal emissive power of the flame). It is possible using the method proposed in this study to predict the spacing distance at which the values of L/D are appropriate for a pool fire for different masses of flammable liquid involved in the pool fire.

Figures 6.16 and 6.17 show that the spacing distances to humans, buildings and process equipment increase linearly with the increase of flammable mass and with increasing L/D value. Also the rates of water requirement to prevent secondary fires increases exponentially with the increase of L/D value and mass of flammable liquid, as illustrated in Figures 6.18 and 6.19. The water rate is consistent with government guideline figures.

The results are useful when designing the layout of plant and restricting access of persons to hazard areas. However, the effect of wind on the envelope of the flame is not taken into account in this study.

All the criteria and information presented so far, illustrate the high degree of uncertaintly associated with the prediction of the occurrence and the damage effects of fireballs and pool fires. It is therefore evident that the determination of spacing distance to ensure



Figure 6.16 Variation of Spacing Distance with L/D Ratio for 5m Diameter Pool Fire.



Figure 6.17 Variation of Spacing Distance with Diameter of Pool fire for 2.0 L/D Ratio.



Figure 6.18 Variation of Cooling Water Rate with L/D Ratio for 5 m Diameter Pool Fire.


Figure 6.19 Variation of Cooling Water Rate with Diameter of Pool fire for 2.0 L/D Ratio.

sufficient safety for the public and the environment cannot be performed with any precision with the present level of knowledge. However, there have been a number of quantitative assessments and several recommendations have been proposed. Most use an empirical form of approach. In all calculations a fundamental decision must be made as to an acceptable criterion for damage. This generally leads to the conclusion that maximum spacing distances are to be preferred.

Undoubtedly the prediction of safe and socially-acceptable spacing distances and determination of positioning of plant and/or storage facilities in relation to public buildings and constructions requires great skill and social responsibility. Therefore the best solution is the attainment of a high degree of safety without any considerable economic loss from spacing and siting policy, by optimising the process and reducing the risk to as low a value as possible.

CHAPTER SEVEN

PREDICTION OF ACCEPTABLE SPACING DISTANCES FOR PLANTS INVOLVING AN UNCONFINED VAPOUR CLOUD EXPLOSION HAZARD

The process industries handle a wide range of materials, a significant proportion of which can be regarded as hazardous. Many of the processes also require careful control to ensure safe operation. Failure of either equipment or operational standards could result in an explosion, for example, the explosive failure of a pressure vessel due to overpressurisation, or a vapour cloud explosion. Therefore, it is necessary to be able to assess the probable consequences of an explosion in terms of damage to the plant itself, to any neighbouring plants and the effects on operating personnel and the general public.

7.1 EXPLOSION

An explosion may be defined as a process whereby a pressure wave is generated in air by a rapid release of energy which causes a shock wave, moving away from the source [70]. An explosion can be produced by the detonation of a high explosive, the rapid burning of a flammable gas cloud, or the sudden failure of a pressure vessel containing high pressure gas. However, irrespective of the source of the released energy a pressure disturbance will be created in the surrounding air, and because of its compressibility, the air becomes heated locally with a corresponding increase in the velocity of sound leading to the front of disturbance becoming steeper as it passes through the air, with consequent increases in density and pressure. The resulting blast wave is largely responsible for the damage caused inside and outside the cloud as the shock wave propagates in the atmosphere beyond the cloud.

7.2 DAMAGE PREDICTIONS FROM OVERPRESSURES

It has been explained above how explosions produce a shock wave. This is a narrow band of high pressure travelling away from the explosion at sonic velocity. This overpressure applies impulsive forces to structures, the effect of which, in structural stress, may be up to twice that of a steadily applied pressure difference of the same magnitude.

Following the shock wave, structures will also be subjected to winds of high velocity. This creates a dynamic pressure which persists for much longer than the overpressure; especially after a UVCE the release of energy from which takes much longer than for a high explosion. Table 7.1 [41] shows the relative values of these pressures and the wind; they have been calculated from published relationships [71].

TABLE 7.1

PRESSURE PEAKS AND WIND VELOCITY FOLLOWING EXPLOSIONS

Overpressure (kN/m ²)	14	35	70	140
Dynamic pressure (kN/m ²)	0.7	4.9	14	56
Wind Velocity (km/h)	112	256	464	752

The response of structures to a dynamic pressure has been

extensively studied for military purposes. The results are generally expressed as the damage to be expected from a shock of specified overpressure.

Table 5.2 shows the likely damage from various peak overpressures. These have been derived from compiled data [42] on the damage resulting from TNT explosions. The estimation of generally acceptable peak pressure values for the safety of constructions is very complicated. It is even more difficult to predict an overpressure value at which people will not be harmed since the majority of injuries to persons outside the perimeter of the vapour cloud are due to building collapse or debris arising from buildings, in particular fragments from windows.

Several methods have been proposed for the calculation of acceptable spacing distances from the epicentre of a UVCE. The most important of them have been used in this study and are presented in the following paragraphs.

7.3 TNT EQUIVALENCE MODEL

The energy released by the explosive is the limiting factor to the magnitude of the blast wave. Therefore it is possible to quantify the potential energy of a conventional explosive, such as TNT, and compare this with that of LPG to arrive at some idea of the potential destructiveness of hydrocarbon explosions.

It has become customary to equate the damage caused by a UVCE to that caused by an equivalent mass of TNT. According to this method, the damage caused by an UVCE due to overpressure can normally be described as a function of a known amount of standard explosive (TNT) and the distance from the blast, ie the "TNT equivalence" system.

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The equivalent amount of TNT is usually calculated by the following steps [26]:

- Estimation of the quantity of vapour which has escaped (or in case of prediction is likely to escape).
- b. An assumption that all of this vapour burns explosively.
- c. Evaluation of the combustion energy released from this quantity of vapour.
- d. Division of this calculated energy release by the energy released upon the detonation of unit mass of TNT (about 1100 kcal/kg).
- e. Evaluation of the actual explosion yield using the explosion yield factor.

Most hydrocarbons display a maximum combustion energy release of around 10,000 to 12,000 kcal/kg compared with 1,100 kcal/kg for the usable energy in TNT. Therefore, on pure energy considerations alone, these hydrocarbons are ten times more potent than TNT. However, the rate of energy release is normally too low to create a shock wave. Therefore, an "efficiency factor" is normally used to determine the equivalent amount of TNT which corresponds to a given amount of hydrocarbon vapour. Various values have been suggested for this factor, eg 3% or 4% is recommended by the Second Report of the Advisory Committee on Major Hazards [72].

For design purposes, Brasie [73] recommends that yields of 2, 5 and 10% may be used to compare the sensitivity of various layout arrangements. Thus the overpressure, and, therefore the damage, can be predicted for different radial ranges. This has been done in the present work for the case of a 10 tonne propane UVCE and the results of calculations for different yields are given in Table 7.3. When the equivalent weight of TNT has been calculated the overpressure and, therefore, the damage can be calculated for a given radius. There are a variety of methods for calculating this; the most common are summarised in the following paragraphs.

7.3.1 Jarrett's Equation

There are a variety of numerical equations, expressing the relation between the TNT equivalent amount and the distance from the epicentre of the UVCE, but the most widely used is the equation proposed by Jarrett [74]. The following equation is used by the UK. Explosives Storage and Transport Committee to assess the range at which different categories of damage may be expected [74]

$$R = \frac{k W^{1/3}}{\left[1 + (6996/w)^2\right]^{1/6}} \qquad \dots 7.1$$

where R = distance from the explosion in feet,

W = weight of TNT in pounds,

K = constant appropriate to various degrees of damage; set out below:

K = 9.6	-	almost complete demolition
K = 13.9	-	50-70% external brickwork destroyed
K = 24	-	houses uninhabitable - partial or total
		collapse of house walls and roofs
K = 70	-	minor structural damage to buildings.
K = 140	-	remaining inhabitable after repair,
		some damage to ceiling and tiling,
		more than 10 percent window glass
		broken.

The constant K, involved in Jarrett's equation takes a different

value for every characteristic value of overpressure according to Figure 7.1 [29].

The quantitative assessment of spacing distance according to the TNT Equivalence Model using Jarrett's equation including the assumption of explosion yield factor of 2%, shows that one tonne of TNT is equivalent to 4.95 tonnes of propane. The constant K, involved in equation 7.1 takes a different value for every characteristic value of overpressure as mentioned earlier. A computer program was written to enable rapid calculations using this method; this is given in Appendix J, and the results of typical calculations are given in Table 7.2.

7.3.2 Hopkinson's Law

Another method to estimate the blast effect is based on the common scaling law of Hopkinson [3] which states that:

"Identical blast waves are produced at identical scaled distances when two explosive charges of the same geometry and explosive material, but differing in size, explode in the same atmosphere".

It is important that only the scale is varied, and it is found that the overpressure at a given radius is proportional to the cube root of the energy or mass content of the charge [3].

This law is the basis of a curve relating scaled ranges $(distance/mass^{1/3})$ to overpressure. The curve of scaled distance versus overpressure which is represented in Figure 7.2 was derived from a large series of trials carried out with surface bursts of hemispherical TNT charges [75]. It should be noted that this curve is only appropriate for conditions where the terrain and atmospheric conditions have no significant effect.



Figure 7.1 The values of constant K (Jarrett's equation) at different overpressures

TABLE 7.2

Overpressure Amount released tonnes H/C equivalent 5 10 20 30 40 50 kN/m² K Values spacing distance (m) 69 3 7.9 12.1 17.4 20.7 23.2 25.2 42 13.9 37 56.4 80.7 96.1 107.5 116.8 34.5 16.3 43.4 66.1 94.6 112.7 126.1 136.9 20.7 19.8 52.7 80.3 114.9 136.9 153.2 166.4 15.9 22.8 60.7 92.5 132.4 157.6 176.4 191.6 10.35 37 98.5 150.1 214.8 255.9 286.3 310.9 6.9 51 137.1 208.9 299 356.2 398.5 432.8 4.8 61 163.4 249.1 356.5 424.6 475.1 516.0 2.76 72 191.7 292.1 418.0 497.9 557.2 605.1 2 99 263.5 401.7 574.8 684.7 766.1 832.0

OVERPRESSURE VERSUS SPACING DISTANCE USING TNT EQUIVALENCE MODEL FOR DIFFERENT TONNAGES OF PROPANE

TABLE 7.3

OVERPRESSURE VERSUS SPACING DISTANCE USING TNT EQUIVALENCE MODEL OF 10 TONNES PROPANE FOR DIFFERENT YIELD FACTORS

Overpressure kN/m ²	Spacing distance (m) for different yield factors				
	2%	5%	10%		
69	12.2	19.2	25.2		
42	56.4	89.1	116.8		
34.5	66.1	104.5	136.9		
20.7	80.3	126.9	166.4		
15.9	92.5	146.2	191.6		
10.35	150.1	237.2	310.9		
6.9	208.9	330.2	432.8		
4.8	249.1	393.6	516.0		
2.76	292.2	461.6	605.1		
2	401.7	634.7	832.0		

In this method values of an explosion efficiency (the ratio of the energy in the blast wave to the energy theoretically available from the heat of combustion, usually expressed as a percentage) of 2, 5 and 10% has been used as recommended by Brasie.

For any overpressure, the scaled distance can be read from Figure 7.2 [70]. The actual distance can be obtained when the equivalent amount of TNT is known.

The quantitative assessment of spacing distance according to TNT Equivalent using Hopkinson's law including the assumption of 2% explosion yield factor shows that 2 tonnes of TNT are equivalent to 10 tonnes of propane in the cloud.

The overpressures developed at various distances can be obtained from Figure 7.2. A computer program was written to enable rapid calculations using this method; this is given in Appendix K and the results of typical calculations with different yield factors are given in Table 7.4.

7.4 GUGAN'S MODEL

This model is based on the concept that percussive combustion initially produces blast waves. These waves are unstable and become shock waves after t_S (seconds) at a radius R_S (metres) where the overpressure will be a maximum. The shock pressure P_S (kN/m²) decreases with range R (metres) according to the semi-empirical equation 7.2 below [41].

$$P_{s} = 37.86 \left[\frac{Eo^{1/3}}{R} \right]^{1.5} \dots 7.2$$

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TABLE 7.4

OVERPRESSURE VERSUS SPACING DISTANCE USING TNT EQUIVALENT MODEL (HOPKINSON'S LAW) FOR DIFFERENT YIELD FACTORS OF 10 TONNES OF PROPANE

Overpressure	Spacing distance (m) for different yield factors				
KIN/III-2	2% yield	5% yield	10% yield		
69	50.4	68.4	86.2		
42	69.3	94.0	118.5		
34.5	75.6	102.6	129.2		
20.7	107.1	145.3	183.1		
15.9	124.7	169.2	213.2		
10.35	176.3	239.3	301.5		
6.9	226.7	307.7	387.7		
4.8	302.3	410.3	516.9		
2.76	440.9	598.3	753.8		
2	579.4	786.4	990.7		

where Eo is the total combustion energy released (MJ)

$$t_s = 1.5 \times 10^{5}/P_c$$
 7.3

$$R_s = 4.95 \times 10^7 / P_c$$
 7.4

In equations 7.3 and 7.4, P_c° is the rate of pressure rise (N/m^2s) at the centre of the initial explosion.

Gugan [26] gives a relationship whereby Pc° may be calculated,

.. 7.6

$$P_c = 1.75 \times 10^6 \text{ Eo/a}^3 t_0^4 \dots 7.5$$

and
$$t_0 = Dc/a$$
 ...

where a = sonic velocity in air; 330 m/s

 t_0 = the duration of explosive release of energy, (s)

Dc = the initial hemispherical diameter of the cloud at NTP, (m)

The quantitative assessment of spacing distance according to Gugan's model is exemplified by the following example:

Consider the explosion of 10 tonnes of propane. Assuming a stoichiometric propane-air mixture of molar air: propane ratio of 23.8 and molecular weight of 44 gives the hemispherical diameter of the cloud as below

volume of cloud at NTP =
$$\frac{(1 + 23.8) \times 22.4 \times 10 \times 10^3}{44}$$
$$= 126 \times 10^3 \text{ m}^3$$

therefore, hemispherical diameter of cloud = 78.4 m Energy release Eo = $46.45 \times 10 \times 10^3 = 464.5 \times 10^3 \text{ MJ}$

> from equation 7.5, $\mathbf{P_c} = \frac{1.75 \times 464.5 \times 10^3 \times 330}{(78.4)^4} = 7.12 \text{ MN/m}^2 \text{ s}$ from equation 7.4, $\mathbf{R_s} = \frac{4.95 \times 10^7}{7.12 \times 10^6} = 6.95 \text{ m}$ from equation 7.6, $\mathbf{t_0} = \frac{78.4}{330} = 0.238 \text{ sec}$

Thus Gugan's model indicates a duration of the explosive release of energy of 0.238 sec and extensive damage about 6.95 m from the epicentre, where the blast transforms into a shock wave.

Using equation 7.2, with Eo = 464.5 x 10³, the range at which the overpressure P_s kN/m² occurs is given in equation 7.2.

Rearranging equation 7.2 gives

 $R = \left[\frac{37.86 \text{ Eo}^{1/2}}{P_{\text{S}}} \right]^{2/3} \dots 7.7$

A computer program was written to enable rapid calculations using this method; this is given in Appendix L and the results of typical calculations are given in Table 7.5.

7.5 CLANCEY'S MODEL

Clancey's model is based upon the concept of a large quantity of vapour being released instantaneously to form a cloud which develops into a hemisphere in which the vapour concentration assumes a Gaussian distribution of the form [76]:

Overpressure (kN/m ²)	Radial range (m)
<i>(</i>)	
69	51.9
42	72.3
34.5	82.4
20.7	116
15.9	138
10.35	184
6.9	241
4.8	307
2.76	444
2	550

TABLE 7.5RANGES OF OVERPRESSURES PREDICTED BY GUGAN'S MODELFOR UVCE OF 10 TONNES PROPANE

$$C_{(r)} = \frac{2W}{(2\pi)^{3/2} \sigma^3} \exp \left[-\frac{r^2}{2\sigma^2} \right] \qquad \dots 7.8$$

The first term is the concentration at the centre, Co, so that

$$C_{(r)} = \operatorname{Co} \exp\left[\frac{-r^2}{2\sigma^2}\right] \qquad \dots 7.9$$

and

$$Co = \frac{2W}{(2\pi)^{3/2} \sigma^3} \qquad \dots 7.10$$

where

- W = total quantity of vapour in cloud (kg)
 - $C_{(r)}$ = vapour concentration at distance r from centre (kg/m³)
 - Co = vapour concentration at centre of cloud (kg/m³)r = radius m

 σ = standard deviation of concentration distribution (m)

If C_2 and C_1 are the concentrations at the upper and lower flammable limits it can be shown that the maximum amount of vapour in the cloud occurs when the concentration at the centre is C_2 or when

$$Co = C_2$$
 7.11

substituting from equation 7.10

$$C_2 = \frac{2W}{(2\pi)^{3/2} \sigma_m^3} \qquad \dots \ 7.12$$

Rearranging equation 7.12 gives,

$$\sigma_{\rm m} = \left[\frac{2W}{(2\pi)^{3/2} C_2}\right]^{1/3} = 0.5 \left[\frac{W}{C_2}\right]^{1/3} \dots 7.13$$

Having obtained σ_m the standard deviation, when the cloud presents the maximum hazard, it is possible to find the radius r_1 , at which the concentration is at the lower flammable limit,

$$C_1 = Co \exp\left[\frac{-r_1^2}{2\sigma_m^2}\right]$$
 7.14

Rearranging equation 7.14 gives,

$$r_{1} = \left[2\sigma_{m}^{2} \ln\left(\frac{C_{2}}{C_{1}}\right) \right]^{1/2} \dots 7.15$$

$$r_{1} = \left[\sigma_{m}^{2} \ln\left(\frac{C_{2}}{C_{1}}\right) \right]^{1/2}$$

$$= 0.71 \left[\frac{W}{C_{2}} \right]^{1/3} \left[\ln\left(\frac{C_{2}}{C_{1}}\right) \right]^{1/2} \dots 7.16$$

This is in the absence of combustion. However, combustion will cause an eight-fold increase in volume. Thus, the radius Re of the edge of the cloud, when it is reached by the combustion wave is $2r_1$,

therefore, Re = 1.42
$$\left[\frac{W}{C_2} \right]^{1/3} \left[\ln \left(\frac{C_2}{C_1} \right) \right]^{1/2}$$
 7.17

Clancey [76] suggests that the maximum pressure of 69 kN/m^2 (10 psi) is produced at ranges of Re, decaying at longer ranges.

The quantitative assessment of spacing distance according to Clancey's model is exemplified by the folloing example:

Consider the catastrophic failure of a tank containing 28 tonnes of propane at 20 °C (293 °K). The atmospheric boiling point of propane is 42 °C (231K). Mean specific heat of liquid propane over the temperature range 293 °K to 231 °K is 2.41 kJ/kg.°K, and latent heat at 231 °K is 410 kJ/kg.

then mass of propane flashing W = $\frac{28 \times 2.41 \times (293-231)}{410}$ = 10 tonnes

Flammable limits for propane in air, $f_1 = 0.024$ and $f_2 = 0.095$.

Converting these into flammable concentration at 293 °K

 $C_1 = \frac{0.024 \text{ x } 44 \text{ x } 273}{22.4 \text{ x } 293} = 0.04 \text{ kg/m}^3$

 $C_2 = \frac{0.095 \text{ x } 44 \text{ x } 273}{22.4 \text{ x } 293} = 0.17 \text{ kg/m}^3$

From equation 7.13, $\sigma_{\rm m} = 0.5 \left[\frac{10,000}{0.17} \right]^{1/3} = 19.23 \text{ m}$

From equation 7.16,
$$r_1 = 0.71 \begin{bmatrix} 10,000 \\ 0.17 \end{bmatrix} {}^{-1/3} \begin{bmatrix} 0.17 \\ 0.04 \end{bmatrix} {}^{1/2}$$

= 32.03 m

From equation 7.17, $Re = 2r_1$

 $= 2 \times 32.03 = 64.06 \text{ m}.$

The overpressure predicted by Clancey's model beyond Re can be calculated as follows:-

Figure 7.3 was redrawn by Lihou [41] in SI units from a similar chart presented by Brasie and Simpson [77]. It shows the overpressure as a function of scaled distance; that is the radial distance (m) from the epicentre of the explosion divided by the cube root of the heat of combustion (MJ) of the total mass of flammable in the cloud. The actual range of any overpressure P_S is Re ($r_S/0.48$), where r_S is the scaled range corresponding to P_S on Figure 7.3.

A computer program was written to enable rapid calculations using this method; this is given in Appendix M and the results of typical calculations are given in Table 7.6.

7.6 ANALYSIS OF BLAST WAVE RESULTS

Overpressure is one of the key parameters in the analysis of explosions. There has been extensive research into the relationship between overpressure and distance, chiefly for military purposes. For



Figure 7.3 Peak pressure distribution from given combustion energy assuming 2% yield.

TABLE 7.6

Overpressure, kN/m ²	Radial range, m
69	64.06
42	93.4
34.5	100.0
20.7	130.8
15.9	149.5
10.35	204.2
6.9	280.3
4.8	373.7
2.76	747.4
2	1281.2

RANGES OF OVERPRESSURE PREDICTED BY CLANCEY'S MODEL FOR A UVCE INVOLVING 10 TONNES PROPANE

example, Figure 7.1-7.3 show the relationship between scaled range and overpressure. These curves show differing values of overpressure at a given scaled distance. There is a great deal of variance in results and the graphs represent "best fit" relationships [38].

The results, as tabulated in Table 7.2 -7.6, are derived in terms of spacing distance (metres) related to the flammable quantity released and to several characteristic values of overpressure which correspond to different levels of damage effect, as shown in Table 5.2. Calculations have been performed, using each of the above methods, and are presented in Appendices J, K, L and M, to enable a quantitative assessment and comparison of all the methods. It must be accepted that the comparison is limited by the considerable numbers of assumptions mentioned earlier. An important assumption for example is that the explosion yield factor does not exceed 2% in Gugan's and Clancey's models. This may not always be true in real incidents in which the yield factor has been over 10% or even below 1%. Nevertheless, these mathematical models with the explosion yield factors of 2%, 5% asnd 10% may be used to compare various layout arrangements.

The results illustrated in Figures 7.4 -7.13 show that the calculated distances according to TNT equivalent methods, Gugan and Clancey have gross deviations due to several assumptions and empirical equations which may effect their accuracy.

The accuracy of each of the above methods may be subject to the following limitations,

i the inherent accuracy is uncertain as demonstrated by the data from real incidents [26] which are shown in Table 7.7, in which topography, atmospheric conditions, etc. cause considerable variations

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TABLE 7.7

DATA FOR SOME PRINCIPAL UVCE's

Date	Place	Flammable	Source	Quantity (kg)	TNT Equiv. (kg)
23.8.1921	Hull England	Hydrogen	Dirigible ZR-2	6,900	450 kg
Yield measu 8.4 km awa	ured: 0.25%. W y. Tremors felt	/indows broken 80 km. Fatalit	(0.5 psi) wit y 1.	hin 3.2 km :	area. Blast heard
16.1.1966	Caltex W. Germany	Methane	Process	500	1000-1200
Yield measu a low lying glass breaka	ured: 18-36%. I dispersion. B ge at 400 m, mi	Liquid methane last damage to inor breakage at	vented at hig plant not pa 1200 m. Fat	h level -substructure of the second s	sequently formed evere. Extensive
9.12.1970	Franklin Co. Port Hudson USA	Propane	Pipeline	70,000	50,000

Yield measured: 7.5%. The overpressure was 1 psi at 585 m and 0.5 psi at 1,100 m. No fatalities.

1974 England Ethylene Process 910-2700 7+

Leak from high pressure piping caused spill lasting about 30 min. before ignition. Windows broken (0.3 psi) in 180 m radius. Roof and ceiling in 4 buildings damaged. No fatalities.

1.6.1974 Flixborough Cyclohexane Process 36,000 18,000 England

Yield measured: 5%. Poorly installed 500 cm temporary pipe failed. Ignition in 25-35 seconds created large overpressures. Fatalities: 28.

20.2.1977	Dallas	Isobutane	Rail	68.200	1600
	Texas		tank car		

Yield measured: 0.25%. Tank car punctured during derailment. Delay to ignition about 2.5 minutes. The open-air explosion was extremely violent and the shock was felt for miles around. Locomotives would appear to have been exposed to about 3.5 kPa at a distance of about 300 m from punctured tank car. No fatalities.

in the overpressure profiles.

ii it may be argued that not only overpressure but also duration is significant when damage potential is considered. Thus the overpressure required to produce a given level of damage may vary in cases where the duration is different [38]. However, in this study, the duration is fixed for a given level of overpressure. In principle this may not be true between one explosion and another.

iii examination of damage at UVCE sites gives two distinct regions, the region enveloped by the cloud and that beyond the periphery of the cloud. This difference should lead to the development of sophisticated mathematical models to explain the blast wave phenomena in UVCE's [3].

iv the statistics on flammable material involved show that they have a very wide range of physical, chemical and combustion properties. The only common factors would appear to be that the materials must be flammable and must be readily able to form mixtures with air within the appropriate flammable range [26].

The results, which are presented in Table 7.2-7.6, Figures 7.4-7.13 and Appendices J, K, L and M show that the four methods always provide spacing distances of the same order for almost every overpressure value, and the difference between results of the four methods increases in proportion to the increase of the flammable quantity (expressed in tonnes of hydrocarbon equivalent).

For flammable quantities less than 5 tonnes, the four methods give similar spacing distances. On the contrary, when the flammable quantity is bigger than 5 tonnes, the four methods can clearly be distinguished from each other. In general the differences between the results are not too large. However, they are significant and cannot be

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neglected.

In Figures 7.10-7.13, the explosion yield factor was assumed to be constant at 2% for every method and the results demonstrate that the Clancey model generally provides the biggest spacing distances and the Jarrett equation usually gives the smallest values of spacing distance.

The 2% yield factor is often not conformed with in practice [26, 38]. In this work, Jarrett's equation and Hopkinson's Law were used to obtain results with 5% and 10% yield factors as well as with 2%, as are shown in Figure 7.5 and 7.7. These results show that as expected the spacing distance increases with increasing yield factor.

The differences between the 4 methods appear to be due to the different theoretical bases used for each method, although some common assumptions have been made in all the methods, eg, explosion yield factors. Also, each method contains several other assumptions that might cause considerable errors in the calculations.

In conclusion, it is possible using these methods to predict the spacing distance at which characteristic values of overpressure are probable from a UVCE. As many factors involved in the potential cloud explosion remain unpredictable (eg wind velocity and direction, weather conditions, degree of confinement, spill rate and operator response), the explosion yield factor is taken at a more conservative value than previously ie 10%. Because all criteria and information illustrate the high degree of uncertainty associated with the prediction of the occurence and the damage effects of UVCE and as none of the four methods of calculation used is more or less credible than the others based on experience from real incidents, this leads to the conclusion that maximum figures are to be preferred.

The best solution is always the attainment of a high degree of safety without any significant economic loss by appropriate spacing and siting policy, in optimizing the process and reducing the risk to as low value as possible. However, the incidents that have arisen in recent years suggest that more importance ought to be given to safety considerations than to economic ones in designing appropriate spacing distances.



Figure 7.4

Variation of spacing distance with hydrocarbon equivalent for 10% explosion yield factor at different overpressures according to TNT equivalent model (Jarrett equation) 209



Figure 7.5 Variation of spacing distance with hydrocarbon equivalent for 6.9 kN/m² overpressure at different explosion yield factors according to TNT equivalent model (Jarrett equation) Explosion yield factor

- ▲ 2%
- o 5% 210
- × 10%



Figure 7.6 Variation of spacing distance with hydrocarbon equivalent for 10% explosion yield factor at different overpressures according to TNT equivalent model (Hopkinson law).



Figure 7.7 Variation of spacing distance with hydrocarbon equivalent for 6.9 kN/m² overpressure at different explosion yield factors according to TNT equivalent model (Hopkinson law)

Explosion yield factor

- x 2%
- 0 5%
- ▲ 10 % ²¹²



Figure 7.8 Variation of spacing distance with hydrocarbon mass at different overpressures according to Gugan's equation.



Figure 7.9 Variation of spacing distance with hydrocarbon mass at different overpressures according to Clancey's equation.



Figure 7.10 Variation of Spacing Distance to 4.8 kN/m² Overpressure based on Different UVCE Calculation Methods.



ure 7.11 Variation of Spacing Distance to 6.9 kN/m² Overpressure based on Different UVCE Calculation Methods.


Figure 7.12 Variation of Spacing Distance to 10.35 kN/m² Overpressure based on Different UVCE Calculation Methods.



Figure 7.13 Comparison between the Different UVCE Calculation Methods for 30 tonnes propane at 2% Yield Factor.

CHAPTER EIGHT

SAFETY MEASURES IN LPG INSTALLATIONS

This chapter is concerned with the control of major LPG hazards, the prevention of their realization, and the limitation of their consequences. It has to be recognised that if a hazard exists, the likelihood of its realization can never be reduced to zero. If it cannot be eliminated entirely, control must aim to reduce the individual or social risk associated with the hazard to a level which is tolerable, acceptable, or justifiable to society as a whole.

Quantification methods are a vital part of the process of assessment of LPG hazards and risks. Some of these techniques have been discussed in detail in Chapters 6 and 7.

Therefore, the possible occurrence of a major LPG fire or explosion can be minimized by sound engineering, plant design and layout, good operating practice and proper training of personnel in both routine operations and actions to be taken in an emergency.

8.1 DESIGN CONSIDERATIONS FOR LPG INSTALLATIONS

8.1.1 Bulk LPG Storage Installations

The following are important factors in plant design (4, 45) :

- a. The maximum weekly offtake, in conjunction with the type and rate of supply, as a guide to determine storage capacity,
- b. The maximum instantaneous and average hourly offtake to determine the size of vaporizer or the capacity of LPG air-plant,

- c. The possibility of future extensions to the plant capacity,
- d. The need for standby facilities,
- e. Access facilities for road delivery vehicles including road widths and suitability of roadways for axle loadings,
- f. Minimum winter temperature, to determine natural vaporization rates from LPG vessels,
- g. The pattern of operation required, i.e continuous or intermittent,
- h. No bund walls are permitted around pressure vessels,
- The design pressure for the top of the vessel should be 110% of maximum operating pressure,
- j. The design pressure used for the bottom of the vessel should be that for the top of the vessel plus the static head of the maximum contents of the vessel,
- Only one product line should be provided at the bottom of the vessel, to be used for filling, discharge and draining,
- A manually-operated fire-safe valve and a remote controlled fail
 -safe emergency valve should be provided at the manifold side,
- m. Separate safety/relief valves should be provided to protect against overpressure due to abnormal operational conditions,
- n. Overfill protection, should also be provided, in addition to relief valves, in the form of instrumentation. This is particularly important for very high thermal expansion material such as LPG.

In addition to the above requirements for the design of LPG installations, guidance is available on the location, spacing, layout etc, in various LPG Codes of Safe Practice, eg, produced in the UK by the Institute of Petroleum [78] LPG ITA [9], the Home Office and the

HSE [11].

8.1.2 Pipework, Valves, Pipe Fittings and LPG Pumps

All materials used, including non-metallic parts for valves, seals, gaskets, diaphragms, etc, must be resistant to the action of LPG under their operating conditions.

i. Fixed Pipework:

Only steel piping may be used for liquid line service, although vapour lines may either be of steel or solid drawn copper. Pipework for liquid and vapour lines adjacent to storage vessels must be flexible enough to accommodate any settling of the storage tanks or other equipment, any thermal expansion or contraction, and any other stresses that may occur in the pipework system. Flexible hoses used for this purpose should not be greater than 230 mm (9 in) long of 25 mm (1 in) bore, and should be used only on installations of up to 2 tonnes nominal LPG capacity. Flexible connections should have a minimum bursting pressure of four times the maximum pressure they will bear in service. Corrosion-resistant braiding should be used [4, 79].

ii Pipe Joints:

Steel joints on pipework with a bore of 50 mm (2 in) or more should be welded or flanged. Joints on pipes with a bore of less than 50 mm (2 in) may be welded, flanged or screwed, although liquid lines should preferably be welded or flanged. Copper pipework up to a maximum size of 12.5 mm (1/2 in) should only be used for vapour lines. All joints should be made using compression-type fittings designed to withstand four times the maximum pressure they will carry

in service. Alternatively they may be silver soldered or brazed using material with a melting point of about 540 °C [4].

Flexible connections and hoses must be resistant to the action of LPG in accordance with the specification of the LPGITA (UK) [9]. Care should be taken to avoid undue stressing when installing flexible hoses. Where hoses are exposed to the possibility of mechanical damage or wear, they should be armoured; where they are likely to be subjected to high temperature, the advice of the manufacturer should be sought. All jointing compounds should be resistant to the action of LPG and should conform to the requirements of BS 5292 or an equivalent specification.

iii Valves and Fittings:

Valves should preferably be made from steel, forged steel, or brass which has been produced by extrusion or rolling. Cast iron valves should not be used. Ball valves are particularly suitable for liquids and retain their seal if involved in fire, ie they have 'fire-safe' characteristics. They should however conform to BS 5351 'Steel Ball-Valves for the Petroleum Industry'.

All steel flanges and flanged fittings used should conform to BS 1560 or equivalent standard. If fittings are 50 mm (2 in) diameter or above, the flanges should be welded, although screwed flanges may be allowed [4].

Steel butt welded fittings, steel socket welded and screwed fittings, and bolting should conform with BS 1965, BS 3799 and BS 4882 respectively or an equivalent specification.

iv LPG Pumps [4]:

Many LPG installations require liquid to be pumped between the storage vessel and the vaporiser. Pumps for this purpose must be suitable for pumping LPG which has no lubrication properties and a low viscosity. They must also be continuously rated and, if electrically-driven, require a flameproof motor. If compressed air is available, air operated reciprocating pumps are available which are very suitable for use in hazardous areas.

Liquid pumps should be provided with a means of pressure relief, discharging back to the storage vessel or to the pump inlet to protect the pump against overload. Pressure relieving devices fitted to pumps are often not designed to maintain a constant delivery pressure, but discharge when a pre-set differential pressure is reached. The inlet pressure at the pump will, however, vary with storage pressure which will in turn depend upon the ambient temperature, thus causing the relief pressure to vary. To maintain a constant outlet pressure, a separate pressure sustaining valve should be used.

Use should be made of adequate suction pipe siting and adequate static head to ensure that a satisfactory NPSH (ie nett positive suction head) is achieved under all conditions. Pumps should always be installed in accordance with manufacturer's instructions, taking special care that the piping places no strain on the pump casing.

Before any pipe run is tested, it must be blown through with nitrogen to purge oxygen from the system and to remove any remaining foreign matter which could foul the pressure regulator or cause trouble with the sealing of valves in the line. After installation, the pipework must be pressure tested to ensure that it is gas tight and free from leaks. A pressure gauge should be fitted in the line and

pressure testing carried out with nitrogen gas to a maximum of 1.5 times the maximum operating pressure of 6.90 bars (100 lb_f/in^2 gauge) whichever is the greater. Joints must always be tested using detergent solution. All pressure testing on underground pipework should be carried out before the pipework is wrapped.

8.2 FIRE PREVENTION, PROTECTION AND CONTROL

Fire prevention and protection measures for process plants, storage tanks and other facilities need to be integrated during design to the degree necessary to prevent a fire occuring, to detect any fire which does occur, and to restrict its spread.

Within the U.K. the safety requirements for the storage and transport of LPG, both in bulk and in cylinders, in factories and other premises to which the Factories Act 1961 applies, is covered by the Highly Flammable Liquids and Liquefied Petroleum Gases Regulations, 1972. It is important that the requirements are not infringed by site modifications or any extensions to existing storage. The Codes of Practice published by The Liquefied petroleum Gas Industry Technical Association - U.K. (LPGITA) [8] should be consulted and their recommendations taken as the minimum standards applicable.

Generally, the design and engineering of LPG process plants requires [45, 8]:

- a. The correct selection and use of materials and methods in the plant construction.
- Establishment of correct process conditions (eg maximum and minimum temperatures and pressures).
- c Incorporation of process control and safe guarding systems, eg protection against over pressure (safety relief valves),

instrumentation providing automatic alarms and trips, systems for safe emergency disposal of excess hydrocarbon gases.

- d. Attention to the layout and spacing of plant and facilities.
- e. Correct choice of electrical equipment for use in hazardous areas by defining such areas in accordance with Area Classification Codes, eg in the U.K. - BS 5345 and IP Electrical Safety Code.
- f. Detection of leakage, eg gas detection on LPG pump seals; or of fire, eg hot oil pumps, or of overheating, eg pump and motor bearings
- g. provision of fire protection to load-bearing elements of steel structures.
- Provision of adequate drainage facilities for the removal of firefighting water.
- i. Provision of fire-fighting water systems, fire detection and alarm systems, fire-fighting fixed and mobile systems, etc.

To minimise the risk of fire, and its spread there are two major requirements for safe operation [45].

- i. The development, and effective use, of comprehensive operating procedures and the training of all personnel in the various aspects of normal and emergency situations; and
- ii. Ensuring that the installed systems continue to perform to the standards of safety specified by the designers.

With regard to (i), Permit-to-Work procedures are an essential part of such training and both operation and engineering personnel must be instructed with regard to the scope of their responsibilities

and the authority levels for safe work. Personnel must be trained in the normal and abnormal conditions of operations and their procedures, and in emergency plans covering fire and explosion, spillage and gas escape.

With regard to (ii), to ensure continuity of safety standards, it is essential that inspection, tests and checks are carried out at regular intervals and that the engineering integrity of the systems is correctly maintained. The frequency of periodic inspections and tests will vary but agreed frequencies and procedures should be laid down as a written Code of Practice, eg an important instrument trip/alarm function may require a monthly check but others may be quarterly.

8.3 FIRE-FIGHTING SYSTEMS

As a matter of company policy, the design and engineering of fire fighting systems should meet national or local requirements. The requirements are based on the assumption that there will be only one major fire at a time. This does not exclude the possiblity of more than one major fire on any site simultaneously, since this has occurred infrequently in the past, but is a reasonable economic judgement.

8.3.1 Fire Fighting Water Systems

a. An open source is preferred for water for fire-fighting.

- A ring main system should be provided which surrounds process plants, loading areas for road and rail, storage facilities, workshops, etc. The capacity of the ring main system must be large enough to provide for the largest single water demand.
- c. Block valves should be provided in the ring main to enable isolation of parts in case of damage and for repair.

- d. Permanent hydrants should be provided and must be readily accessible from the LPG plant roads.
- e. The capacity of the ring main system must not be less than 720, m³/hr (or 2600 gal./min). In practice LPG installations have been provided with capacities ranging up to 10,500 gal./min depending on their requirements for water quantities for making foam and for exposure protection of adjacent facilities and installations.
- f. The water distribution system should be maintained liquid full and permanently pressurised to 3 bar by a small pressurising pump or by connection to the plant cooling water system [45, 8].

8.3.2 Pumps for Fire-fighting Water

At least two identical submerged pumps are required, one electrical motor-driven and one diesel-driven as a spare, taking suction from open water. These pumps must be capable of starting against an open discharge so the power drives must be rated accordingly. The pump drives should be provided with automatic starting devices. Manual start/stop should be provided from the control room or fire station and of course at the pumps. To maintain a system pressure of 10 bar at the furthest hydrant under full flow conditions, the pump discharge pressure could be 15 bar. A pressure indicator controller should be installed at the common discharge to enable adjustment to the pressure required when fire-fighting water is taken from close to the pump area. This makes it safe for personnel handling fire hoses [45].

8.3.3 Fixed Automatic Water Spray Systems for LPG Spheres and Horizontal Vessels

LPG pressure storage should be provided with fixed automatic water spray facilities to protect the sphere or horizontal vessel against radiation during a fire.

None-clogging type open sprinklers are to be fitted over the entire surface to ensure an even distribution of water rgardless of wind direction or wind force. The design criteria includes [45, 8, 80]

- i Piping supports must not be welded to the vessel.
- ii Heavy quality galvanised piping to BS 1387 is normally to be specified.
- iii The water spray system on each vessel is to be divided into two separate sections; each section being opposite to an adjacent storage vessel.
- iv The supply of water should be possible from two sides, preferably opposite each other. Strainers at the ring main shall be fitted between block valves.
- v The block valves and strainers should be placed at least 30 m from the periphery of the vessel.
- vi The system downstream of the valves shall slope upwards for self-draining via an 8 mm diameter hole near the valves.
- vii The capacity of the water spray systems fitted to the LPG storage is to be 10 litre/min/m² of surface to be protected.
- viii The automatic sprinkler systems must be installed over the pumps. Also, the sprinkler heads are to be located to give effective cover of the pump area and a surrounding area of 1.5 m width.

The automatic fire detector installed should consist of a plastic tube kept under instrument air pressure. In the event of a fire this plastic tubing will melt at approximately 90 °C causing the air pressure to fall, which will actuate an alarm in the control room and start the fire-water pump [45, 80].

8.4 SAFETY OF PERSONNEL

There are many hazards in LPG installations, especially when a fire or explosion occurs, and the following are rules which should be observed by safety officers and personnel in the interests of general safety [79].

- a. Any crew of fire-fighting personnel stationed in a hazardous position should have an observer attached to it whose task is to warn the men when danger is imminent, and a prearranged signal for this purpose should be agreed upon. This observer should preferably be a safety officer whose technical knowledge is such that he will be intimate with the main dangers, and a fire brigade officer should act as a liaison officer.
- Men should not work inside a bund wall if there is any leakage of LPG, or if the work can be done adequately from outside.
 When it is necessary to work within a bund, points of egress should be agreed upon.
- c. Where necessary, men should be protected from radiated heat by improvised shields, and reliefs, ie relief of personnel, should be arranged at frequent intervals. This protection should be given even if the heat does not make it immediately necessary, as a change of wind may easily increase the heat.
- d. Men should not be allowed on tank roofs unless absolutely

necessary, and never on empty tanks which are on fire. If men have to go on to a tank roof (eg for a vent fire), they should take lines with them in case they are cut-off from the ladder.

- e Breathing apparatus should be worn when necessary to protect against toxic or asphyxiating vapours or gases.
- f Personnel should not be allowed to smoke. It may not be dangerous in the vicinity of the fire, yet highly so if the men move within the range of vapours elsewhere in the LPG installation.

g Personnel should not approach the fire or gas leak from upwind.

- h All persons should be kept out of any vapour cloud area. Any area which is in the path of the vapour cloud should be evacuated immediately, eliminating all sources of ignition at the same time.
- i All persons except those necessary to cope with the incident should be kept away from the human hazard range by policing the dangerous area.

8.5 PREVENTION OF, AND PROTECTION FROM, VAPOUR CLOUD EXPLOSIONS

If LPG is allowed to mix with air and form a concentration between 2%-10% in an enclosed space and is subsequently ignited it will burn and very probably give a pressure wave in front of the flame. In these incidents the pressure can rise up to about 100 psi and destroy vessels and buildings. This explosion risk is common to all fuel gases including LPG and is most likely to occur inside items of plant such as storage tanks etc [26]. The quantification of LPG vapour cloud explosions has been discussed in detail in Chapter 7.

8.5.1 Prevention

Prevention in this context means prevention of CVCE and UVCE, not necessarily prevention of a fireball, still less of escape of flammable material. Whilst the latter undoubtedly is the desirable objective, it is largely a matter of assiduous but ordinary operating and engineering competence [26].

Prevention of ignition may be possible. There are many instances on record [81] where ignition has not occured. But sources of ignition are legion and those cases of non-ignition of major clouds must be considered to have happened by chance rather than by design. Moreover, the emission itself my generate ignition sources, especially if it is forceful. Therefore, ignition at any escape of vapour must be considered likely. Experimental work [82, 83] suggests that conventional water sprays at best have no effect. The criteria for extinction are not as yet precisely quantified. Therefore, techniques of prevention by other means may be envisaged [26]. At the present time prevention can be achieved economically and with certainty only by assuring plant integrity and sound engineering and design, as described in Section 8.1.

8.5.2 Protection

Unconfined vapour cloud explosions may be predicted and their effects forecast by the methods described in detail in Chapter 7. At present they cannot be prevented. Given a major escape the probability is that ignition will occur rather more frequently than 14 times in every 15 incidents [81], and that blast will occur in rather more than half of those occasions.

Therefore, protection is desirable. Unfortunately, like many desirable objectives, it is not easy to achieve. Protection is two-fold,

of the process workers and plant, and of the surrounding community and its buildings. Because most communities and industries are independent entities, despite their obvious dependence upon each other, their contiguous development can only be prevented by law. This can be the only socially certain method of community protection whatever view is taken of the real risks [26].

Buildings can be built to withstand blast. It is desirable, if these buildings are built and designed to accommodate personnel, that they also protect persons within them,. Therefore, all openings and glazed areas should be sealed and should be of no less strength to pressure than the structure itself. Also, self-contained atmospheres within buildings occupied by personnel would be essential to eliminate the possibility of invasion of these atmospheres by hot, and probably toxic, combustion products resulting from vapour cloud explosions.

There are no new concepts involved in protection within process plants. However, civil engineering will enable the design of structures for the process plant of any reasonable process size to withstand statically applied pressure. The pressure involved will be inversely related to the distance of the structures from the explosion centre. Therefore, a degree of compromise exists in most practical situations [26].

8.6 OTHER SAFETY PRECAUTIONS IN THE RUNNING OF LPG PLANT

8.6.1 Inspection of LPG Installations

The following stages of inspection of the installation should be applied [4, 14]:

a <u>During Manufacture</u>:

All vessels, process equipment, pipework etc must be inspected during maufacture, modification or repair by an approved inspection authority. Such inspection and testing should be in accordance with the Code to which the equipment, vessel or pipework is constructed.

b <u>Before service</u>:

All equipment, vessels, valves and accessory joints must be tested by a competent person for soundness at the maximum working pressure and a suitable record should be kept.

c <u>Periodically</u> [14]:

After being brought into service, all equipment, vessel and pipework shall be subjected to one of the following tests, at least every five years by an approved inspection authority:

- i A hydraulic pressure test
- ii Uninsulated equipment shall be fully-examined internally and externally, or
- iii Alternatively, all welds of the equipment, vessels and pipework may be subjected to an ultrasonic or a radiographic test.

Finally, a certificate should be issued, and kept on record, for each piece of equipment, vessel, etc, after each of the tests mentioned above.

8.6.2 Maintenance

The aim of maintenance is to ensure that an LPG installation is always in good working order and will not give unpleasant surprises.

Surprises that are most likely to upset safety. Therefore, in addition to a rigidly-applied Permit-to-Work system, the following points should be taken into consideration in the maintenance of any LPG plant [9, 84].

- i Setting up system for sufficient stock of spare parts with maximum and minimum stock figures, and keeping the stock up to date.
- ii Keeping the plant clean and tidy with no unnecessary obstructions upsetting the work, free access to emergency exits and free access to all operations. It is very difficult to maintain a high degree of safety when things are in a mess.
- iii Keeping all guarding in place, it might be necessary during maintenance to erect fencing around certain equipment or parts of installations.
- iv Coloured paint on the floor and walls can be used to give information to the staff about risks when they cross the painted line on the floor, such as when there is danger to the eyes or the need to use face masks.
- During maintenance, staff should always use work gloves, helmets and safety footwear when these make work safer.
 Synthetic fibres can create static electricity, therfore, these fibres should be avoided in clothing.
- vi Finally, maintenance should be carried out at frequent intervals on all installations including vehicles, pipework, hoses and ancilliary equipment [9].

8.6.3 Training of Staff

The training of staff must be regarded as most vital. Many managers consider, quite wrongly that anybody is capable of running

an LPG plant. Good workers are absolutely necessary for sound performance of the various operations involved in an LPG plant and maintenance or repair requires a really expert technician. Such qualities as care and intelligence are imperative for anybody who handles LPG, or is present within the hazardous area.

One must educate and train the staff, not merely by teaching them how to operate the equipment and how to carry out various operations, but by providing a thorough understanding of the nature of LPG. This should cover how LPG 'reacts' under various conditions and how they themselves should react under certain circumstances. Also the staff should be made completely familiar with the safety regulations.

Many accidents at LPG plants are caused not only by carelessness or non-observance of the precautionary measures, but primarily because of insufficient knowledge of the product handled, lack of education or training in the performance of the actual operations, or ignorance and insufficient understanding of the safety regulations. Such accidents are absolutely unnecessary and it should be emphasised that an LPG plant is not a risky or dangerous place of work provided the regulations are observed. The safety level at any LPG plant is directly proportional to the staff's knowledge of the product, and their training in running of LPG plant [84].

A brief survey of the practical subjects that should be included in the training of operational staff beyond the information about the nature of LPG and the ways it reacts, and the training in understanding and observing the safety regulations follows [84]:

i Daily start-up

ii

Knowledge of the installation and function of the machinery

iii General, practical maintenance and service.

iv Trouble-shooting and repair.

v Knowledge of spare parts and their specifications

vi Knowledge of the instruction manual for the plant.

Such a training course should be held in the LPG plant and should be a practical and thorough instruction in the use of all machinery and equipment in order to make sure that the staff will really familiarise themselves with the equipment and the operations they have to perform.

Finally, the drivers of LPG vehicles should be mature with a high sense of responsibility. They must be able to work correctly without supervision and must possess the ability to evaluate emergency situations and to reach sensible decisions. Also, the driver of a vehicle conveying LPG and any other person engaged in conveyance should be trained in the properties of LPG and the handling methods applicable to the grades being carried, in the use of the fire extinguishing equipment carried on the vehicle, and in the emergency procedure to be followed.

8.6.4 Emergency Planning

Emergency planning with LPG incidents should include the following:

i Teaching:

The emergency procedures should be comprehensively covered in the classroom by both civilian tutor and Fire Service instructor. These procedures are to be then put to the test in two practical ways. Firstly, a tactical exercise should be staged using a floor map where the staff discuss the problems and the

subsequent tactics they will adopt for a carefully planned situation. Secondly, a major exercise is staged on site which may involve a tank fire or LPG spillage. The objects of these exercises include confirmation for the staff that the previous instruction has been correctly assimilated; the checking of the command and control capabilities of staff nominated to be safety officers in charge of the exercise, and to give all staff the necessary confidence in themselves and their equipment to deal with a potentially hazardous situation [85].

ii Pre-planning:

While it is not possible to conduct operations according to a precise set of rules it is essential that proper emergency preplanning is carried out. The success of operations at any incident will largely depend upon the adequacies of pre-planning and training directly related to each of the situations likely to be encountered. These pre-planning operations may include the police, ambulance and local authority departments [85].

iii Water Supplies:

To protect an LPG vessel from fire it is necessary to apply sufficient cooling water. The Technical Bulletin [45] recommends a minimum rate of 0.1 min/m². Because of difficulties in obtaining uniform complete coverage of tank services with water, much higher rates of application will often be required in practice.

It must be remembered that peak requirements in terms of water will necessarily occur at the earlier stage of emergency operations and may need to be sustained for a prolonged period. Pre-planning must therefore include consideration as to the

availability of adequate water supplies and to ensure they do not fail during such operations [85, 86].

iv

Communications and Emergency Control Centre:

The ideal method is a detection sytem which gives an alarm and possibly activates a suppression system. This may not always be practicable on a large open site. Therefore, some combination of break-glass alarms, telephones connected to a multi-line emergency panel, and pocket radios is required [85].

In order to cope with any major incident, provision should be made for a permanent Control and Communication Centre near a main entrance and remote from any operating area [88]. Its function is to receive information and to transmit it to the centres and services concerned, eg Works and City Fire Services, Police, Work Security, other plants on the site, Medical Authorities, etc. The control centre should obviously be provided with detailed site and plant layouts, drawings and a 'call out' list of essential personnel for expert technical assistance for each plant or area. There should also be spare telephones and pocket radios for incident control and an information room to cope with press and radio [87].

Finally, as in any emergency situation, it is of paramount importance to avoid endangering human life in the event of fire involving or seriously exposing LPG equipment, or serious leakage of LPG without a fire. Also, the evacuation of people around the hazardous area should be pre-planned as an emergency evacuation plan.

CHAPTER NINE

CONCLUSIONS AND RECOMMENDATIONS

9.1 CONCLUSIONS

9.1.1 Main Conclusions

The main conclusions arising from this work are:

- Numerous accidents have occurred at LPG Installations leading to a fireball, blast wave or pool fire. These incidents have involved the release of the contents from bulk storage vessels, or rail and road tankers; a list of such incidents is shown in Table 6.1. the three main hazards associated with major LPG releases are:
 - i A Major Fire, including a pool fire
 - ii An Unconfined Vapour Cloud Explosion (UVCE), or,
 - iii A Boiling Liquid Expanding Vapour Explosion (BLEVE)
- In any evaluation of thermal radiation hazards resulting from fireballs, there are five primary parameters which should be considered:
 - i mass of fuel released
 - ii fireball equivalent spherical diameter
 - iii duration time
 - iv fireball height
 - v fireball surface temperature

3. New correlations for thermal radiation damage criteria have been

proposed to fit the data of Stoll and Green [63], which include the exposure of a human forearm to different radiation intensities, using a least square curve fitting technique, as shown in Appendix C, to give three relationships which enables the thermal radiation intensity on human skin to be related to the time of exposure as:

Pain Threshold	$q = 28.2 (t)^{-0.73}$
Threshold Blister	$q = 60.5 (t)^{-0.75}$
Full Blister	$q = 61.8 (t)^{-0.73}$

Furthermore the data from their work which represent tissue damage for the threshold and full blister have been correlated using the same technique to give two relationships between the damage to tissue and the thermal radiation incident on human skin as:

Threshold Blister	$Z = 2.61 (q)^{1.09}$
Full Blister	$Z = 3.65 (q)^{1.31}$

The results predicted from the above correlations exhibited a maximum error of 8% from the Stoll and Green experimental data.

4. Using an isothermal model for different authors' parameters shows that both Roberts' and Marshall's correlations fit the model, giving values very close to those recommended by them in Table 6.3. Also, it appears that Roberts' and Marshall's parameters give values of the rate of growth of fireball diameter close to that achieved by an isothermal model and prove that only these parameters fit an isothermal model.

- 5. The comparison between the values of BLEVE fireball diameter obtained by applying various authors' correlations mentioned in Table 6.7 and the value of fireball diameter obtained from real incidents shows that Crowley's correlation gives a close fit of fireball diameter as shown in Table 6.8 to those observed from real BLEVE incidents mentioned in Section 6.6.
- 6. Whilst it is not possible to deduce from the pool fire thermal radiation calculation procedure the validity of the results obtained, comparison of the results with those published in the literature showed good agreement. However, many factors involved in the pool fire remain unpredictable. It is possible using the method proposed for pool fires in this study as described in Section 6.10 to predict the spacing distance for different values of L/D and mass of flammable material involved for humans, buildings and process equipment. The water rate obtained by the proposed method is consistent with a government guideline and is useful when designing the layout of plant and to keep people away from hazardous areas.
- 7. The TNT equivalent model including Jarrett's equation and Hopkinson's law, Gugan's and Clancey's models have been used to predict the spacing distance at which characteristic values of overpressure are probable from a UVCE. As many

factors involved in the UVCE remain unpredictable, the explosion yield factor is taken at a more conservative value than previously, ie 10%.

None of the four methods of prediction for the spacing distance used is more or less credible than the others using experience from real incidents. Therefore, in general, it is concluded that maximum figures are to be preferred for conservative estimates for use in site layout, emergency planning, etc.

- 8. The blast wave results demonstrate that the Clancey model generally provides the biggest spacing distances and Jarrett's equation usually gives the shortest spacing distance. Also, these results show that, as expected, the spacing distance increases with increasing yield factor.
- 9. Quantification methods, which are recommended in this study, are a vital part of the process of assessment of LPG hazards and risks. These techniques have been discussed in detail in Chapter 6 and 7. The possibility of an LPG major fire and explosion occurrence can be minimised by sound engineering in plant design and layout, good operating practice, proper fire-fighting systems including spray water, proper inspection methods of LPG installations, good maintenance and proper training of personnel in both routine operations and on actions to be taken in an emergency as described in detail in Chapter 8.
- The safety recommendations and restrictions outlined in detail in Chapter 8 may often have considerable economic consequences.

This economic obstacle should generally be put aside in favour of public safety. Certainly the best solution is always the attainment of a high degree of safety without any significant economic loss from spacing, by optimising the process and reducing the risk as much as possible.

9.1.2 Minor Conclusions

In addition the following minor conclusions arose:

- Correlations which were used to predict fireball diameter and duration: it is interesting to note that the average of all exponents is 1/3 (ie the value of a and b in equations 6.1 and 6.2). It is recommended that Robert's parameters (A, B) be used to evaluate the fireball diameter and duration because maximum figures, which Robert's correlations provided, are to be preferred.
- The fireball diameter, duration, height and hazard ranges for humans, buildings and process equipment increase in proportion to the increase of the flammable quantity of fuel.
- 3. The results of blast waves are presented in terms of spacing distance related to the flammable quantity released and to several characteristic values of overpressure which correspond to different levels of damage effect, as shown in Table 5.2. Also these results show that predicted spacing distances according to TNT equivalent methods, Gugan and Clancey have gross deviations due to several assumptions and empirical equations which may affect their accuracy.

4. In the case of fireballs resulting from UVCE, the hazard ranges based on fireball surface temperature predicted using the thermodynamic equation gives a slightly smaller hazard range than the results based on temperature predicted using isothermal equation. However, the difference between isothermal and thermodynamic temperature results are not significant and can be neglected in large scale fireballs resulting from UVCE. Either method could be used to predict the theoretical surface temperature of a fireball.

But in the case of a fireball resulting from a BLEVE, the hazard ranges based on temperature obtained from the thermodynamic equation have large deviations compared with the hazard ranges predicted by using the temperature obtained from the BLEVE equation.

- 5. No precise method can be proposed for full hazard assessment. Each method has its advantages and disadvantages depending on the assumptions and experimental observations. Therefore, the choice of a quantified method depends entirely on the purpose, urgency and accuracy of results and no method can fully identify all the hazards in LPG plants, such as human error which is a major source of risk.
- 6. Fireball diameter increases linearly with the duration of combustion both in the isothermal and BLEVE models, and that is because it has been assumed that the equivalent spherical diameter of fireball increases at constant rate throughout

combustion.

9.2 RECOMMENDATIONS FOR FURTHER WORK

As a continuation of this work, the following areas appear worth studying:

- The shape of the fireball changes continuously during its rise. At ignition, it is hemispherical, then spherical and after lift-off takes several shapes, eg oblate, spheroidal or mushroom. Therefore models could be derived which take account of the change in shape as a function of time.
- 2. All the present models and correlations for UVCE and BLEVE effects are based upon small-scale experimental work. Therefore, the accuracy and the validity of this work upon scale-up by a factor of several thousands remains unproven. Large scale experiments would therefore be desirable in order to approach more closely the size of real incidents.

For BLEVE experiments as a first attempt, masses of approximately 1 tonne are recommended. These BLEVE'S could be engineered using explosively-ruptured tanks (as at SMRE) or using large piston releases at atmospheric pressure. The amount required to generate a realistic >5 tonnes UVCE may be impracticable on economic and environmental grounds. However, release of a premixed vapour air mixture could be engineered using plastic balloons.

3. The quantification and assessment of hazards by mathematical

models described in Chapters 6 and 7, shows that prediction of safe and socially acceptable spacing distances require further study. Social and economic aspects should be considered by optimising the process and reducing the risk to the public and the environment as far as possible because these aspects cannot be determined with any precision with the present level of knowledge.

- 4. Further investigations and studies are required to cover many factors involved in a vapour cloud explosion, the effects of which are still difficult to predict (eg wind velocity and direction, weather conditions, degree of confinement, spill rate and operator response).
- 5. Further experimental and theoretical studies are recommended to reduce the lack of accuracy of many factors affecting the thermal radiation hazard ranges related to fireballs or pool fires (eg fraction of fuel cracked to produce soot, air relative humidity, wind speed and direction, and local topography).

APPENDIX A

Computer program for evaluation of initial reduced overpressure (\overline{P}_{SO})

```
С
C
c
       EVALUATION OF INITIAL REDUCED OVERPRESSURE (PSO):
000
       *****
                     ******
                                                    *****
C
       DIMENSION PS0(500), PSNC(10), DIFF(500), RS(500), FS(500)
        PRINT 15
       FORMAT(//10X, 'INITIAL REDUCED OVERPRESSURE (PSO) ',
  15
     @'RESULTS:',/10X,43('*')/)
          READ *.N
         READ *, (PSNC(11), 11=1, N)
С
         DO 3 K=1,N
С
C
         PRINT 40,K,PSNC(K)
       FORMAT(/10X, 'ESTIMATED STORAGE PRESSURE (', 11, ')=', F8.1,
  40
     @/10X,26('-')/)
        1=1
        PSO(1)=1.0
        RS(|)=(1.0+PSNC(K))*((1.0-((0.2*1.08*PSO(|))/((2.8*(2.8+2.4
  10
     @*PSO(1)))**0.5)))**12.0)
C
       FS(1)=1.0+PSO(1)
       DIFF(1) = FS(1) - RS(1)
        IF (ABS(DIFF(1)).GT.0.3) THEN
      PSO(1+1)=PSO(1)+0.2
       |=|+1
        GO TO 10
          ELSE
C
      PRINT 300, I, DIFF(1)
  300
       FORMAT(10X, 'DIFFERENCE(', 12, ')=', F8.4/)
С
C
      PRINT 400, 1, PSO(1)
     FORMAT(10X, 'THE VALUE OF PSO(', 12, ')=', F8.1/)
 400
С
С
        ENDIF
C
  3
            CONTINUE
C
           STOP
C
              END
```

INITIAL REDUCED OVERPRESSURE (PSO) RESULTS:

EST IMATED STORAGE PRESSURE (1)= 16.0 DIFFERENCE (10)= -0.0119 THE VALUE OF PSO(10)= 2.8 EST IMATED STORAGE PRESSURE (2)= 20.0 DIFFERENCE (12)= 0.1185 THE VALUE OF PSO(12)= 3.2 EST IMATED STORAGE PRESSURE (3)= 23.0 DIFFERENCE (13)= 0.0447 THE VALUE OF PSO(13)= 3.4 EST IMATED STORAGE PRESSURE (4)= 25.0

DIFFERENCE(14)= 0.1870

THE VALUE OF PSO(14)= 3.6

APPENDIX B

Computer program for evaluation of fireball duration and diameter using different authors' parameters

```
C
C
C PROGRAM FOR EVALUATION OF FIREBALL DURATION AND DIAMETER
C USING GAYLE, BRASIE, MARSHALL, ROBERT, HARDEE, AND FAY EQUATIONS:
C ==
C
C
C
      DIMENSION W(30), DG(30), DB(30), DM(30), DR(30), DH(30), DF(30),
     @TG(30), TB(30), TM(30), TR(30), TH(30), TF(30)
С
C
       PRINT 21
 21
      FORMAT(/10X, 'FIREBALL DURATION & DIAMETER RESULTS: ', /
     @10X,37('*')//)
С
С
       READ *.N
       READ *, (W(11), 11=1, N)
С
C
       DO 3 I=1.N
       TG(1)=0.245*(W(1)**0.356)
       TB(1)=0.300*(W(1)**0.333)
       TM(1)=0.380*(W(1)**0.333)
       TR(1)=0.450*(W(1)**0.333)
       TH(1)=2.570*(W(1)**0.166)
       TF(1)=2.530*(W(1)**0.166)
C
C
       DG(1)=3.680*(W(1)**0.326)
       DB(1)=3.800*(W(1)**0.333)
       DM(1)=5.500*(W(1)**0.333)
       DR(1)=5.800*(W(1)**0.333)
       DH(1)=6.360*(W(1)**0.325)
       DF(1)=6.280*(W(1)**0.333)
C
  3
          CONTINUE
C
С
      PRINT 100
      FORMAT(10X, 'FIREBALL DURATION (SECONDS): ',/,10X,25('-')//)
 100
С
      PRINT 200
      FORMAT(8X, 'MASS GAYLE BRASIE MARSH ROBERT HARDEE FAY', /
 200
     @5X,50('-'))
С
      DO 5 J=1,N
      PRINT 300,W(J),TG(J),TB(J),TM(J),TR(J),TH(J),TF(J)
 300
     FORMAT(5X, F9.1, 6(F6.1, 1X))
C
      CONTINUE
 5
С
```

C	
	PRINT 400
400	FORMAT(///10X, 'FIREBALL DIAMETER (METERS): './10X.27('-')//)
С	
	PRINT 500
500	FORMAT(8X, 'MASS GAYLE BRASIE MARSH ROBERT HARDEF FAY' /
	@5X,50('-'))
С	
	DO 10 M=1,N
	PRINT 600, W(M), DG(M), DB(M), DM(M), DR(M), DH(M), DF(M)
600	FORMAT(5X, F9.1, 6(F6.1, 1X))
С	
10	CONTINUE
С	
	STOP
	END
FIREBALL DURATION & DIAMETER RESULTS:

FIREBALL DURATION (SECONDS):

MASS	GAYLE	BRASIE	MARSH	ROBERT	HARDEE	FAY
0.1	0.1	0.1	0.2	0.2	1.8	1.7
1.0	0.2	0.3	0.4	0.5	2.6	2.5
5.0	0.4	0.5	0.6	0.8	3.4	3.3
10.0	0.6	0.6	0.8	1.0	3.8	3.7
25.0	0.8	0.9	1.1	1.3	4.4	4.3
50.0	1.0	1.1	1.4	1.7	4.9	4.8
75.0	1.1	1.3	1.6	1.9	5.3	5.2
100.0	1.3	1.4	1.8	2.1	5.5	5.4
250.0	1.7	1.9	2.4	2.8	6.4	6.3
500.0	2.2	2.4	3.0	3.6	7.2	7.1
750.0	2.6	2.7	3.4	4.1	7.7	7.6
1000.0	2.9	3.0	3.8	4.5	8.1	8.0
5000.0	5.1	5.1	6.5	7.7	10.6	10.4
10000.0	6.5	6.4	8.2	9.7	11.9	11.7
25000.0	9.0	8.7	11.1	13.1	13.8	13.6
50000.0	11.5	11.0	13.9	16.5	15.5	15.2
75000.0	13.3	12.6	16.0	18.9	16.6	16.3
100000.0	14.8	13.9	17.6	20.8	17.4	17.1
250000.0	20.5	18.8	23.8	28.2	20.2	19.9
500000.0	26.2	23.7	30.0	35.6	22.7	22.3
750000.0	30.2	27.1	34.4	40.7	24.3	23.9
1000000.0	33.5	29.9	37.8	44.8	25.5	25.1

FIREBALL DIAMETER (METERS):

MASS	GAYLE	BRASIE	MARSH	ROBERT	HARDEE	FAY
0.1	1.7	1.8	2.6	2.7	3.0	2.9
1.0	3.7	3.8	5.5	5.8	6.4	6.3
5.0	6.2	6.5	9.4	9.9	10.7	10.7
10.0	7.8	8.2	11.8	12.5	13.4	13.5
25.0	10.5	11.1	16.1	16.9	18.1	18.3
50.0	13.2	14.0	20.2	21.3	22.7	23 1
75.0	15.0	16.0	23.2	24.4	25.9	26 4
100.0	16.5	17.6	25.5	26.9	28.4	29 1
250.0	22.3	23.9	34.6	36.5	38 3	30 5
500.0	27.9	30.1	43.6	45 9	47 9	10 7
750.0	31.9	34.4	49 9	52 6	54 7	49.7
1000.0	35.0	37.9	54.9	57.9	60.0	62.7

5000.0	59.1	64.8	93.8	98.9	101.3	107.1	
10000.0	74.1	81.6	118.1	124.6	126.9	134.9	
25000.0	99.9	110.7	160.3	169.0	170.9	183.0	
50000.0	125.2	139.5	201.9	212.9	214.1	230.5	
75000.0	142.9	159.7	231.1	243.7	244.3	263.8	
100000.0	157.0	175.7	254.3	268.2	268.2	290.4	
250000.0	211.6	238.4	345.0	363.9	361.2	394.0	
500000.0	265.3	300.3	434.6	458.3	452.5	496.3	
750000.0	302.8	343.7	497.5	524.6	516.2	568.0	
000000.0	332.5	378.3	547.5	577.3	566.8	625.1	

APPENDIX C

Regression Analysis Computer Program

CPR	YRAM FOR MUTIPLE LINEAR RECRASSION ANALYOLO
C ===	A A A A A A A A A A A A A A A A A A A
С	
C	
C	MAIN PROCEAM
C	MATN PROGRAM
č	
C	
	DIMENSION X(16,2), Y(16), A(4), B(2), XBAR(2), YHAT(16),
	@AA(2,2),T(16),F21(16)
C	
C	
C	READ *, NB, NX, M
U	NN=NB*NB
	IF(NB.LE.O) THEN
	GO TO 555
	ELSE
~	READ *, (F21(KK), KK=1, M)
C	D 1 K 1 M
С	W I K=I,M
c	
	X(K, 1) = ALOG(F21(K))
С	
С	
1	CONTINUE
c	
U	BEAD * (T(J J) J =1 M)
С	······································
	DO 3 I=1,M
5	Y(1)=ALOG(T(1))
3	CONTINUE
C	
C	CALL LESOLY Y NE M & E YEAR YHAT AA NOL
555	STOP
	ENDIF
	END
C	
C	
c	
	SUBROUTINE LESQ(X,Y,N,M,A,B,XBAR, YHAT, AA N2)
С	
	DIMENSION X(M,N), Y(M), A(N2), B(N), XBAR(N), YHAT(M), AA(N,N)
C	
C	PRINT 101
101	FORMAT (10X, 'MULTIPLE LINEAR RECRESSION ANALYSIS) (10Y 10(1 1))
C	AVALISIS , TOLITILE ETHEAN REGRESSION ANALISIS , / TOX, 40('-')//)

```
C CALCULAT AVERAGE X AND Y VALUES
С
С
        DO 8 |=1,N
         SLMX=0.0
        DO 13 J=1,M
         SLMX=SLMX+X(J, I)
  13
  8
         XBAR(I)=SUMX/FLOAT(M)
          SUMY=0.0
         DO 15 K=1,M
  15
          SUMY=SUMY+Y(K)
         YBAR=SUMY/FLOAT(M)
       PRINT 180
 180
       FORMAT(//,6X, 'VARIABLE AVERAGE VALUES', /,6X,23('-')/)
       PRINT 240, (11, XBAR(11), 11=1, N)
       FORMAT(6(6X, 'XBAR(', 12, ')=', 1PE14.7)/)
 240
        PRINT 260, YBAR
 260
       FORMAT(6X, 'YBAR=', 1PE14.7, /)
C
C
C
    CALCULATE REGRESSION MATRICES
С
С
C
      KK=1
      DO 25 I=1,N
      DO 25 J=1,N
      SLMA=0.0
      SUMB=0.0
       DO 30 K=1,M
      SUMA=SUMA+(X(K,I)-XBAR(I))*(X(K,J)-XBAR(J))
 30
       SUMB=SUMB+(Y(K)-YBAR)*(X(K, I)-XBAR(I))
      AA(I, J) = SUMA
      A(KK)=SUMA
        KK=KK+1
 25
      B(1)=SUMB
      PRINT 310
 310
      FORMAT(//, 10X, ' A MATRIX', /, 10X, 9('-')/)
       DO 55 ||=1,N
       PRINT 360, (AA(11, JJ), JJ=1, N)
  55
 360
       FORMAT(/,50(2X,E10.4))
       PRINT 410
 410
       FORMAT(//,6X, 'B MATRIX',/,6X,9('-'))
       PRINT 440, (B(KK), KK=1, N)
 440
      FORMAT(/, 50(2X, E10.4))
С
C
C
   SOLVE REGRESSION MATRICES FOR COEFFICIENTS
С
        С
С
     CALL STMG(A, B, N, KS, N2)
       SUMX=0.0
      DO 60 I=1,N
```

```
SUMX=SUMX+B(1)*XBAR(1)
  60
       AZERO=YBAR-SUMX
        PRINT 460
        FORMAT(//10X, 'VALUES OF THE REGRESSION COEFFICIENTS',/,
  460
      @10X,40('-')/)
       PRINT 470, (JJ, B(JJ), JJ=1, N)
  470 FORMAT(5(6X, 'AHAT(', 12, ')=', 1PE16.8, /))
        PRINT 480, AZERO
  480
        FORMAT(6X, ' AZERO=', 1PE16.8, /)
C
          AHATO=EXP(AZERO)
         PRINT 490, AHATO
  490
        FORMAT(6X, ' AHAT(0)=', 1PE16.8, /)
C
С
С
C
      CALCULATE S AND R VALUES
C
C
C
       STEST=0.0
       DO 80 J=1.M
      SUMS1=0.0
      DO 90 K=1,N
       SLMS1=SLMS1+B(K)*X(J,K)
  90
       YHAT(J)=AZERO+SUMS1
      DIFF=(Y(J)-YHAT(J))**2
  80 STEST=STEST+DIFF
      SUMST=0.0
      DO 92 1=1,M
  92 SUMST=SUMST+(Y(I)-YBAR)**2
      SUMSR=SUMST-STEST
      RTEST=SUMSR/SUMST
       PRINT 520
 520 FORMAT(////,6X, 'EXPERIMENTAL VALUES', 11X, 'REGRESSION VALUES',
     @/,6X,50('-'))
С
C
      DO 115 KK=1,M
 115 PRINT 555, KK, Y(KK), KK, YHAT(KK)
     FORMAT(/7X, 'Y(', 12, ')=', 1PE16.6, 12X, 'YHAT(', 12, ')=', 1PE16.6)
 555
C
       PRINT 560, SUMST, STEST, RTEST
       FORMAT(///,6X, 'SLMST=', 1PE16.8, /,6X, 'STEST=', 1PE16.6, /,
 560
     @6X, 'RTEST=', 1PE16.6,/)
C
        RETURN
        END
С
С
С
C
      SUBROUTINE STMG(A, B, N, KS, NS)
```

```
DIMENSION A(NS), B(N)
      TOL=0.0
      KS=0
      JJ=-N
      DO 65 J=1,N
      JY=J+1
      JJ=JJ+N+1
      BIGA=0.0
       II=JJ-J
     DO 33 I=J,N
С
000
   SEARCH FOR MAXIMUM COEFFICIENT IN COLUMN:
          С
      |J=||+1
      IF((ABS(BIGA)-ABS(A(IJ))).GE.O.O) THEN
      GO TO 33
       ELSE
      BIGA=A(IJ)
       IMAX=1
            ENDIF
       CONTINUE
  33
С
С
  TEST FOR PIVOT LESS THAN TOLERANC
С
       IF((ABS(BIGA)-TOL).GT.0.0) THEN
       GO TO 40
        ELSE
      KS=1
      RETURN
С
CC
    INTERCHANE ROWS IF NECEEARY
        C
     11=J+N^{*}(J-2)
  40
         ENDIF
     II=IMAX-J
      DO 51 K=J,N
     11=11+N
     12=11+11
     SAVE=A(11)
     A(|1)=A(|2)
     A(12)=SAVE
С
С
    DIVIDE EQUATION BY LEADING COEFFICIENT
C
51
      A(11)=A(11)/BIGA
     SAVE=B(IMAX)
     B(IMAX)=B(J)
     B(J)=SAVE/BIGA
С
С
    ELIMINATE NEXT VARIABLE
С
```

```
IF((J-N).EQ.O) THEN
      GO TO 70
       ELSE
      IQS=N*(J-1)
         ENDIF
      DO 65 IX=JY,N
      IXJ=IQS+IX
      ||=J-|X|
      DO 61 JX=JY,N
      |XJX=N*(JX-1)+|X
      JJX=IXJX+II
61
      A(IXJX)=A(IXJX)-(A(IXJ)*A(JJX))
      B(IX)=B(IX)-(B(J)*A(IXJ))
65
          CONTINUE
     BACK SOLUTION
 70
      NY=N-1
      | |=N*N
       DO 81 J=1,NY
       IA=II-J
       IB=N-J
       IC=N
      DO 81 K=1, J
       B(IB)=B(IB)-A(IA)*B(IC)
       IA=IA-N
 81
        IC=IC-1
```

IC=IC-RETURN

END

000

С

(FOR HUMAN PAIN)

VARIABLE AVERAGE VALUES

XBAR(1)= 1.6177132E+00 XBAR(2)= 0.0000000E+00 XBAR(YBAR= 2.1478679E+00

A MATRIX

0.2184E+01 0.0000E+01

0.0000E+01 0.0000E+01

B MATRIX ------

-.1607E+01 0.0000E+01

VALUES OF THE REGRESSION COEFFICIENTS

AHAT(1)= -7.35716443E-01 AHAT(2)= 0.0000000E+00 AHAT(AZERO= 3.33804613E+00

AHAT(0)= 2.81640445E+01

EXP	ERIMEN	ITAL VALUES	REGRESSION VALU	ES
Y(1)=	2.817204E+00	YHAT(1)=	2.757965E+00
Y(2)=	2.529721E+00	YHAT(2)=	2.635061E+00
Y(3)=	2.124415E+00	YHAT(3)=	2.070578E+00

Y(4)=	1.836733E+00	YHAT(4) =	1.790000E+00
Y(5)=	1.431268E+00	YHAT(5)=	1.485735E+00

SUMST=	1.20469258E+00
STEST=	2.265476E-02
RTEST=	9.811946E-01

(FOR HUMAN THRESHOLD BLISTER) ______

VARIABLE AVERAGE VALUES

XBAR(1)= 2.5855118E+00 XBAR(2)= 0.0000000E+00 XBAR(YBAR= 2.1478679E+00

A MATRIX -----

0.2103E+01 0.0000E+01

0.0000E+01 0.0000E+01

B MATRIX -----

-.1590E+01 0.0000E+01

VALUES OF THE REGRESSION COEFFICIENTS

AHAT(1)= -7.56293498E-01 AHAT(2)= 0.0000000E+00 AHAT(AZERO= 4.10327367E+00

AHAT(0)= 6.05381519E+01

EXP	ERIMEN	ITAL VALUES	REGRESSION	VALUE	s
Y(1)=	2.817204E+00	YHAT (1)=	2.800356E+00
Y(2)=	2.529721E+00	YHAT (2)=	2.549753E+00
Y(3)=	2.124415E+00	YHAT (3)=	2.140499E+00

Y(4)=	1.836733E+00	YHAT(4) =	1.807958E+00
Y(5)=	1.431268E+00	YHAT(5)=	1.440772E+00

SUMST=	1.20469258E+00
STEST=	1.862134E-03
RTEST=	9.984543E-01

(FOR HUMAN FULL BLISTER)

VARIABLE AVERAGE VALUES

XBAR(1)= 2.9875514E+00 XBAR(2)= 0.0000000E+00 XBAR(YBAR= 1.9325737E+00

A MATRIX -----

0.1137E+01 0.0000E+01

0.0000E+01 0.0000E+01

B MATRIX ------

-.8341E+00 0.0000E+01

VALUES OF THE REGRESSION COEFFICIENTS

AHAT(1)= -7.33791171E-01 AHAT(2)= 0.0000000E+00 AHAT(AZERO= 4.12481256E+00

AHAT(0)= 6.18562200E+01

EXPI	ERIMEN	ITAL VALUES	REGRESSION	VALUE	S
Y(1)=	2.529721E+00	YHAT(1)=	2.545868E+00
Y(2)=	1.836733E+00	YHAT (2)=	1.780656E+00
Y(3)=	1.431268E+00	YHAT (3)=	1.471197E+00

SUMST=	6.17077655E-01
STEST=	4.999757E-03
RTEST=	9.918977E-01

(HUMAN TISSUE DAMAGE FOR THE THRESHOLD BLISTER)

VARIABLE AVERAGE VALUES

XBAR(1) = 2.1537312E+00 XBAR(2) = 0.0000000E+00 XBAR(YBAR= 3.3139853E+00

A MATRIX -----

0.1204E+01 0.0000E+01

0.0000E+01 0.0000E+01

B MATRIX -----

0.1316E+01 0.0000E+01

VALUES OF THE REGRESSION COEFFICIENTS

AHAT(1)= 1.09334027E+00 AHAT(2)= 0.0000000E+00 AHAT(AZERO= 9.59224212E-01

AHAT(0)= 2.60967137E+00

EXPERIMENTAL VALUES			REGRESSION VALUES		
Y(1)=	2.708050E+00	YHAT (1)=	2.524087E+00
Y(2)=	2.833213E+00	YHAT (2)=	2.967398E+00
Y(3)=	3.295837E+00	YHAT (3)=	3.725070E+00

Y(4) = 4.418841E+00 YHAT(4) = 4.039386E+00

SUMST=	1.81933363E+00
STEST=	3.800745E-01
RTEST=	7.910914E-01

(HUMAN TISSUE DAMAGE FOR THE FULL BLISTER)

VARIABLE AVERAGE VALUES

XBAR(1)= 1.9325737E+00 XBAR(2)= 0.0000000E+00 XBAR(YBAR= 3.8318586E+00

A MATRIX

0.6171E+00 0.0000E+01

0.0000E+01 0.0000E+01

B MATRIX

0.8096E+00 0.0000E+01

VALUES OF THE REGRESSION COEFFICIENTS

AHAT(1)= 1.31192471E+00 AHAT(2)= 0.0000000E+00 AHAT(AZERO= 1.29646737E+00

AHAT(0)= 3.65635754E+00

EXPER IME	NTAL VALUES	REGRESSION VALUES		
Y(1)=	2.995732E+00	YHAT (1)=	3.174183E+00
Y(2)=	3.988984E+00	YHAT(2)=	3.706123E+00
Y(3)=	4.510860E+00	YHAT(3)=	4.615270E+00

SUMST=	1.18483787E+00
STEST=	1.227569E-01
RTEST=	8.963935E-01

APPENDIX D

Computer program for evaluation of fireball surface temperatures using isothermal model equation for different authors' parameters

00000	
С	PROGRAM FOR EVALUATION OF FIREBALL SURFACE TEMPERATURES:
C	
c	(USTING TSU-THERIVIAL MODEL EQUATION) ************************************
С	
C	
c	
	PRINT 11
c	FORMAT(5X, 'METHANE FIREBALL SURFACE TEMPERATURE: ',/,5X,42('*')//)
	CALL TEMP(GN, FC, V, SECMA, TS)
C	
•	PRINT 33
33	FORMAT(5X, 'ETHANE FIREBALL SURFACE TEMPERATURE: ',/,5X,42('*')//)
C	CALL TEMP (GN26, FC26, V26, SECMA26, TS26)
С	
С	PRINT 12
12	FORMAT(5X, 'PROPANE FIREBALL SURFACE TEMPERATURE: '. /. 5X. 43('*')//)
С	
с	CALL TEVIP (GNT, FCT, VT, SECMAT, TST)
С	
13	PRINT 13 FORMAT(5X 'BUTANE ELPERALL SUBFACE TEMPERATURE ((5X 10(1)))
C	POINT (0x, BOTALE PIREBALE SURFACE TEMPERATURE: , /, 5x, 43('*')//)
~	CALL TEMP(GN2, FC2, V2, SECMA2, TS2)
c	
	PRINT 14
C	FORMAT(5X, 'PENTANE FIREBALL SURFACE TEMPERATURE: ', /, 5X, 43('*')//)
	CALL TEMP(GN3, FC3, V3, SECMA3, TS3)
C	
-	STOP
~	END
c	
С	
С	SUBBOUTINE TEMP (GN EC V SECMA TS)
С	
C	
c	

```
DIMENSION T(1000), R1(1000), R2(1000), DIF(1000)
C
       READ *, WTM, GN, FC, V, SECMA, A, ZH, TS
C
         1=1
       T(1)=TS
  5
      R1(1)=(17.26*A*SEQMA)*((T(1)/1000.0)**4.0)
      R2(1)=(((WTM*ZH*FC)-((31.61*V)*(T(1)-290.0)))/((T(1)*(V+
     @(GN*FC)))))*((((V+(GN+1.0)*FC)*(T(1)-290.0))/WTM)**0.33333)
С
CC
        DIF(1)=R2(1)-R1(1)
         IF (ABS(DIF(1)).GT.1.0) THEN
        T(|+1)=T(|)+2.0
        |=|+1
           GO TO 5
            ELSE
      PRINT 66, 1, R1(1)
      FORMAT(10X, ' THE VALUE OF RIGHT TERM (', 13, ')=', F10.5)
  66
C
      PRINT 67, 1, R2(1)
      FORMAT(10X, ' THE VALUE OF LEFT TERM (', 13, ')=', F10.5)
  67
C
      PRINT 68, I, DIF(1)
      FORMAT(10X, ' THE DIFFERANCE (', 13, ')=', F10.5)
 68
C
       PRINT 100, 1, T(1)
       FORMAT(10X, 'SURFACE TEMPERATURE (', 13, ')=', F10.3, ' K.DEGREE')
 100
         ENDIF
              PRINT 16
          FORMAT(3X, ///)
  16
         RETURN
         END
С
C
```

ISOTHERMAL FIREBALL SUREFACE TEMPERATURE RESULS

THESE RESULTS ARE BASED ON GAYLE PARAMETERS

> THE VALUE OF RIGHT TERM (270)= 88.35633 THE VALUE OF LEFT TERM (270)= 89.26959 THE DIFFERANCE (270)= 0.91326 SURFACE TEMPERATURE (270)= 2138.000 K.DEGREE

> THE VALUE OF RIGHT TERM (292)= 95.85745 THE VALUE OF LEFT TERM (292)= 96.61901 THE DIFFERANCE (292)= 0.76156 SURFACE TEMPERATURE (292)= 2182.000 K.DEGREE

> THE VALUE OF RIGHT TERM (295)= 96.91615 THE VALUE OF LEFT TERM (295)= 97.78817 THE DIFFERANCE (295)= 0.87202 SURFACE TEMPERATURE (295)= 2188.000 K.DEGREE

> THE VALUE OF RIGHT TERM (295)= 96.91615 THE VALUE OF LEFT TERM (295)= 97.69197 THE DIFFERANCE (295)= 0.77582 SURFACE TEMPERATURE (295)= 2188.000 K.DEGREE

PENTANE FIREBALL SURFACE TEMPERATURE:

THE VALUE OF RIGHT TERM (296)= 97.27099 THE VALUE OF LEFT TERM (296)= 98.07716 THE DIFFERANCE (296)= 0.80617 SURFACE TEMPERATURE (296)= 2190.000 K.DEGREE ISOTHERMAL FIREBALL SURFACE TEMPERATURE RESULTS

THESE RESULTS ARE BASED ON BRASIE PARAMETERS

METHANE FIREBALL SURFACE TEMPERATURE:

THE VALUE OF RIGHT TERM (244)= 98.04360 THE VALUE OF LEFT TERM (244)= 98.93226 THE DIFFERANCE (244)= 0.88866 SURFACE TEMPERATURE (244)= 2086.000 K.DEGREE

ETHANE FIREBALL SURFACE TEMPERATURE:

THE VALUE OF RIGHT TERM (264)= 105.78281 THE VALUE OF LEFT TERM (264)= 106.54972 THE DIFFERANCE (264)= 0.76691 SURFACE TEMPERATURE (264)= 2126.000 K.DEGREE

PROPANE FIREBALL SURFACE TEMPERATURE:

THE VALUE OF RIGHT TERM (267)= 106.98203 THE VALUE OF LEFT TERM (267)= 107.54098 THE DIFFERANCE (267)= 0.55895 SURFACE TEMPERATURE (267)= 2132.000 K.DEGREE

> THE VALUE OF RIGHT TERM (267)= 106.98203 THE VALUE OF LEFT TERM (267)= 107.35042 THE DIFFERANCE (267)= 0.36839 SURFACE TEMPERATURE (267)= 2132.000 K.DEGREE

THE VALUE OF RIGHT TERM (268)= 107.38403 THE VALUE OF LEFT TERM (268)= 107.67764 THE DIFFERANCE (268)= 0.29361 SURFACE TEMPERATURE (268)= 2134.000 K.DEGREE ISOTHERMAL FIREBALL SURFACE TEMPERATURE RESULTS

THESE RESULTS ARE BASED ON MARSHALL PARAMETERS

> THE VALUE OF RIGHT TERM (213)= 110.06935 THE VALUE OF LEFT TERM (213)= 110.86079 THE DIFFERANCE (213)= 0.79143 SURFACE TEMPERATURE (213)= 2024.000 K.DEGREE

ETHANE FIREBALL SURFACE TEMPERATURE:

THE VALUE OF RIGHT TERM (231)= 118.11179 THE VALUE OF LEFT TERM (231)= 118.69158 THE DIFFERANCE (231)= 0.57978 SURFACE TEMPERATURE (231)= 2060.000 K.DEGREE

PROPANE FIREBALL SURFACE TEMPERATURE:

THE VALUE OF RIGHT TERM (233)= 119.03184 THE VALUE OF LEFT TERM (233)= 119.83356 THE DIFFERANCE (233)= 0.80171 SURFACE TEMPERATURE (233)= 2064.000 K.DEGREE

BUTANE FIREBALL SURFACE TEMPERATURE:

THE VALUE OF RIGHT TERM (233)= 119.03184 THE VALUE OF LEFT TERM (233)= 119.52421 THE DIFFERANCE (233)= 0.49237 SURFACE TEMPERATURE (233)= 2064.000 K.DEGREE

PENTANE FIREBALL SURFACE TEMPERATURE:

THE VALUE OF RIGHT TERM (234)= 119.49388 THE VALUE OF LEFT TERM (234)= 119.77812 THE DIFFERANCE (234)= 0.28424 SURFACE TEMPERATURE (234)= 2066.000 K.DEGREE ISOTHERMAL FIREBALL SURFACE TEMPERATURE RESULTS

THESE RESULTS ARE BASED ON ROBERT PARAMETERS

> THE VALUE OF RIGHT TERM (191)= 119.37519 THE VALUE OF LEFT TERM (191)= 119.61138 THE DIFFERANCE (191)= 0.23619 SURFACE TEMPERATURE (191)= 1980.000 K.DEGREE

> THE VALUE OF RIGHT TERM (207)= 127.28148 THE VALUE OF LEFT TERM (207)= 127.83855 THE DIFFERANCE (207)= 0.55707 SURFACE TEMPERATURE (207)= 2012.000 K.DEGREE

> THE VALUE OF RIGHT TERM (209)= 128.29668 THE VALUE OF LEFT TERM (209)= 128.82691 THE DIFFERANCE (209)= 0.53023 SURFACE TEMPERATURE (209)= 2016.000 K.DEGREE

> THE VALUE OF RIGHT TERM (209)= 128.29668 THE VALUE OF LEFT TERM (209)= 128.43076 THE DIFFERANCE (209)= 0.13408 SURFACE TEMPERATURE (209)= 2016.000 K.DEGREE

PENTANE FIREBALL SURFACE TEMPERATURE:

THE VALUE OF RIGHT TERM (209)= 128.29668 THE VALUE OF LEFT TERM (209)= 129.00556 THE DIFFERANCE (209)= 0.70889 SURFACE TEMPERATURE (209)= 2016.000 K.DEGREE

APPENDIX E

Computer program for evaluation of fireball surface temperatures using BLEVE's model equation

00000	
CC	PROGRAM FOR EVALUATION OF FIREBALLS SURFACE TEMPERATURES:
0000000	(OF B.L.E.V.E. FIREBALLS) ************************************
11	PRINT 11 FORMAT(5X, 'METHANE FIREBALL SURFACE TEMPERATURE: ',/,5X,42('*')//)
c c	CALL TEMP(TS1,RS1,C1)
	PRINT 26
26 C	FORMAT(5X, 'ETHANE FIREBALL SURFACE TEMPERATURE: ', /, 5X, 42('*')//)
cc	CALL TEMP(TS2,RS2,C2)
	PRINT 12
12 C	FORMAT(5X, 'PROPANE FIREBALL SURFACE TEMPERATURE: ', /, 5X, 43('*')//)
cc	CALL TEMP(TS3,RS3,C3)
13 C	PRINT 13 FORMAT(5X, 'BUTANE FIREBALL SURFACE TEMPERATURE: ',/,5X,43('*')//)
c	CALL TEMP(TS4,RS4,C4)
14 C	PRINT 14 FORMAT(5X, 'PENTANE FIREBALL SURFACE TEMPERATURE: ',/,5X,43('*')//)
C	CALL TEMP(TS5,RS5,C5)
~	STOP END
0000	
0000	SUBROUTINE TEMP(TS,RS,C)
0	DIMENSION T(900), R1(900), R2(900), DIF(900)

```
С
       READ *, WTM, ZH, TS, RS, C
С
         1=1
       T(I)=TS
  5
       R1(1)=(69700/WTM)*((T(1)/1000.0)**4.0)
       R2(1)=ZH-(((1000.0*RS*C)*(((T(1)/1000.0)-0.290))))
С
CC
С
         DIF(1)=R2(1)-R1(1)
         IF (ABS(DIF(1)).GT.200.0) THEN
         T(|+1)=T(|)+2.0
        1=1+1
           GO TO 5
            ELSE
      PRINT 66, 1, R1(1)
      FORMAT(10X, ' THE VALUE OF RIGHT TERM (', 13, ')=', F12.3)
  66
C
      PRINT 67, 1, R2(1)
      FORMAT(10X, ' THE VALUE OF LEFT TERM (', 13, ')=', F12.3)
  67
С
      PRINT 68, I, DIF(I)
 68
      FORMAT(10X, ' THE DIFFERANCE (', 13, ')=', F12.3)
C
       PRINT 100, 1, T(1)
       FORMAT(10X, 'B.L.E.V.E. SURFACE TEMPERATURE (', 13, ')=', F10.3,
 100
     @' K.DEGREE',/)
С
         ENDIF
              PRINT 16
  16
           FORMAT(3X, ///)
         RETURN
         END
C
C
```

BLEVE FIREBALL SURFACE TEMPERATURE RESULTS

METHANE FIREBALL SURFACE TEMPERATURE:

THE VALUE OF RIGHT TERM (142)= 21013.849 THE VALUE OF LEFT TERM (142)= 21138.395 THE DIFFERANCE (142)= 124.546 B.L.E.V.E. SURFACE TEMPERATURE (142)= 1482.000 K.DEGREE

> THE VALUE OF RIGHT TERM (222)= 16889.008 THE VALUE OF LEFT TERM (222)= 17024.280 THE DIFFERANCE (222)= 135.272 B.L.E.V.E. SURFACE TEMPERATURE (222)= 1642.000 K.DEGREE

PROPANE FIREBALL SURFACE TEMPERATURE:

THE VALUE OF RIGHT TERM (221)= 14520.350 THE VALUE OF LEFT TERM (221)= 14677.600 THE DIFFERANCE (221)= 157.250 B.L.E.V.E. SURFACE TEMPERATURE (221)= 1740.000 K.DEGREE

> THE VALUE OF RIGHT TERM (210)= 13127.448 THE VALUE OF LEFT TERM (210)= 13248.904 THE DIFFERANCE (210)= 121.456 B.L.E.V.E. SURFACE TEMPERATURE (210)= 1818.000 K.DEGREE

THE VALUE OF RIGHT TERM (243)= 12196.181 THE VALUE OF LEFT TERM (243)= 12310.800 THE DIFFERANCE (243)= 114.619 B.L.E.V.E. SURFACE TEMPERATURE (243)= 1884.000 K.DEGREE

APPENDIX F

Computer program for evaluation of fireball surface temperature using thermodynamic equation

С C C С С PROGRAM FOR EVALUATION OF FIREBALL SURFACE TEMPERATURE С the set of the set of the set of the ******* С USING THERMODYNAMIC EQUATION: C ***** С C C С PRINT 15 15 FORMAT(5X, 'METHANE FIREBALL SURFACE TEMPERATURE : ', /, 5X, 42('=')//) С CALL TEMTHR(Q,G1,G2,G3) С C PRINT 17 FORMAT(5X, 'ETHANE FIREBALL SURFACE TEMPERATURE: ', /, 5X, 42('*')//) 17 С CALL TEMTHR(Q,G1,G2,G3) С C PRINT 20 FORMAT(5X, 'PROPANE FIREBALL SURFACE TEMPERATUE : ', /, 5X, 43('=')//) 20 C CALL TEMTHR(Q,G1,G2,G3) C C PRINT 30 FORMAT(5X, 'BUTANE FIREBALL SURFACE TEMPERATURE : ', /, 5X, 43('=')//) 30 С CALL TEMTHR(Q,G1,G2,G3) С С PRINT 35 FORMAT(5X, 'PENTANE FIREBALL SURFACE TEMPERATURE : ', /, 5X, 43('=')//) 35 С CALL TEMTHR(Q,G1,G2,G3) С С С C C STOP END С С C SUBROUTINE TEMTHR(Q,GN1,GN2,GN3) C C C
```
DIMENSION TS(400), T(400), CPN2(400), CPCO2(400),
      @CPO2(400), TSN(400), CPH2O(400), DIF(400)
 C
         READ *, GN1, GN2, GN3, GN4, Q, FC
 C
            Q1=Q-(GN3*10520.0)
 C
         | = 1
         TS(1)=1600.0
  5
         T(1)=TS(1)+273.0
        CPN2(1)=6.83+(0.0009*T(1))-(12000.0/(T(1)**2.0))
        CPCO2(1)=10.57+(0.0021*T(1))-(206000.0/(T(1)*2.0))
        CPH20(1)=7.30+(0.00246*T(1))
       CPO2(1)=7.16+(0.001*T(1))-(40000.0/(T(1)**2.0))
         TSN(1)=((FC*Q1)/((GN1*CPN2(1))+(GN2*CPCO2(1))+(GN3*CPH2O(1)))
      @+(GN4*CPO2(1)))+25.0
 C
C
        DIF(1)=TSN(1)-TS(1)
         IF (ABS(DIF(1)).GT.1.0) THEN
         TS(1+1)=TS(1)+1.0
          |=|+1
         GO TO 5
         ELSE
         PRINT 55, 1, TS(1)
   55
        FORMAT(5X, 'ASSUME SURFACE TEMPERATURE (', 13, ')=',
      @F10.3, ' C.DEGREE')
С
        PRINT 56, 1, CPN2(1)
  56
        FORMAT(5X, ' THE VALUE OF CPN2(', 13, ')=', F7.3)
. C
        PRINT 57, 1, CPCO2(1)
        FORMAT(5X, ' THE VALUE OF CPCO2 (', 13, ')=', F7.3)
  57
C
        PRINT 58, 1, CPH20(1)
         FORMAT(5X, ' THE VALUE OF CPH20 (', 13, ')=', F7.3)
  58
C
       PRINT 79, 1, CPO2(1)
       FORMAT(5X, 'THE VALUE OF CPO2(', 13, ')=', F7.3)
   79
C
        PRINT 100, 1, TSN(1)
        FORMAT(10X, ' SURFACE TEMPERATURE (', 13, ')=', F10.3,
  100
      @' C.DEGREE ')
       TSK=TSN(1)+273.0
       PRINT 150, I, TSK
  150 FORMAT(/, 10X, 'SURFACE TEMPERATURE OF FIRE BALL (', 13, ')=',
      @F10.3,' BY K.DEGREE',////)
          ENDIF
           RETURN
                END
```

THERMODYNAMIC FIREBALL SURFACE TEMPERATURE RESULTS

METHANE FIREBALL SURFACE TEMPERATURE :

ASSUME SURFACE TEMPERATURE (142)= 1741.000 C.DEGREE THE VALUE OF CPN2(142)= 8.640 THE VALUE OF CPC02 (142)= 14.749 THE VALUE OF CPH2O (142)= 12.254 THE VALUE OF CPO2(142)= 9.164 SURFACE TEMPERATURE (142)= 1741.468 C.DEGREE

SURFACE TEMPERATURE OF FIRE BALL (142)= 2014.468 BY K.DEGREE

ASSUME SURFACE TEMPERATURE (220)= 1819.000 C.DEGREE THE VALUE OF CPN2(220)= 8.710 THE VALUE OF CPC02 (220)= 14.916 THE VALUE OF CPH20 (220)= 12.446 THE VALUE OF CPO2(220)= 9.243 SURFACE TEMPERATURE (220)= 1819.953 C.DEGREE

SURFACE TEMPERATURE OF FIRE BALL (220)= 2092.953 BY K.DEGREE

PROPANE FIREBALL SURFACE TEMPERATUE :

ASSUME SURFACE TEMPERATURE (241)= 1840.000 C.DEGREE THE VALUE OF CPN2(241)= 8.729 THE VALUE OF CPC02 (241)= 14.961 THE VALUE OF CPH2O (241)= 12.498 THE VALUE OF CPO2(241)= 9.264 SURFACE TEMPERATURE (241)= 1840.337 C.DEGREE

SURFACE TEMPERATURE OF FIRE BALL (241) = 2113.337 BY K.DEGREE

BUTANE FIREBALL SURFACE TEMPERATURE :

ASSUME SURFACE TEMPERATURE (248)= 1847.000 C.DEGREE THE VALUE OF CPN2(248)= 8.735 THE VALUE OF CPC02 (248)= 14.976 THE VALUE OF CPH2O (248)= 12.515 THE VALUE OF CPO2(248)= 9.271 SURFACE TEMPERATURE (248)= 1847.308 C.DEGREE

SURFACE TEMPERATURE OF FIRE BALL (248)= 2120.308 BY K.DEGREE

PENTANE FIREBALL SURFACE TEMPERATURE :

ASSUME SURFACE TEMPERATURE (255)= 1854.000 C.DEGREE THE VALUE OF CPN2(255)= 8.742 THE VALUE OF CPC02 (255)= 14.991 THE VALUE OF CPH2O (255)= 12.532 THE VALUE OF CPO2(255)= 9.278 SURFACE TEMPERATURE (255)= 1854.224 C.DEGREE

SURFACE TEMPERATURE OF FIRE BALL (255)= 2127.224 BY K.DEGREE

CC	PROGRAM FOR EVALUATION OF BLEVE FIREBALL SURFACE TEMERATURE								
C C C C	US **	ING THERMODYNAMIC EQUATION: ************************************							
-		PRINT 15							
c ¹	15	FORMAT(5X, 'METHANE FIREBALL SURFACE TEMPERATURE : ', /, 5X, 42('=')//)							
с		CALL TEMTHR(Q,G1,G2,G3)							
С									
17	,	PRINT 17							
c		CALL TRATE FIREBALL SURFACE TEMPERATURE: ',/,5X,42('*')//)							
С		CALL TEMTHR(Q,G1,G2,G3)							
C		PRINT 20							
20 C)	FORMAT(5X, 'PROPANE FIREBALL SURFACE TEMPERATUE : ', /, 5X, 43('=')//)							
-		CALL TEMTHR(Q,G1,G2,G3)							
с с									
		PRINT 30							
30 C)	FORMAT(5X, 'BUTANE FIREBALL SURFACE TEMPERATURE : ', /, 5X, 43('=')//)							
с		CALL TEMTHR(Q,G1,G2,G3)							
С		DRINT of							
35	5	FORMAT(5X, 'PENTANE FIREBALL SURFACE TEMPERATURE : ', /, 5X, 43('=')//)							
с с		CALL TEMTHR(Q,G1,G2,G3)							
c									
c									
С									
С									
		STOP							
C		ENU							
č									
С									
		SUBROUTINE TEMTHR (Q, GN1, GN2, GN3)							
C									
C									
C		DIMENSION TS(999) T(999) CPN2(999) CPCC2(999)							
с	0	CPCH(999), CPO2(999), TSN(999), CPH2O(999), DIF(999)							
с	•	READ *, GN1, GN2, GN3, GN4, GN5, Q, FC							
с		READ *, A, B, C							

	Q1=Q-(GN3*10520.0)
С	
	I=1
	TS(1)=700.0
5	T(1)=TS(1)+273.0
	CPN2(1)=6.83+(0.0009*T(1))-(12000.0/(T(1))**2.0))
	$CPOO2(1) = 10.57 \pm (0.0021 \pm 7(1)) - (206000.0)((7(1) \pm 2.0))$
	$CPH20(1) = 7.30 \pm (0.00246 \pm 7(1))$
	$CPO2(1) = 7.16+(0.00240^{-1}(1))$
	CPCH(1) = 1.10+(0.001+1(1))=(4000.0/(1(1)+2.0))
	$C_{F}C_{F}(1) = A_{F}(B^{+}(1)) = (C^{+}(1(1)^{+}2.0))$
	SN(1) = ((FC + Q1)) / ((GN1 + CPN2(1)) + (GN2 + CPCO2(1)) + (GN3 + CPH2O(1)))
~	(GV4*CPO2(1))+(GN5*CPCH(1)))+25.0
č	
C	
	DIF(1)=ISN(1)-IS(1)
	IF (ABS(DIF(I)).GT.1.0) THEN
	1S(1+1)=TS(1)+1.0
	1=1+1
	GO TO 5
	ELSE
	PRINT 55, 1, TS(1)
55	FORMAT(5X, 'ASSUME SURFACE TEMPERATURE (', 13, ')=',
-	@F10.3, 'C.DEGREE')
C	
	PRINT 56, 1, CPN2(1)
56	FORMAT(5X, ' THE VALUE OF CPN2(', 13, ')=', F7.3)
C	
	PRINT 57, 1, CPCO2(1)
57	FORMAT(5X, ' THE VALUE OF CPCO2 (', 13, ')=', F7.3)
C	
	PRINT 58, I, CPH2O(I)
58	FORMAT(5X, ' THE VALUE OF CPH2O (', 13, ')=', F7.3)
С	
	PRINT 79, 1, CPO2(1)
79	FORMAT(5X, 'THE VALUE OF CPO2(', 13, ')=', F7.3)
С	
	PRINT 88, I, CPCH(I)
88	FORMAT(5X, 'THE VALUE OF CPCH(', 13, ')=', F8.3, /)
С	
	PRINT 100, I, TSN(I)
100	FORMAT(10X, ' SURFACE TEMPERATURE (', 13, ')=', F10.3,
	@'C.DEGREE ')
	TSK=TSN(1)+273.0
	PRINT 150, I, TSK
150	FORMAT(/, 10X, 'SURFACE TEMPERATURE OF FIRE BALL (', 13, ')='.
	@F10.3,' BY K.DEGREE',////)
	ENDIF
	RETURN
	END

THERMODYNAMIC BLEVE FIREBALL SURFACE TEMPERATURE RESULTS

METHANE FIREBALL SURFACE TEMPERATURE :

ASSUME SURFACE TEMPERATURE (235)= 934.000 C.DEGREE THE VALUE OF CPN2(235)= 7.908 THE VALUE OF CPC02 (235)= 12.963 THE VALUE OF CPH2O (235)= 10.269 THE VALUE OF CPC2(235)= 8.340 THE VALUE OF CPCH(235)= 18.896

SURFACE TEMPERATURE (235)= 934.654 C.DEGREE

SURFACE TEMPERATURE OF FIRE BALL (235)= 1207.654 BY K.DEGREE

ASSUME SURFACE TEMPERATURE (247)= 946.000 C.DEGREE THE VALUE OF CPN2(247)= 7.919 THE VALUE OF CPO02 (247)= 12.991 THE VALUE OF CPH20 (247)= 10.299 THE VALUE OF CPO2(247)= 8.352 THE VALUE OF CPCH(247)= 32.396

SURFACE TEMPERATURE (247)= 946.642 C.DEGREE

SURFACE TEMPERATURE OF FIRE BALL (247)= 1219.642 BY K.DEGREE

PROPANE FIREBALL SURFACE TEMPERATUE :

ASSUME SURFACE TEMPERATURE (248)= 947.000 C.DEGREE THE VALUE OF CPN2(248)= 7.920 THE VALUE OF CPC02 (248)= 12.994 THE VALUE OF CPH2O (248)= 10.301 THE VALUE OF CPO2(248)= 8.353 THE VALUE OF CPCH(248)= 46.092

SURFACE TEMPERATURE (248)= 947.490 C.DEGREE

SURFACE TEMPERATURE OF FIRE BALL (248)= 1220.490 BY K.DEGREE

BUTANE FIREBALL SURFACE TEMPERATURE :

ASSUME SURFACE TEMPERATURE (248)= 947.000 C.DEGREE THE VALUE OF CPN2(248)= 7.920 THE VALUE OF CPO02 (248)= 12.994 THE VALUE OF CPH20 (248)= 10.301 THE VALUE OF CPO2(248)= 8.353 THE VALUE OF CPCH(248)= 59.611

SURFACE TEMPERATURE (248)= 947.024 C.DEGREE

SURFACE TEMPERATURE OF FIRE BALL (248)= 1220.024 BY K.DEGREE

PENTANE FIREBALL SURFACE TEMPERATURE :

ASSUME SURFACE TEMPERATURE (248)= 947.000 C.DEGREE THE VALUE OF CPN2(248)= 7.920 THE VALUE OF CPC2 (248)= 12.994 THE VALUE OF CPH2O (248)= 10.301 THE VALUE OF CPC2(248)= 8.353 THE VALUE OF CPCH(248)= 73.100

SURFACE TEMPERATURE (248)= 947.514 C.DEGREE

SURFACE TEMPERATURE OF FIRE BALL (248)= 1220.514 BY K.DEGREE

APPENDIX G

Two computer programs for evaluation of

- i. View Factors, and the ratio of height to diameter of the fireball
- Fireball dimensions and hazard ranges using different authors' parameters, hot shell-cold core, isothermal and BLEVE models.

C С C C PROGRAM FOR EVALUATION OF VIEW FACTOR AND THE RATIO OF THE C = C HEIGHT TO DIAMETER OF THE FIREBALL: C _____ DIMENSION W(25) C READ *.N READ *, (W(1), I=1,N) С C PRINT 77 77 FORMAT(//12X, 'VIEW FACTOR AND (H/DS) RESULTS: ', @/12X,30('-')//) С С C PRINT 13 FORMAT(5X, 'FIREBALL RESULTS USING PREVIOUS WORK', 13 @' EQUATIONS & PARAMETERS: ', /5X, 60('=') //) С C С С CALCULATION OF FIREBALL DIMENSIONS BY USING PREVIOUS WORK. С 000 С C READ *, THETA, V, V2, V3, V4, V5, GN, GN2, GN3, GN4, GN5, FC, FC2, @FC3, FC4, FC5, WTM, WTM2, WTM3, WTM4, WTM5 C PRINT 19 FORMAT(10X, 'USING GAYLE PARAMETERS (A, AS, B & BS)', /, 10X, 37('=')//) 19 С READ *, SECMA, TEMS1, TEMS2, TEMS3, TEMS4, TEMS5 С C READ *, A, AS, B, BS, A2, AS2, B2, BS2, A3, AS3, B3, BS3, A4, AS4, B4, BS4 @, A5, AS5, B5, BS5 C CALL FIRBAL (N,W, THETA, V, V2, V3, GN, GN2, GN3, FC, FC2, FC3, WTM, WTM2 +,WTM3,A,A2,A3,AS,AS2,AS3,B,B2,B3,BS,BS2,BS3,A4,AS4,B4,BS4, @A5, AS5, B5, BS5, SECMA, V4, GN4, FC4, WTM4, V5, GN5, FC5, WTM5, @TEMS1, TEMS2, TEMS3, TEMS4, TEMS5) C C C PRINT 21 FORMAT(10X, 'USING BRASIE PARAMETERS (A, AS, B & BS)', /, 10X, 21 +40('=')//)

0	
	READ *, SECMA, TEMS1, TEMS2, TEMS3, TEMS4, TEMS5
C	
C	
-	PEAD * A AS P PS A2 AS2 P2 P22 A2 A22 P2 P22 A2 A22 P2 P22 A2 A22 P2 P22 P2
	A, A, A, B, B, B, A, A, A, B,
	@,A5,A55,B5,B55
	CALL FIRBAL(N,W, THETA, V, V2, V3, GN, GN2, GN3, FC, FC2, FC3, WTM, WTM2,
	@WTM3, A, A2, A3, AS, AS2, AS3, B, B2, B3, BS, BS2, BS3, A4, AS4, B4, BS4
	@A5, AS5, B5, BS5, SECMA, V4, GN4, FC4 WTM4, V5, GN5, FC5, WTM5
	@TEMS1. TEMS2. TEMS3. TEMS4. TEMS5.)
C	
č	
č	
C	
	PRINT 22
22	FORMAT(10X, 'USING MARSHALL PARAMETERS (A, AS, B & BS) . 1 (10X
	@40('=')//)
С	
-	PEAD & SEGMA TRACI TRACA TRACA TRACA TRACA
~	ALAD , SECURA, TEVIST, TEVISZ, TEVISZ, TEVISZ, TEVISZ, TEVISZ, TEVISZ,
C	
С	
	READ *, A, AS, B, BS, A2, AS2, B2, BS2, A3, AS3, B3, BS3, A4, AS4, B4, BS4
	CALL FIRBAL (N,W, THETA, V, V2, V3, GN, GN2, GN3, EC, EC2, EC3, WITM WITM2
	GWTM3, A, A2, A3, A5, A52, A53, B, B2, B3, B5, B52, B53, A4, A54, B4, B4, B4, B4, B4, B4, B4, B4, B4, B
	@45 AS5 B5 B5 SECAN VA COM ECA WITH VE COM STATUS
	a T5/61 T5/62, T
~	(, TEV51, TEV52, TEV53, TEV54, TEV55)
C	
С	
С	
	PRINT 23
23	FORMAT(10X, 'LSING BOBERTS PARAMETERS (A AS B & BS), (/ 10)
	@40('=')//)
C	
č	
C	
	READ *, SEOMA, TEMS1, TEMS2, TEMS3, TEMS4, TEMS5
С	
	READ *, A, AS, B, BS, A2, AS2, B2, BS2, A3, AS3, B3, BS3, A4, AS4, B4, B54
	@, A5, AS5, B5, BS5
	CALL ELEBRAL (N.W. THETA V/ V/2 V/2 CALCAR CAR AND TO THE THE AND
	WTM2 A A2 A2 A2 AC AC A2
	ewilws, A, A2, A3, A5, A52, AS3, B, B2, B3, BS, BS2, BS3, A4, AS4, B4, BS4,
	@A5,AS5,B5,BS5,SECMA,V4,GN4,FC4,WTM4,V5,GN5,FC5,WTM5,
	@TEMS1, TEMS2, TEMS3, TEMS4, TEMS5)
С	
С	
С	
-	PDINT 21
21	
31	FORWAT (TOX, USING HOT SHELL-COLD CORE MODEL: ', /, 10X,
	@35('=')//)
С	
С	
	PRINT 201
201	FORMAT (10X ' METHANE RESULTS OF LOT OUT !! ON D CORE .
201	at 10% decision and hesoers of hor shell-cold core MODEL:',
~	w/, 10,,40(*)/)
C	
C	

C	
~	CALL HOIMET (N,W, THETA)
C	
C	
	PRINT 203
203	FORMAT (//10X, ' BUTANE RESULTS OF HOT SHELL-COLD CORE MODEL: ',
	@/,10X,46('*')/)
С	
	CALL HOTSHE (N, W, THETA, SECMA)
С	
С	
С	
	PRINT 32
32	FORMAT(10X, 'USING ISO-THERMAL MODEL: ', /, 10X, 23('=')//)
С	
С	
С	
	PRINT 204
204	FORMAT (10X, ' METANE RESULTS OF ISO-THERMAL MODEL .'
	a/.10X.37('=')/)
C	
•	READ * SECMAS VS ONS WIMS ECS TEMES 1
	CALL ISOTHERN W THETA SECMAS VE ONE WINE ECE TEMOLIN
C	CALL 150 (11, 11, 11, 11, 520 (45, 15, 345, 11))
č	
C	DRINT 1999
1264	FRINI 1200
1200	PORVATION, ETHANE RESULTS OF ISU-THERMAL MODEL:
~	@/, IUX, 3/('=')/)
C	
~	READ *, SECWA26, V26, GN26, WIM26, FC26, TEMS26
C	
-	CALL ISOTHR(N,W, THE TA, SECMA26, V26, GN26, WTM26, FC26, TEMS26)
C	
C	
	PRINT 206
206	FORMAT(10X, ' PROPANE RESULTS OF ISO-THERMAL MODEL: ',
	@/,10X,39('=')/)
С	
	READ *, SECMA6, V6, GN6, WTM6, FC6, TEMS12
	CALL ISOTHR(N,W, THETA, SECMA6, V6, GN6, WTM6, FC6, TEMS12)
С	
С	
	PRINT 207
207	FORMAT (10X, ' BUTANE RESULTS OF ISO-THERMAL MODEL: '.
	@/,10X,38('=')/)
С	
	READ *, SECMA7, V7, GN7, WTM7, FC7, TEMS13
	CALL ISOTHR (N.W. THETA, SEGMAT, VT. GNT, WTMT, FCT, TEMS13)
C	
C	
	PRINT 208
208	FORMAT (10X ' PENTANE RESULTS OF ISO_THEDMAL MODDEL .'
200	a/ 10X 37('-')/)
	S7, 107, 57 (=)/)

С

READ *, SECMA8, V8, GN8, WTM8, FC8, TEMS14 CALL ISOTHR (N, W, THETA, SECMA8, V8, GN8, WTM8, FC8, TEMS14) C С C PRINT 901 901 FORMAT(///10X, ' B.L.E.V.E FIREBALL RESULTS: ', /, 10X, 30('*')//) С C С PRINT 902 FORMAT(10X, ' METHANE BLEVE FIREBALL RESULTS: ', /, 10X, 35('=')/) 902 С С READ *, SECMA, WTM, TEMS6 CALL BLEVE (N, W, THETA, SECMA, TEMS6, WTM) С **PRINT 1623** 1623 FORMAT(10X, 'ETHANE BLEVE FIREBALL RESULTS: ', /, 10X, 35('=')/) C READ *, SECMA, WTM, TEMS27 CALL BLEVE (N, W, THETA, SECMA, TEMS27, WTM) C C PRINT 903 FORMAT(10X, 'PROPANE BLEVE FIREBALL RESULTS: ',/, 10X, 35('=')/) 903 С READ *, SECMA, WTM, TEMS7 CALL BLEVE (N, W, THETA, SECMA, TEMS7, WTM) C C PRINT 904 FORMAT(10X, ' BUTANE BLEVE FIREBALL RESULTS: ',/10X,35('=')/) 904 C READ *, SECMA, WTM, TEMS8 CALL BLEVE (N, W, THETA, SECMA, TEMS8, WTM) C C PRINT 906 FORMAT(10X, ' PENTANE BLEVE FIREBALL RESULTS: ', /, 10X, 35('=')/) 906 C READ *, SECMA, WTM, TEMS9 CALL BLEVE (N,W, THETA, SECMA, TEMS9, WTM) С C STOP END С 00000 C

c	Q1,Q2PIG,TEMP.OF FIREBALL SURFACE, DISTANCE OF THE TARGETS, TAW, Q2.
C C	BY USING PREVIOUS WORK:
c	
C	
C	SUBROUTINE FIRBAL(II, W, THETA, V, V2, V3, GN, GN2, GN3, FC, FC2, FC3, +WTM, WTM2, WTM3, A1, A2, A3, AS1, AS2, AS3, B1, B2, B3, BS1, BS2, BS3, @A4, AS4, B4, BS4, A5, AS5, B5, BS5, SECMA, V4, GN4, FC4, WTM4, @V5, GN5, EC5, WTM5, TEMS1, TEMS2, TEMS2, TEMS4, TEMS1)
С	evo, evo, i co, windo, i evo i, i evoz, i evoz, i evoz, i evoz, i evoz, i evoz,
c	
C C	
C	
	DIMENSION Q1(10),DSH(10),F21(10),Y(10),DS(10), @Q2HLM(10),F21HLM(10),F21PIG(10),TC(10),F21B(10), @W(10),Q2PIG(10),F21PE(10)
C	
	TAW=1.0 QB=12.6
C	QPE=37.8
	CALCULATION OF METHANE FIREBALL DIMENSIONS & HAZARDS
11	PRINT 11
2	CONVERTIGA, METHANE FIREBALL RESULTS: ,/, 10X, 26('*')/)
2	
2	DO 5 1=1,11
	TC(1)=A1*(W(1)**AS1) DS(1)=B1*(W(1)**BS1)
	Y(1)=10*TC(1) Q1(1)=SEQMA*THETA*(TEMS1**4.0)
	Q2HLM(1)=60.53/(TC(1)**0.756) Q2PIG(1)=50.0/(TC(1)**0.71)

```
DSH(1)=Y(1)/DS(1)
         F21HUM(1)=Q2HUM(1)/(TAW*Q1(1))
         F21PIG(1)=Q2PIG(1)/(TAW*Q1(1))
         F21B(1)=QB/(TAW*Q1(1))
         F21PE(1)=QPE/(TAW*Q1(1))
   5
            CONTINUE
С
C
       PRINT 100
  100
      FORMAT(1X,
     @'
           MASS
                    TEMP
                             H/DS F21-HLMAN
                                               F21-ANIM F21-BUILD',
     @'
          F21-PROCESS',/,72('-'))
С
      DO 6 N=1, 11
      PRINT 105, W(N), TEMS1, DSH(N), F21HLM(N), F21PIG(N), F21B(N), F21PE(N)
 105
        FORMAT(2F9.2,5(1X,F9.5))
C
С
  6
         CONTINUE
С
C
C
C
C
   CALCULATION OF ETHANE FIREBALL DIMENSIONS & HAZARDS:
С
С
С
         PRINT 5000
           FORMAT(10X,///)
 5000
C
С
      PRINT 261
      FORMAT(10X, ' ETHANE FIREBALL RESULTS: ', /, 10X, 26('*')/)
 261
С
0000
C
        DO 51 J=1,11
       TC(J)=A2*(W(J)**AS2)
       DS(J)=B2*(W(J)**BS2)
       Y(J)=10*TC(J)
       Q1(J)=SECMA*THETA*(TEMS2**4.0)
       Q2HUM(J)=28.16/(TC(J)**0.735)
       Q2PIG(J)=50.0/(TC(J)**0.71)
       DSH(J)=Y(J)/DS(J)
       F21HUM(J)=Q2HUM(J)/(TAW*Q1(J))
       F21PIG(J)=Q2PIG(J)/(TAW*Q1(J))
       F21B(J)=QB/(TAW*Q1(J))
       F21PE(J)=QPE/(TAW*Q1(J))
C
  51
         CONTINUE
C
```

```
С
       PRINT 500
 500 FORMAT(1X,
      @'
           MASS
                    TEMP
                              H/DS
                                    F21-HUMAN F21-ANIM F21-BUILD',
      @'
          F21_PROCESS',/,1X,72('-'))
С
С
         DO 76 M=1,11
      PRINT 1051, W(M), TEMS2, DSH(M), F21HUM(M), F21PIG(M), F21B(M), F21PE(M)
 1051 FORMAT(2F9.2, 1X, 5(1X, F9.5))
C
  76
           CONTINUE
С
С
С
С
C
   CALCULATION OF PROPANE FIREBALL DIMENSIONS & HAZARDS:
C
C
   -
        PRINT 260
 260
        FORMAT(10X, ///)
С
        PRINT 150
 150
         FORMAT(10X, 'PROPANE FIREBALL RESULTS: ', /, 10X, 24('*')/)
С
С
С
        DO 10 M=1, 11
        TC(M)=A3*(W(M)**AS3)
       DS(M) = B3*(W(M)**BS3)
       Y(M)=10*TC(M)
C
      Q1(M)=SECMA*THETA*(TEMS3**4.0)
       Q2HUM(M)=28.16/(TC(M)**0.735)
       Q2PIG(M)=50.0/(TC(M)**0.71)
        DSH(M) = Y(M) / DS(M)
        F21HUM(M) = Q2HUM(M) / (TAW*Q1(M))
        F21PIG(M) = Q2PIG(M) / (TAW*Q1(M))
        F21B(M) = QB/(TAW*Q1(M))
        F21PE(M) = QPE/(TAW*Q1(M))
 10
        CONTINUE
С
С
C
       PRINT 200
     FORMAT(1X,
 200
     @'
           MASS
                    TEMP
                                      F21-HUMAN F21-ANIM F21-BULID',
                               H/DS
     @'
         F21-PROCESS',/,1X,72('-'))
        DO 15 MM=1, 11
       PRINT 210,W(MM), TEMS3, DSH(MM), F21HUM(MM), F21PIG(MM),
     @F21B(MM),F21PE(MM)
 210
          FORMAT(2F9.2,5(1X,F9.5))
  15
         CONTINUE
```

```
C
```

```
0000
     CALCULATION OF BUTANE FIREBALL DIMENSIONS & HAZARDS:
C
C
C
        PRINT 360
        FORMAT(2X, ///)
 360
C
         PRINT 255
 255
         FORMAT(10X, 'BUTANE FIREBALL RESULTS: ',/, 10X, 25('*')/)
С
00000
C
         DO 20 J=1,11
        TC(J)=A4*(W(J)**AS4)
        DS(J)=B4*(W(J)**BS4)
        Y(J) = 10.0*TC(J)
       Q1(J)=SECMA*THETA*(TEMS4**4.0)
         Q2HLM(J)=28.16/(TC(J)**0.735)
       Q2PIG(J)=50.0/(TC(J)**0.71)
          DSH(J)=Y(J)/DS(J)
          F21HUM(J)=Q2HUM(J)/(TAW*Q1(J))
          F21PIG(J)=Q2PIG(J)/(TAW*Q1(J))
          F21B(J)=QB/(TAW*Q1(J))
          F21PE(J)=QPE/(TAW*Q1(J))
 20
          CONTINUE
C
С
С
       PRINT 300
 300 FORMAT(1X,
     @'
           MASS
                   TEMP
                             H/DS
                                    F21-HUMAN F21-ANIM F21-BUILD ',
     @' F21-PROCESS',/1X,72('-'))
C
          DO 25 L=1,11
       PRINT 320,W(L), TEMS4, DSH(L), F21HUM(L), F21PIG(L),
     @F21B(L),F21PE(L)
  320
         FORMAT(2F9.2,5(1X,F9.5))
С
   25
          CONTINUE
C
C
С
         PRINT 811
 811
         FORMAT(5X, ///)
С
CC
```

```
C
   CALCULATION OF PENTANE FIREBALL DIMENSIONS & HAZARDS:
C
                     C
С
С
      PRINT 301
      FORMAT(10X, 'PENTANE FIREBALL RESULTS: ', /, 10X, 26('*')/)
 301
C
C
      DO 77 L=1,11
      TC(L) = A5*(W(L)**AS5)
      DS(L)=B5*(W(L)**BS5)
      Y(L)=10.0*TC(L)
       Q1(L)=SECMA*THETA*(TEMS5**4.0)
       Q2HLM(L)=28.16/(TC(L)**0.735)
      Q2PIG(L)=50.0/(TC(L)**0.71)
        DSH(L)=Y(L)/DS(L)
        F21HUM(L)=Q2HUM(L)/(TAW*Q1(L))
        F21PIG(L)=Q2PIG(L)/(TAW*Q1(L))
        F21B(L)=QB/(TAW*Q1(L))
        F21PE(L)=QPE/(TAW*Q1(L))
  77
        CONTINUE
С
C
C
      PRINT 303
 303 FORMAT(1X,
     @'
          MASS
                   TEMP
                           H/DS F21-HUMAN F21-ANIM F21-BUILD',
         F21-PROCESS', /, 1X, 72('-'))
     @'
С
С
C
     DO 27 K=1,11
       PRINT 306, W(K), TEMS5, DSH(K), F21HUM(K), F21PIG(K),
     @F21B(K),F21PE(K)
С
C
  306
          FORMAT(2F9.2,5(1X,F9.5))
C
  27
       CONTINUE
С
С
          PRINT 99
        FORMAT(10X,////)
  99
C
           RETURN
           END
C
C
C
C SUBROUTINE FOR METHANE ONLY (USING HOT SELL-COLD CORE MODEL):
C
          C
С
```

C

```
SUBROUTINE HOTMET (M, W, THETA)
      DIMENSION Q1(25), Y(25), DC(25), F21HLM(25), F21PIG(25), F21B(25),
     @TC(25),W(25),DO(25),Q2PIG(25),F21PE(25),DSH(25)
     @,Q2HUM(25)
C
C
         TAW=1.0
      SECMA=0.5
      QB=12.6
      QPE=37.8
C
        READ *, WTM, TEMS, ZH
C
C
      DO 140 I=1,M
      Q1(1)=SECMA*THETA*(TEMS**4.0)
      DO(1)=1.42*(W(1)**0.3334)
      DC(1)=D0(1)*((((273.0*WTM*ZH)/(44.8*290*Q1(1)))+1.0)**0.3334)
      TC(1)=(DC(1)-DO(1))/1.0
      Y(1)=1.0*TC(1)
       Q2HLM(1)=28.16/(TC(1)**0.735)
      Q2PIG(1)=50.0/(TC(1)**0.71)
      DSH(1)=Y(1)/DC(1)
        F21HUM(1)=Q2HUM(1)/(TAW*Q1(1))
        F21PIG(1)=Q2PIG(1)/(TAW*Q1(1))
С
        F21B(1)=QB/(TAW*Q1(1))
        F21PE(1)=QPE/(TAW*Q1(1))
 140
     CONTINUE
С
C
      PRINT 1421
 1421 FORMAT(1X,
     @'
         MASS
                                 F21-HUMAN F21-ANIM F21-BUILD',
                   TEMP
                           H/DS
     @'
         F21-PROCESS',/,1X,72('-'))
C
C
      DO 145 J=1,M
      PRINT 1400,W(J), TEMS, DSH(J), F21HLM(J), F21PIG(J),
     @F21B(J),F21PE(J)
 1400 FORMAT(2F9.2,5(1X,F9.5))
  145
         CONTINUE
C
C
C
        RETURN
         END
C
000
С
C
     SUBROUTINE PROGRAM FOR CALCULATION OF FIREBALLS
```

C C	DIMENSIONS BY USING HOT SHELL-COLD CORE MODEL:						
C C							
000	SUBROUTINE HOTSHE (MM, W, THETA, SECMA)						
c	DIMENSION Q1(25),Y(25),DC(25),F21HUM(25),F21PIG(25),F21B(25), @F21PE(25),DSH(25),TC(25),DO(25),Q2PIG(25),D1(25), @W(25),Q2HUM(25)						
000000	CALCULATION OF FIREBALL BY USING HOT SHELL-COLD CORE MODEL:						
c							
	TAW=1.0 QB=12.6						
	QPE=37.8						
C							
C	READ *, TEMS, WTM.C. RS. ZH						
С							
C							
c							
C							
С							
	F1=((1.0-(29.0/WTM))*(93.05*SECMA*((TEMS/1000.0)**4.0)))/(ZH) @-BS*C*(TEMS-290.0)-93.05*SECMA*((TEMS/1000.0)**4.0)))/(ZH)						
С	0 10 0 (1200-200.0)-33.00 SEQVA+((1205/1000.0)**4.0)*RS)						
С							
С	PRINT 222 E1						
333	FORMAT(/10X, 'THE VALUE OF E1 =' E10 2 ()						
С							
С							
С							
	PRINT 1222						
1222	2 FORMAT(//,5X, 'THIS MODEL SHOUD NOT USED IN THESE CONDITIONS'						
-	@/,5X, 'BECAUSE F1 VALUE IS NEGATIVE & IT MUST BE POSTIVE',//)						
C							
č							
	ELSE						
C							
C							
C	DO 40 K=1.MM						
	Q1(K)=SECMA*THETA*(TEMS**4.0)						

```
С
      D1(K)=(1.567*(1.0+(RS*F1))*W(K))*0.3334
      DO(K)=3.57*((W(K)/WTM)**0.3334)
      DC(K)=((((273.0*ZH*(1.0-F1))/(448.0*Q1(K)*(1.0+RS*F1)))
     @+1.0)**0.3334)*D1(K)
C
       TC(K) = (DC(K) - DO(K))/1.0
С
С
C
         Y(K)=10.0*TC(K)
        Q2HLM(K)=28.16/(TC(K)**0.735)
        Q2PIG(K)=50.0/(TC(K)**0.71)
             DSH(K)=Y(K)/DC(K)
        F21HUM(K)=Q2HUM(K)/(TAW*Q1(K))
        F21PIG(K)=Q2PIG(K)/(TAW*Q1(K))
        F21B(K)=QB/(TAW*Q1(K))
        F21PE(K)=QPE/(TAW*Q1(K))
С
 40
        CONTINUE
С
С
С
      PRINT 421
 421
      FORMAT(1X,
     0'
          MASS
                     TEMP H/DS
                                   F21-HUMAN F21-ANIM F21-BUILD',
         F21-PROCESS',/,1X,72('-'))
     @'
С
         DO 45 KI=1,MM
        PRINT 493, W(KI), TEMS, DSH(KI), F21HUM(KI), F21PIG(KI)
     @,F21B(KI),F21PE(KI)
 493
        FORMAT(2F9.2,5(1X,F9.5))
C
  45
        CONTINUE
C
С
С
      IF(DSH(1).GT.2.5) THEN
C
        PRINT 221, DSH(1)
        FORMAT(/10X, 'THE VALUE OF H/DS =', F10.3, /)
  221
C
C
С
      PRINT 1223
 1223 FORMAT(//, 5X, 'THE VALUE OF F21 BECOMES VERY SMALL AND THAT MEAN',
     @/,5X, 'THE VALUE OF F21 APPROACHES ZERO.',
     @/,5X, 'THEN THIS MODEL MUST NOT USED IN THESE CONDITIONS. ',///)
C
C
С
         ELSE
С
C
```

29	PRINT 29 FORMAT(3X,76('*')////)
С	
	ENDIF
C	
c	
	RETURN
с	
C	
c	
C C	SUBROUTINE PROGRAM FOR CALCULATION OF FIREBALL
C	BY USING ISO-THERMAL MODEL EQUATIONS:
C C	
С	
С	SUBROUTINE ISOTHR(LL,W, THETA, SECMA, V, GN, WTM, FC, TEMS)
C	
C	DIMENSION W(25), F21HUM(25), TC(25), DC(25), DO(25),
	@F21PIG(25),F21B(25),F21PE(25),Y(25),Q1(25),Q2PIG(25), @O2HIM(25),DSH(25)
с	
C C	
c	
C C	CALCULATION OF FIREBALL BY USING ISOTHERMAL MODEL .
с	
	IAW=1.0 QB=12.6
~	QPE=37.8
c	
C	
c	
	DO 50 L=1,LL DO(1) = 0.532 * ((220, 0***(1)) (*T)() * * 0.2020)
	DC(L)=0.539*((V+(GN+1.0)*FC)**0.3333)*((TEMS*W(L)/WTM)**0.3333)
c	TC(L)=0.45*(W(L)**0.3334)
-	Y(L)=10.0*TC(L)
	Q1(L)=SECMA*THETA*(TEMS**4.0) Q2HLM(L)=28.16/(TC(L))**0.735)
	Q2PIG(L)=50.0/(TC(L)**0.71)
	DSH(L)=Y(L)/DC(L) $F21HUM(L)=Q2HUM(L)/(TAW*Q1(L))$
	F21PIG(L)=Q2PIG(L)/(TAW*Q1(L))

```
F21B(L)=QB/(TAW*Q1(L))
        F21PE(L) = QPE/(TAW*Q1(L))
С
 50
         CONTINUE
С
C
      PRINT 752
 752 FORMAT(1X,
     @'
          MASS
                  TEMP
                           H/DC F21-HUMAN
                                              F21-ANIM F21-BUILD ',
     @'
        F21-PROCESS',/,1X,74('-'))
С
CC
      DO 55 N=1,LL
      PRINT 500, W(N), TEMS, DSH(N), F21HLM(N), F21PIG(N),
     @F21B(N),F21PE(N)
 500
       FORMAT(2F9.2,5(2X,F9.5))
 55
        CONTINUE
С
       PRINT 43
 43
       FORMAT(2X,/)
С
С
                RETURN
                END
С
C
CC
С
С
   SUBROTINE USED TOCALCULATE B.L.E.V.E. FIREBALL
С
     С
   DIMENSIONS AND PROPERTIES:
С
     -
CC
С
С
      SUBROUTINE BLEVE (M, W, THETA, SECMA, TEMS, WTM)
C
C
С
     DIMENSION W(25), F21HLM(25), TC(25), DC(25), F21PIG(25),
    @Y(25),Q1BL(25),F21B(25),F21PE(25),DSH(25),
    @Q2PIG(25),Q2HUM(25)
С
C
C
         TAW=1.0
      QB=12.6
      QPE=37.8
С
CC
```

```
С
С
      DO 83 N=1,M
      TC(N)=0.31*((44.8*W(N)/WTM)**0.3333)
      DC(N)=3.44*((44.8*W(N)/WTM)**0.3333)
C
      Y(N) = 0.5 * DC(N)
      Q1BL(N)=SECMA*THETA*(TEMS**4.0)
        Q2HLM(N)=28.16/(TC(N)**0.735)
        Q2PIG(N)=50.0/(TC(N)**0.71)
         DSH(N)=Y(N)/DC(N)
        F21HUM(N)=Q2HUM(N)/(TAW*Q1BL(N))
        F21PIG(N)=Q2PIG(N)/(TAW*Q1BL(N))
        F21B(N)=QB/(TAW*Q1BL(N))
        F21PE(N)=QPE/(TAW*Q1BL(N))
  83
       CONTINUE
С
C
      PRINT 703
 703 FORMAT(1X,
     @'
          MASS
                    TEMP
                             H/DC
                                      F21-HUMAN F21-ANIM F21-BUILD
     @' F21_PROCESS',/,1X,74('-'))
C
С
      DO 87 K=1,M
      PRINT 808, W(K), TEMS, DSH(K), F21HLM(K), F21PIG(K),
     @F21B(K),F21PE(K)
 808
         FORMAT(2F9.2,5(2X,F9.5))
С
 87
          CONTINUE
```

CC

000

819 C PRINT 819

FORMAT(2X,//)

RETURN

VIEW FACTOR AND (H/DS) RESULTS:

(FOR ISOTHERMAL FIREBALL TEMPERATURES)

FIREBALL RESULTS USING PREVIOUS WORK EQUATIONS & PARAMETERS:

USING GAYLE PARAMETERS (A, AS, B & BS)

METHANE FIREBALL RESULTS:

MASS	TEMP	H/DS F	21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2138.00	0.81906	0.02305	0.01999	0.01064	0.03191
5000.00	2138.00	0.85958	0.01495	0.01331	0.01064	0.03191
10000.00	2138.00	0.87764	0.01241	0.01117	0.01064	0.03191
25000.00	2138.00	0.90210	0.00969	0.00886	0.01064	0.03191
50000.00	2138.00	0.92106	0.00804	0.00744	0.01064	0.03191
75000.00	2138.00	0.93233	0.00721	0.00671	0.01064	0.03191
100000.00	2138.00	0.94041	0.00668	0.00624	0.01064	0.03191
500000.00	2138.00	0.98693	0.00433	0.00416	0.01064	0.03191

ETHANE FIREBALL RESULTS:

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2182.00	0.81906	0.01011	0.01842	0.00980	0.02941
5000.00	2182.00	0.85958	0.00663	0.01227	0.00980	0.02941
10000.00	2182.00	0.87764	0.00553	0.01029	0.00980	0.02941
25000.00	2182.00	0.90210	0.00435	0.00817	0.00980	0.02941
50000.00	2182.00	0.92106	0.00363	0.00685	0.00980	0.02941
75000.00	2182.00	0.93233	0.00327	0.00619	0.00980	0.02941
100000.00	2182.00	0.94041	0.00303	0.00575	0.00980	0.02941
500000.00	2182.00	0.98693	0.00199	0.00383	0.00980	0.02941

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BULID	F21-PROCESS
1000.00	2188.00	0.81906	0.01000	0.01822	0.00970	0.02909
5000.00	2188.00	0.85958	0.00656	0.01213	0.00970	0.02909
10000.00	2188.00	0.87764	0.00547	0.01018	0.00970	0.02909
25000.00	2188.00	0.90210	0.00431	0.00808	0.00970	0.02909
50000.00	2188.00	0.92106	0.00359	0.00678	0.00970	0.02909
75000.00	2188.00	0.93233	0.00323	0.00612	0.00970	0.02909
100000.00	2188.00	0.94041	0.00300	0.00569	0.00970	0.02909
500000.00	2188.00	0.98693	0.00197	0.00379	0.00970	0.02909

MASS	TEMP	H/DS I	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2188.00	0.81906	0.01000	0.01822	0.00970	0.02909
5000.00	2188.00	0.85958	0.00656	0.01213	0.00970	0.02909
10000.00	2188.00	0.87764	0.00547	0.01018	0.00970	0.02909
25000.00	2188.00	0.90210	0.00431	0.00808	0.00970	0.02909
50000.00	2188.00	0.92106	0.00359	0.00678	0.00970	0.02909
75000.00	2188.00	0.93233	0.00323	0.00612	0.00970	0.02909
100000.00	2188.00	0.94041	0.00300	0.00569	0.00970	0.02909
500000.00	2188.00	0.98693	0.00197	0.00379	0.00970	0.02909

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2190.00	0.81906	0.00996	0.01816	0.00966	0.02898
5000.00	2190.00	0.85958	0.00654	0.01209	0.00966	0.02898
10000.00	2190.00	0.87764	0.00545	0.01015	0.00966	0.02898
25000.00	2190.00	0.90210	0.00429	0.00805	0.00966	0.02898
50000.00	2190.00	0.92106	0.00358	0.00675	0.00966	0.02898
75000.00	2190.00	0.93233	0.00322	0.00610	0.00966	0.02898
100000.00	2190.00	0.94041	0.00298	0.00567	0.00966	0.02898
500000.00	2190.00	0.98693	0.00196	0.00377	0.00966	0.02898

USING BRASIE PARAMETERS (A,AS,B & BS)

METHANE FIREBALL RESULTS:

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2086.00	0.78947	0.02461	0.02138	0.01174	0.03521
5000.00	2086.00	0.78947	0.01641	0.01462	0.01174	0.03521
10000.00	2086.00	0.78947	0.01379	0.01241	0.01174	0.03521
25000.00	2086.00	0.78947	0.01095	0.00999	0.01174	0.03521
50000.00	2086.00	0.78947	0.00919	0.00848	0.01174	0.03521
75000.00	2086.00	0.78947	0.00830	0.00770	0.01174	0.03521
100000.00	2086.00	0.78947	0.00772	0.00720	0.01174	0.03521
500000.00	2086.00	0.78947	0.00515	0.00492	0.01174	0.03521

ETHANE FIREBALL RESULTS:

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2126.00	0.78947	0.01086	0.01982	0.01088	0.03263
5000.00	2126.00	0.78947	0.00732	0.01355	0.01088	0.03263
10000.00	2126.00	0.78947	0.00618	0.01150	0.01088	0.03263
25000.00	2126.00	0.78947	0.00494	0.00926	0.01088	0.03263
50000.00	2126.00	0.78947	0.00417	0.00786	0.01088	0.03263
75000.00	2126.00	0.78947	0.00377	0.00714	0.01088	0.03263
100000.00	2126.00	0.78947	0.00352	0.00667	0.01088	0.03263
500000.00	2126.00	0.78947	0.00237	0.00456	0.01088	0.03263

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BULID	F21-PROCESS
1000.00	2132.00	0.78947	0.01074	0.01960	0.01076	0.03227
5000.00	2132.00	0.78947	0.00724	0.01339	0.01076	0.03227
10000.00	2132.00	0.78947	0.00611	0.01137	0.01076	0.03227
25000.00	2132.00	0.78947	0.00488	0.00916	0.01076	0.03227
50000.00	2132.00	0.78947	0.00412	0.00777	0.01076	0.03227
75000.00	2132.00	0.78947	0.00373	0.00706	0.01076	0.03227
100000.00	2132.00	0.78947	0.00348	0.00660	0.01076	0.03227
500000.00	2132.00	0.78947	0.00235	0.00451	0.01076	0.03227

MASS TEMP H/DS F21-HLMAN F21-ANIM F21-BUILD	F21_DDOCESS
---	-------------

		the same party when your party when they want a	THE OWNER AND ADDRESS ADDRE	THE OWNER WHEN THE OWNER AND ADDRESS OF THE OWNER WHEN THE	the same want there want they are start and the same the		
1000.00	2132.00	0.78947	0.01074	0.01960	0.01076	0.03227	
5000.00	2132.00	0.78947	0.00724	0.01339	0.01076	0.03227	
10000.00	2132.00	0.78947	0.00611	0.01137	0.01076	0.03227	
25000.00	2132.00	0.78947	0.00488	0.00916	0.01076	0.03227	
50000.00	2132.00	0.78947	0.00412	0.00777	0.01076	0.03227	
75000.00	2132.00	0.78947	0.00373	0.00706	0.01076	0.03227	
100000.00	2132.00	0.78947	0.00348	0.00660	0.01076	0.03227	
500000.00	2132.00	0.78947	0.00235	0.00451	0.01076	0.03227	

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2134.00	0.78947	0.01070	0.01952	0.01072	0.03215
5000.00	2134.00	0.78947	0.00722	0.01334	0.01072	0.03215
10000.00	2134.00	0.78947	0.00609	0.01133	0.01072	0.03215
25000.00	2134.00	0.78947	0.00487	0.00912	0.01072	0.03215
50000.00	2134.00	0.78947	0.00411	0.00774	0.01072	0.03215
75000.00	2134.00	0.78947	0.00372	0.00703	0.01072	0.03215
100000.00	2134.00	0.78947	0.00347	0.00657	0.01072	0.03215
500000.00	2134.00	0.78947	0.00234	0.00449	0.01072	0.03215

USING MARSHALL PARAMETERS (A, AS, B & BS):

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2024.00	0.69091	0.02323	0.02040	0.01324	0.03973
5000.00	2024.00	0.69091	0.01549	0.01394	0.01324	0.03973
10000.00	2024.00	0.69091	0.01301	0.01184	0.01324	0.03973
25000.00	2024.00	0.69091	0.01033	0.00953	0.01324	0.03973
50000.00	2024.00	0.69091	0.00868	0.00809	0.01324	0.03973
75000.00	2024.00	0.69091	0.00783	0.00735	0.01324	0.03973
100000.00	2024.00	0.69091	0.00729	0.00687	0.01324	0.03973
500000.00	2024.00	0.69091	0.00486	0.00469	0.01324	0.03973

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2060.00	0.69091	0.01036	0.01901	0.01234	0.03702
5000.00	2060.00	0.69091	0.00698	0.01299	0.01234	0.03702
10000.00	2060.00	0.69091	0.00589	0.01103	0.01234	0.03702
25000.00	2060.00	0.69091	0.00471	0.00888	0.01234	0.03702
50000.00	2060.00	0.69091	0.00398	0.00754	0.01234	0.03702
75000.00	2060.00	0.69091	0.00360	0.00685	0.01234	0.03702
100000.00	2060.00	0.69091	0.00335	0.00640	0.01234	0.03702
500000.00	2060.00	0.69091	0.00226	0.00437	0.01234	0.03702

PROPANE FIREBALL RESULTS:

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BULID	F21-PROCESS
1000.00	2064.00	0.69091	0.01028	0.01886	0.01224	0.03673
5000.00	2064.00	0.69091	0.00693	0.01289	0.01224	0.03673
10000.00	2064.00	0.69091	0.00585	0.01094	0.01224	0.03673
25000.00	2064.00	0.69091	0.00467	0.00881	0.01224	0.03673
50000.00	2064.00	0.69091	0.00394	0.00748	0.01224	0.03673
75000.00	2064.00	0.69091	0.00357	0.00680	0.01224	0.03673
100000.00	2064.00	0.69091	0.00333	0.00635	0.01224	0.03673
500000.00	2064.00	0.69091	0.00225	0.00434	0.01224	0.03673

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS	
1000.00	2064.00	0.69091	0.01028	0.01886	0.01224	0.03673	-
5000.00	2064.00	0.69091	0.00693	0.01289	0.01224	0.03673	
10000.00	2064.00	0.69091	0.00585	0.01094	0.01224	0.03673	
25000.00	2064.00	0.69091	0.00467	0.00881	0.01224	0.03673	
50000.00	2064.00	0.69091	0.00394	0.00748	0.01224	0.03673	
75000.00	2064.00	0.69091	0.00357	0.00680	0.01224	0.03673	
100000.00	2064.00	0.69091	0.00333	0.00635	0.01224	0.03673	
500000.00	2064.00	0.69091	0.00225	0.00434	0.01224	0.03673	

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2066.00	0.78947	0.01218	0.02222	0.01220	0.03659

5000.00	2066.00	0.78947	0.00821	0.01519	0.01220	0.03659
10000.00	2066.00	0.78947	0.00693	0.01289	0.01220	0.03659
25000.00	2066.00	0.78947	0.00554	0.01038	0.01220	0.03659
50000.00	2066.00	0.78947	0.00467	0.00881	0.01220	0.03659
75000.00	2066.00	0.78947	0.00423	0.00801	0.01220	0.03659
100000.00	2066.00	0.78947	0.00395	0.00748	0.01220	0.03659
500000.00	2066.00	0.78947	0.00266	0.00511	0.01220	0.03659

USING ROBERTS PARAMETERS (A, AS, B & BS):

METHANE FIREBALL RESULTS:

MASS	TEMP	H/DS F	21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	1980.00	0.77586	0.02232	0.01975	0.01446	0.04338
5000.00	1980.00	0.77586	0.01488	0.01350	0.01446	0.04338
10000.00	1980.00	0.77586	0.01250	0.01146	0.01446	0.04338
25000.00	1980.00	0.77586	0.00992	0.00923	0.01446	0.04338
50000.00	1980.00	0.77586	0.00834	0.00783	0.01446	0.04338
75000.00	1980.00	0.77586	0.00753	0.00712	0.01446	0.04338
100000.00	1980.00	0.77586	0.00700	0.00665	0.01446	0.04338
500000.00	1980.00	0.77586	0.00467	0.00455	0.01446	0.04338

ETHANE FIREBALL RESULTS:

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2012.00	0.77586	0.01005	0.01853	0.01356	0.04068
5000.00	2012.00	0.77586	0.00678	0.01266	0.01356	0.04068
10000.00	2012.00	0.77586	0.00572	0.01075	0.01356	0.04068
25000.00	2012.00	0.77586	0.00457	0.00866	0.01356	0.04068
50000.00	2012.00	0.77586	0.00386	0.00735	0.01356	0.04068
75000.00	2012.00	0.77586	0.00349	0.00668	0.01356	0.04068
100000.00	2012.00	0.77586	0.00326	0.00624	0.01356	0.04068
500000.00	2012.00	0.77586	0.00220	0.00426	0.01356	0.04068

PROPANE FIREBALL RESULTS:

MASS TEMP H/DS F21-HUMAN F21-ANIM F21-BULI) F21_PROCESS
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					the second states where some states where the			
	1000.00	2016.00	0.77586	0.00997	0.01838	0.01345	0.04036	
	5000.00	2016.00	0.77586	0.00672	0.01256	0.01345	0.04036	
	10000.00	2016.00	0.77586	0.00567	0.01066	0.01345	0.04036	
	25000.00	2016.00	0.77586	0.00453	0.00859	0.01345	0.04036	
	50000.00	2016.00	0.77586	0.00383	0.00729	0.01345	0.04036	
	75000.00	2016.00	0.77586	0.00347	0.00662	0.01345	0.04036	
	100000.00	2016.00	0.77586	0.00323	0.00619	0.01345	0.04036	
5	500000.00	2016.00	0.77586	0.00218	0.00423	0.01345	0.04036	

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS	
1000.00	2016.00	0.77586	0.00997	0.01838	0.01345	0.04036	
5000.00	2016.00	0.77586	0.00672	0.01256	0.01345	0.04036	
10000.00	2016.00	0.77586	0.00567	0.01066	0.01345	0.04036	
25000.00	2016.00	0.77586	0.00453	0.00859	0.01345	0.04036	
50000.00	2016.00	0.77586	0.00383	0.00729	0.01345	0.04036	
75000.00	2016.00	0.77586	0.00347	0.00662	0.01345	0.04036	
100000.00	2016.00	0.77586	0.00323	0.00619	0.01345	0.04036	
500000.00	2016.00	0.77586	0.00218	0.00423	0.01345	0.04036	

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2016.00	0.77586	0.00997	0.01838	0.01345	0.04036
5000.00	2016.00	0.77586	0.00672	0.01256	0.01345	0.04036
10000.00	2016.00	0.77586	0.00567	0.01066	0.01345	0.04036
25000.00	2016.00	0.77586	0.00453	0.00859	0.01345	0.04036
50000.00	2016.00	0.77586	0.00383	0.00729	0.01345	0.04036
75000.00	2016.00	0.77586	0.00347	0.00662	0.01345	0.04036
100000.00	2016.00	0.77586	0.00323	0.00619	0.01345	0.04036
500000.00	2016.00	0.77586	0.00218	0.00423	0.01345	0.04036

USING HOT SHELL-COLD CORE MODEL:

METHANE RESULTS OF HOT SHELL-COLD CORE MODEL:

MASS	TEMP	H/DS F2	1-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS	
1000.00	1500.00	0.79645	0.01024	0.02010	0.08779	0.26337	-
5000.00	1500.00	0.79645	0.00690	0.01373	0.08779	0.26337	
10000.00	1500.00	0.79645	0.00582	0.01165	0.08779	0.26337	
25000.00	1500.00	0.79645	0.00465	0.00938	0.08779	0.26337	
50000.00	1500.00	0.79645	0.00392	0.00796	0.08779	0.26337	
75000.00	1500.00	0.79645	0.00355	0.00723	0.08779	0.26337	
100000.00	1500.00	0.79645	0.00331	0.00676	0.08779	0.26337	
500000.00	1500.00	0.79645	0.00223	0.00462	0.08779	0.26337	

BUTANE RESULTS OF HOT SHELL-COLD CORE MODEL:

THE VALUE OF F1 = 0.007

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS	
1000.00	1300.00	9.97053	0.00047	0.00102	0.07781	0.23342	-
5000.00	1300.00	9.98992	0.00014	0.00032	0.07781	0.23342	
10000.00	1300.00	9.99365	0.00009	0.00020	0.07781	0.23342	
25000.00	1300.00	9.99655	0.00004	0.00010	0.07781	0.23342	
50000.00	1300.00	9.99783	0.00003	0.00006	0.07781	0.23342	
75000.00	1300.00	9.99834	0.00002	0.00005	0.07781	0.23342	
100000.00	1300.00	9.99863	0.00002	0.00004	0.07781	0.23342	
500000.00	1300.00	9.99953	0.00000	0.00001	0.07781	0.23342	

THE VALUE OF H/DS = 9.971

THE VALUE OF F21 BECOMES VERY SMALL AND THAT MEAN THE VALUE OF F21 APPROACHES ZERO. THEN THIS MODEL MUST NOT USED IN THESE CONDITIONS.

USING ISO-THERMAL MODEL:

METANE RESULTS OF ISO-THERMAL MODEL:

MASS	TEMP	H/DC	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	1980.00	0.76761	0.01069	0.01972	0.01446	0.04338
5000.00	1980.00	0.76773	0.00721	0.01347	0.01446	0.04338
10000.00	1980.00	0.76779	0.00608	0.01143	0.01446	0.04338
25000.00	1980.00	0.76786	0.00486	0.00920	0.01446	0.04338
50000.00	1980.00	0.76791	0.00410	0.00781	0.01446	0.04338
75000.00	1980.00	0.76794	0.00371	0.00709	0.01446	0.04338

100000.00	1980.00	0.76796	0.00346	0.00663	0.01446	0.04338
500000.00	1980.00	0.76809	0.00233	0.00453	0.01446	0.04338

ETHANE RESULTS OF ISO-THERMAL MODEL:

MASS	TEMP	H/DC	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2012.00	0.78380	0.01003	0.01849	0.01356	0.04068
5000.00	2012.00	0.78393	0.00676	0.01263	0.01356	0.04068
10000.00	2012.00	0.78398	0.00570	0.01072	0.01356	0.04068
25000.00	2012.00	0.78405	0.00456	0.00863	0.01356	0.04068
50000.00	2012.00	0.78411	0.00385	0.00732	0.01356	0.04068
75000.00	2012.00	0.78414	0.00348	0.00665	0.01356	0.04068
100000.00	2012.00	0.78416	0.00324	0.00622	0.01356	0.04068
500000.00	2012.00	0.78429	0.00219	0.00425	0.01356	0.04068

PROPANE RESULTS OF ISO-THERMAL MODEL:

	MASS	TEMP	H/DC	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
	1000.00	2016.00	0.79137	0.00995	0.01834	0.01345	0.04036
	5000.00	2016.00	0.79150	0.00671	0.01253	0.01345	0.04036
	10000.00	2016.00	0.79155	0.00566	0.01064	0.01345	0.04036
	25000.00	2016.00	0.79163	0.00452	0.00856	0.01345	0.04036
	50000.00	2016.00	0.79168	0.00382	0.00727	0.01345	0.04036
	75000.00	2016.00	0.79171	0.00345	0.00660	0.01345	0.04036
1	00000.00	2016.00	0.79174	0.00322	0.00617	0.01345	0.04036
5	00000.00	2016.00	0.79186	0.00217	0.00421	0.01345	0.04036

BUTANE RESULTS OF ISO-THERMAL MODEL:

MASS	TEMP	H/DC	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2016.00	0.79566	0.00995	0.01834	0.01345	0.04036
5000.00	2016.00	0.79579	0.00671	0.01253	0.01345	0.04036
10000.00	2016.00	0.79584	0.00566	0.01064	0.01345	0.04036
25000.00	2016.00	0.79591	0.00452	0.00856	0.01345	0.04036
50000.00	2016.00	0.79597	0.00382	0.00727	0.01345	0.04036
75000.00	2016.00	0.79600	0.00345	0.00660	0.01345	0.04036
100000.00	2016.00	0.79602	0.00322	0.00617	0.01345	0.04036
500000.00	2016.00	0.79615	0.00217	0.00421	0.01345	0.04036

PENTANE RESULTS OF ISO-THERMAL MODDEL:

MASS	TEMP	H/DC	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS

1000.00	2016.00	0.79837	0.00995	0.01834	0.01345	0.04036
5000.00	2016.00	0.79850	0.00671	0.01253	0.01345	0.04036
10000.00	2016.00	0.79855	0.00566	0.01064	0.01345	0.04036
25000.00	2016.00	0.79863	0.00452	0.00856	0.01345	0.04036
50000.00	2016.00	0.79868	0.00382	0.00727	0.01345	0.04036
75000.00	2016.00	0.79871	0.00345	0.00660	0.01345	0.04036
100000.00	2016.00	0.79874	0.00322	0.00617	0.01345	0.04036
500000.00	2016.00	0.79887	0.00217	0.00421	0.01345	0.04036

METHANE BLEVE FIREBALL RESULTS:

 	10000	

MASS	TEMP	H/DC	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	1482.00	0.50000	0.03484	0.06418	0.04607	0.13820
5000.00	1482.00	0.50000	0.02349	0.04385	0.04607	0.13820
10000.00	1482.00	0.50000	0.01982	0.03722	0.04607	0,13820
25000.00	1482.00	0.50000	0.01583	0.02996	0.04607	0.13820
50000.00	1482.00	0.50000	0.01336	0.02543	0.04607	0.13820
75000.00	1482.00	0.50000	0.01210	0.02310	0.04607	0.13820
100000.00	1482.00	0.50000	0.01127	0.02158	0.04607	0.13820
500000.00	1482.00	0.50000	0.00760	0.01475	0.04607	0.13820

ETHANE BLEVE FIREBALL RESULTS:

MASS	TEMP	H/DC	F21-HLMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	1642.00	0.50000	0.02697	0.04942	0.03057	0.09171
5000.00	1642.00	0.50000	0.01818	0.03377	0.03057	0.09171
10000.00	1642.00	0.50000	0.01534	0.02866	0.03057	0.09171
25000.00	1642.00	0.50000	0.01226	0.02307	0.03057	0.09171
50000.00	1642.00	0.50000	0.01034	0.01958	0.03057	0.09171
75000.00	1642.00	0.50000	0.00936	0.01779	0.03057	0.09171
100000.00	1642.00	0.50000	0.00873	0.01662	0.03057	0.09171
500000.00	1642.00	0.50000	0.00588	0.01136	0.03057	0.09171

PROPANE BLEVE FIREBALL RESULTS:

TEMP	H/DC	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
	TEMP	TEMP H/DC	TEMP H/DC F21-HUMAN	TEMP H/DC F21-HUMAN F21-ANIM	TEMP H/DC F21-HUMAN F21-ANIM F21-BUILD

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1000.00	1740.00	0.50000	0.02349	0.04291	0.02424	0.07273
5000.00	1740.00	0.50000	0.01584	0.02932	0.02424	0.07273
10000.00	1740.00	0.50000	0.01336	0.02488	0.02424	0.07273
25000.00	1740.00	0.50000	0.01068	0.02003	0.02424	0.07273
50000.00	1740.00	0.50000	0.00901	0.01700	0.02424	0.07273
75000.00	1740.00	0.50000	0.00816	0.01545	0.02424	0.07273
100000.00	1740.00	0.50000	0.00760	0.01443	0.02424	0.07273
500000.00	1740.00	0.50000	0.00512	0.00986	0.02424	0.07273

BUTANE BLEVE FIREBALL RESULTS:

MASS	TEMP	H/DC	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	1818.00	0.50000	0.02109	0.03844	0.02034	0.06103
5000.00	1818.00	0.50000	0.01422	0.02626	0.02034	0.06103
10000.00	1818.00	0.50000	0.01200	0.02229	0.02034	0.06103
25000.00	1818.00	0.50000	0.00959	0.01794	0.02034	0.06103
50000.00	1818.00	0.50000	0.00809	0.01523	0.02034	0.06103
75000.00	1818.00	0.50000	0.00732	0.01384	0.02034	0.06103
100000.00	1818.00	0.50000	0.00683	0.01293	0.02034	0.06103
500000.00	1818.00	0.50000	0.00460	0.00883	0.02034	0.06103

PENTANE BLEVE FIREBALL RESULTS:

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MASS	TEMP	H/DC	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	1884.00	0.50000	0.01928	0.03508	0.01764	0.05292
5000.00	1884.00	0.50000	0.01300	0.02397	0.01764	0.05292
10000.00	1884.00	0.50000	0.01097	0.02034	0.01764	0.05292
25000.00	1884.00	0.50000	0.00876	0.01638	0.01764	0.05292
50000.00	1884.00	0.50000	0.00739	0.01390	0.01764	0.05292
75000.00	1884.00	0.50000	0.00670	0.01263	0.01764	0.05292
100000.00	1884.00	0.50000	0.00624	0.01180	0.01764	0.05292
500000.00	1884.00	0.50000	0.00421	0.00806	0.01764	0.05292

VIEW FACTOR AND (H/DS) RESULTS:

FIREBALL RESULTS USING PREVIOUS WORK EQUATIONS & PARAMETERS:

USING GAYLE PARAMETERS (A,AS,B & BS)

METHANE FIREBALL RESULTS:

MASS	TEMP	H/DS H	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2014.47	0.81906	0.02925	0.02536	0.01349	0.04048
5000.00	2014.47	0.85958	0.01897	0.01688	0.01349	0.04048
10000.00	2014.47	0.87764	0.01574	0.01417	0.01349	0.04048
25000.00	2014.47	0.90210	0.01230	0.01124	0.01349	0.04048
50000.00	2014.47	0.92106	0.01021	0.00943	0.01349	0.04048
75000.00	2014.47	0.93233	0.00915	0.00852	0.01349	0.04048
100000.00	2014.47	0.94041	0.00847	0.00792	0.01349	0.04048
500000.00	2014.47	0.98693	0.00549	0.00527	0.01349	0.04048

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2092.95	0.81906	0.01194	0.02177	0.01158	0.03474
5000.00	2092.95	0.85958	0.00784	0.01449	0.01158	0.03474
10000.00	2092.95	0.87764	0.00654	0.01216	0.01158	0.03474
25000.00	2092.95	0.90210	0.00514	0.00965	0.01158	0.03474
50000.00	2092.95	0.92106	0.00429	0.00810	0.01158	0.03474
75000.00	2092.95	0.93233	0.00386	0.00731	0.01158	0.03474
100000.00	2092.95	0.94041	0.00358	0.00680	0.01158	0.03474
500000.00	2092.95	0.98693	0.00235	0.00452	0.01158	0.03474

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BULID	F21-PROCESS
1000.00	2113.34	0.81906	0.01149	0.02094	0.01114	0.03342
5000.00	2113.34	0.85958	0.00754	0.01394	0.01114	0.03342
10000.00	2113.34	0.87764	0.00629	0.01170	0.01114	0.03342
25000.00	2113.34	0.90210	0.00495	0.00928	0.01114	0.03342
50000.00	2113.34	0.92106	0.00413	0.00779	0.01114	0.03342
75000.00	2113.34	0.93233	0.00371	0.00703	0.01114	0.03342
100000.00	2113.34	0.94041	0.00344	0.00654	0.01114	0.03342
500000.00	2113.34	0.98693	0.00226	0.00435	0.01114	0.03342

BUTANE FIREBALL RESULTS:

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS	
1000.00	2120.31	0.81906	0.01134	0.02066	0.01099	0.03298	-
5000.00	2120.31	0.85958	0.00744	0.01376	0.01099	0.03298	
10000.00	2120.31	0.87764	0.00621	0.01155	0.01099	0.03298	
25000.00	2120.31	0.90210	0.00488	0.00916	0.01099	0.03298	
50000.00	2120.31	0.92106	0.00407	0.00769	0.01099	0.03298	
75000.00	2120.31	0.93233	0.00366	0.00694	0.01099	0.03298	
100000.00	2120.31	0.94041	0.00340	0.00645	0.01099	0.03298	
500000.00	2120.31	0.98693	0.00223	0.00430	0.01099	0.03298	

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2127.22	0.81906	0.01119	0.02040	0.01085	0.03256
5000.00	2127.22	0.85958	0.00734	0.01358	0.01085	0.03256
10000.00	2127.22	0.87764	0.00613	0.01140	0.01085	0.03256
25000.00	2127.22	0.90210	0.00482	0.00904	0.01085	0.03256
50000.00	2127.22	0.92106	0.00402	0.00759	0.01085	0.03256
75000.00	2127.22	0.93233	0.00362	0.00685	0.01085	0.03256
100000.00	2127.22	0.94041	0.00335	0.00637	0.01085	0.03256
500000.00	2127.22	0.98693	0.00220	0.00424	0.01085	0.03256

USING BRASIE PARAMETERS (A,AS,B & BS)

METHANE FIREBALL RESULTS:

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2014.47	0.78947	0.02830	0.02459	0.01349	0.04048
5000.00	2014.47	0.78947	0.01887	0.01681	0.01349	0.04048
10000.00	2014.47	0.78947	0.01585	0.01426	0.01349	0.04048
25000.00	2014.47	0.78947	0.01259	0.01149	0.01349	0.04048
50000.00	2014.47	0.78947	0.01057	0.00975	0.01349	0.04048
75000.00	2014.47	0.78947	0.00954	0.00886	0.01349	0.04048
100000.00	2014.47	0.78947	0.00888	0.00828	0.01349	0.04048
500000.00	2014.47	0.78947	0.00592	0.00566	0.01349	0.04048

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2092.95	0.78947	7 0.01156	0.02110	0.01158	0.03474
5000.00	2092.95	0.78947	0.00780	0.01442	0.01158	0.03474
10000.00	2092.95	0.78947	0.00658	0.01224	0.01158	0.03474
25000.00	2092.95	0.78947	0.00526	0.00986	0.01158	0.03474
50000.00	2092.95	0.78947	0.00444	0.00837	0.01158	0.03474
75000.00	2092.95	0.78947	0.00402	0.00760	0.01158	0.03474
100000.00	2092.95	0.78947	0.00375	0.00710	0.01158	0.03474
500000.00	2092.95	0:78947	0.00253	0.00485	0.01158	0.03474

PROPANE FIREBALL RESULTS:

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BULID	F21-PROCESS
1000.00	2113.34	0.78947	0.01112	0.02030	0.01114	0.03342
5000.00	2113.34	0.78947	0.00750	0.01387	0.01114	0.03342
10000.00	2113.34	0.78947	0.00633	0.01178	0.01114	0.03342
25000.00	2113.34	0.78947	0.00506	0.00948	0.01114	0.03342
50000.00	2113.34	0.78947	0.00427	0.00805	0.01114	0.03342
75000.00	2113.34	0.78947	0.00387	0.00731	0.01114	0.03342
100000.00	2113.34	0.78947	0.00360	0.00683	0.01114	0.03342
500000.00	2113.34	0.78947	0.00243	0.00467	0.01114	0.03342

MASS TEMP H/DS F21-HUMAN F21-ANIM F21-BUILD F21-PROCESS

							the second
1000.00	2120.31	0.78947	0.01098	0.02003	0.01099	0.03298	
5000.00	2120.31	0.78947	0.00740	0.01369	0.01099	0.03298	
10000.00	2120.31	0.78947	0.00625	0.01162	0.01099	0.03298	
25000.00	2120.31	0.78947	0.00499	0.00936	0.01099	0.03298	
50000.00	2120.31	0.78947	0.00421	0.00794	0.01099	0.03298	
75000.00	2120.31	0.78947	0.00382	0.00722	0.01099	0.03298	
100000.00	2120.31	0.78947	0.00356	0.00674	0.01099	0.03298	
500000.00	2120.31	0.78947	0.00240	0.00461	0.01099	0.03298	

PENTANE FIREBALL RESULTS:

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2127.22	0.78947	0.01084	0.01977	0.01085	0.03256
5000.00	2127.22	0.78947	0.00731	0.01352	0.01085	0.03256
10000.00	2127.22	0.78947	0.00617	0.01147	0.01085	0.03256
25000.00	2127.22	0.78947	0.00493	0.00924	0.01085	0.03256
50000.00	2127.22	0.78947	0.00416	0.00784	0.01085	0.03256
75000.00	2127.22	0.78947	0.00377	0.00712	0.01085	0.03256
100000.00	2127.22	0.78947	0.00351	0.00666	0.01085	0.03256
500000.00	2127.22	0.78947	0.00237	0.00455	0.01085	0.03256

USING MARSHALL PARAMETERS (A, AS, B & BS):

METHANE FIREBALL RESULTS:

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2014.47	0.69091	0.02367	0.02079	0.01349	0.04048
5000.00	2014.47	0.69091	0.01578	0.01421	0.01349	0.04048
10000.00	2014.47	0.69091	0.01326	0.01206	0.01349	0.04048
25000.00	2014.47	0.69091	0.01053	0.00971	0.01349	0.04048
50000.00	2014.47	0.69091	0.00884	0.00824	0.01349	0.04048
75000.00	2014.47	0.69091	0.00798	0.00749	0.01349	0.04048
100000.00	2014.47	0.69091	0.00742	0.00700	0.01349	0.04048
500000.00	2014.47	0.69091	0.00495	0.00478	0.01349	0.04048

ETHANE FIREBALL RESULTS:

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2092.95	0.69091	0.00972	0.01784	0.01158	0.03474
5000.00	2092.95	0.69091	0.00655	0.01219	0.01158	0.03474
10000.00	2092.95	0.69091	0.00553	0.01035	0.01158	0.03474
25000.00	2092.95	0.69091	0.00442	0.00833	0.01158	0.03474
50000.00	2092.95	0.69091	0.00373	0.00707	0.01158	0.03474
75000.00	2092.95	0.69091	0.00338	0.00643	0.01158	0.03474
100000.00	2092.95	0.69091	0.00315	0.00601	0.01158	0.03474
500000.00	2092.95	0.69091	0.00212	0.00410	0.01158	0.03474

PROPANE FIREBALL RESULTS:

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BULID	F21-PROCESS
1000.00	2113.34	0.69091	0.00935	0.01716	0.01114	0.03342
5000.00	2113.34	0.69091	0.00631	0.01173	0.01114	0.03342
10000.00	2113.34	0.69091	0.00532	0.00996	0.01114	0.03342
25000.00	2113.34	0.69091	0.00425	0.00802	0.01114	0.03342
50000.00	2113.34	0.69091	0.00359	0.00681	0.01114	0.03342
75000.00	2113.34	0.69091	0.00325	0.00618	0.01114	0.03342
100000.00	2113.34	0.69091	0.00303	0.00578	0.01114	0.03342
500000.00	2113.34	0.69091	0.00204	0.00395	0.01114	0.03342

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS	
1000.00	2120.31	0.69091	0.00923	0.01694	0.01099	0.03298	-
5000.00	2120.31	0.69091	0.00622	0.01158	0.01099	0.03298	
10000.00	2120.31	0.69091	0.00525	0.00983	0.01099	0.03298	
25000.00	2120.31	0.69091	0.00420	0.00791	0.01099	0.03298	
50000.00	2120.31	0.69091	0.00354	0.00672	0.01099	0.03298	
75000.00	2120.31	0.69091	0.00321	0.00610	0.01099	0.03298	
100000.00	2120.31	0.69091	0.00299	0.00570	0.01099	0.03298	
500000.00	2120.31	0.69091	0.00202	0.00390	0.01099	0.03298	

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2127.22	0.78947	0.01084	0.01977	0.01085	0.03256

5000.00	2127.22	0.78947	0.00731	0.01352	0.01085	0.03256
10000.00	2127.22	0.78947	0.00617	0.01147	0.01085	0.03256
25000.00	2127.22	0.78947	0.00493	0.00924	0.01085	0.03256
50000.00	2127.22	0.78947	0.00416	0.00784	0.01085	0.03256
75000.00	2127.22	0.78947	0.00377	0.00712	0.01085	0.03256
100000.00	2127.22	0.78947	0.00351	0.00666	0.01085	0.03256
500000.00	2127.22	0.78947	0.00237	0.00455	0.01085	0.03256

USING ROBERTS PARAMETERS (A, AS, B & BS):

METHANE FIREBALL RESULTS:

MASS	TEMP	H/DS F	21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2014.47	0.77586	0.02083	0.01844	0.01349	0.04048
5000.00	2014.47	0.77586	0.01389	0.01260	0.01349	0.04048
10000.00	2014.47	0.77586	0.01167	0.01070	0.01349	0.04048
25000.00	2014.47	0.77586	0.00926	0.00861	0.01349	0.04048
50000.00	2014.47	0.77586	0.00778	0.00731	0.01349	0.04048
75000.00	2014.47	0.77586	0.00702	0.00664	0.01349	0.04048
100000.00	2014.47	0.77586	0.00653	0.00621	0.01349	0.04048
500000.00	2014.47	0.77586	0.00436	0.00424	0.01349	0.04048

ETHANE FIREBALL RESULTS:

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2092.95	0.77586	0.00858	0.01582	0.01158	0.03474
5000.00	2092.95	0.77586	0.00579	0.01081	0.01158	0.03474
10000.00	2092.95	0.77586	6 0.00489	0.00918	0.01158	0.03474
25000.00	2092.95	0.77586	0.00390	0.00739	0.01158	0.03474
50000.00	2092.95	0.77586	6 0.00329	0.00627	0.01158	0.03474
75000.00	2092.95	0.77586	0.00298	0.00570	0.01158	0.03474
100000.00	2092.95	0.77586	6 0.00278	0.00533	0.01158	0.03474
500000.00	2092.95	0.77586	0.00188	0.00364	0.01158	0.03474

PROPANE FIREBALL RESULTS:

MASS TEMP H/DS F21-HUMAN F21-ANIM F21-BULID F21-PROCESS

					tions which there were same times there which the	the same want when some state and state when	A REAL PROPERTY AND ADDRESS OF
1000.00	2113.34	0.77586	0.00826	0.01522	0.01114	0.03342	
5000.00	2113.34	0.77586	0.00557	0.01040	0.01114	0.03342	
10000.00	2113.34	0.77586	0.00470	0.00883	0.01114	0.03342	
25000.00	2113.34	0.77586	0.00376	0.00711	0.01114	0.03342	
50000.00	2113.34	0.77586	0.00317	0.00604	0.01114	0.03342	
75000.00	2113.34	0.77586	0.00287	0.00548	0.01114	0.03342	
100000.00	2113.34	0.77586	0.00267	0.00512	0.01114	0.03342	
500000.00	2113.34	0.77586	0.00180	0.00350	0.01114	0.03342	

BUTANE FIREBALL RESULTS:

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS	
1000.00	2120.31	0.77586	0.00815	0.01502	0.01099	0.03298	-
5000.00	2120.31	0.77586	0.00550	0.01027	0.01099	0.03298	
10000.00	2120.31	0.77586	0.00464	0.00872	0.01099	0.03298	
25000.00	2120.31	0.77586	0.00371	0.00702	0.01099	0.03298	
50000.00	2120.31	0.77586	0.00313	0.00596	0.01099	0.03298	
75000.00	2120.31	0.77586	0.00283	0.00541	0.01099	0.03298	
100000.00	2120.31	0.77586	0.00264	0.00506	0.01099	0.03298	
500000.00	2120.31	0.77586	0.00178	0.00346	0.01099	0.03298	

PENTANE FIREBALL RESULTS:

MASS	TEMP	H/DS	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2127.22	0.77586	0.00804	0.01483	0.01085	0.03256
5000.00	2127.22	0.77586	0.00542	0.01013	0.01085	0.03256
10000.00	2127.22	0.77586	0.00458	0.00860	0.01085	0.03256
25000.00	2127.22	0.77586	0.00366	0.00693	0.01085	0.03256
50000.00	2127.22	0.77586	0.00309	0.00588	0.01085	0.03256
75000.00	2127.22	0.77586	0.00280	0.00534	0.01085	0.03256
100000.00	2127.22	0.77586	0.00261	0.00499	0.01085	0.03256
500000.00	2127.22	0.77586	0.00176	0.00341	0.01085	0.03256

USING HOT SHELL-COLD CORE MODEL:

METHANE RESULTS OF HOT SHELL-COLD CORE MODEL:

MASS	TEMP	H/DS F2	1-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS	
1000.00	1500.00	0.79645	0.01024	0.02010	0.08779	0.26337	
5000.00	1500.00	0.79645	0.00690	0.01373	0.08779	0.26337	
10000.00	1500.00	0.79645	0.00582	0.01165	0.08779	0.26337	
25000.00	1500.00	0.79645	0.00465	0.00938	0.08779	0.26337	
50000.00	1500.00	0.79645	0.00392	0.00796	0.08779	0.26337	
75000.00	1500.00	0.79645	0.00355	0.00723	0.08779	0.26337	
100000.00	1500.00	0.79645	0.00331	0.00676	0.08779	0.26337	
500000.00	1500.00	0.79645	0.00223	0.00462	0.08779	0.26337	

BUTANE RESULTS OF HOT SHELL-COLD CORE MODEL:

THE VALUE OF F1 = 0.007

MASS	TEMP	H/DS F	21-HUMAN	F21-ANIM F	21-BUILD	F21-PROCESS	
1000.00	1300.00	9.97053	0.00047	0.00102	0.07781	0.23342	
5000.00	1300.00	9.98992	0.00014	0.00032	0.07781	0.23342	
10000.00	1300.00	9.99365	0.00009	0.00020	0.07781	0.23342	
25000.00	1300.00	9.99655	0.00004	0.00010	0.07781	0.23342	
50000.00	1300.00	9.99783	0.00003	0.00006	0.07781	0.23342	
75000.00	1300.00	9.99834	0.00002	0.00005	0.07781	0.23342	
100000.00	1300.00	9.99863	0.00002	0.00004	0.07781	0 23342	
500000.00	1300.00	9.99953	0.00000	0.00001	0.07781	0.23342	

THE VALUE OF H/DS = 9.971

THE VALUE OF F21 BECOMES VERY SWALL AND THAT MEAN THE VALUE OF F21 APPROACHES ZERO. THEN THIS MODEL MUST NOT USED IN THESE CONDITIONS.

USING ISO-THERMAL MODEL:

METANE RESULTS OF ISO-THERMAL MODEL:

MASS	TEMP	H/DC	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2014.47	0.76321	0.00998	0.01840	0.01349	0.04048
5000.00	2014.47	0.76333	0.00673	0.01257	0.01349	0.04048
10000.00	2014.47	0.76338	0.00568	0.01067	0.01349	0.04048
25000.00	2014.47	0.76345	0.00454	0.00859	0.01349	0.04048
50000.00	2014.47	0.76350	0.00383	0.00729	0.01349	0.04048
75000.00	2014.47	0.76354	0.00346	0.00662	0.01349	0.04048

100000.00	2014.47	0.76356	0.00323	0.00619	0.01349	0.04048
500000.00	2014.47	0.76368	0.00218	0.00423	0.01349	0.04048

ETHANE RESULTS OF ISO-THERMAL MODEL:

MASS	TEMP	H/DC	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2092.95	0.77356	0.00857	0.01579	0.01158	0.03474
5000.00	2092.95	0.77369	0.00577	0.01079	0.01158	0.03474
10000.00	2092.95	0.77374	0.00487	0.00916	0.01158	0.03474
25000.00	2092.95	0.77381	0.00389	0.00737	0.01158	0.03474
50000.00	2092.95	0.77386	0.00328	0.00626	0.01158	0.03474
75000.00	2092.95	0.77390	0.00297	0.00568	0.01158	0.03474
100000.00	2092.95	0.77392	0.00277	0.00531	0.01158	0.03474
500000.00	2092.95	0.77404	0.00187	0.00363	0.01158	0.03474

PROPANE RESULTS OF ISO-THERMAL MODEL:

MASS	TEMP	H/DC	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2113.34	0.77903	0.00824	0.01519	0.01114	0.03342
5000.00	2113.34	0.77916	0.00555	0.01038	0.01114	0.03342
10000.00	2113.34	0.77921	0.00469	0.00881	0.01114	0.03342
25000.00	2113.34	0.77928	0.00374	0.00709	0.01114	0.03342
50000.00	2113.34	0.77934	0.00316	0.00602	0.01114	0.03342
75000.00	2113.34	0.77937	0.00286	0.00547	0.01114	0.03342
100000.00	2113.34	0.77939	0.00267	0.00511	0.01114	0.03342
500000.00	2113.34	0.77952	0.00180	0.00349	0.01114	0.03342

BUTANE RESULTS OF ISO-THERMAL MODEL:

MASS	TEMP	H/DC	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	2120.31	0.78239	0.00813	0.01499	0.01099	0.03298
. 5000.00	2120.31	0.78252	0.00548	0.01024	0.01099	0.03298
10000.00	2120.31	0.78257	0.00463	0.00869	0.01099	0.03298
25000.00	2120.31	0.78264	0.00370	0.00700	0.01099	0.03298
50000.00	2120.31	0.78270	0.00312	0.00594	0.01099	0.03298
75000.00	2120.31	0.78273	0.00282	0.00540	0.01099	0.03298
100000.00	2120.31	0.78275	0.00263	0.00504	0.01099	0.03298
500000.00	2120.31	0.78288	0.00177	0.00344	0.01099	0.03298

PENTANE RESULTS OF ISO-THERMAL MODDEL:

MASS	TEMP	H/DC	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS

1000.00 2127.22 0.78421 0.00803 0.01480 0.01085	0.03256
5000.00 2127.22 0.78433 0.00541 0.01011 0.01085	0.03256
10000.00 2127.22 0.78439 0.00457 0.00858 0.01085	0.03256
25000.00 2127.22 0.78446 0.00365 0.00691 0.01085	0.03256
50000.00 2127.22 0.78451 0.00308 0.00586 0.01085	0.03256
75000.00 2127.22 0.78455 0.00279 0.00533 0.01085	0.03256
100000.00 2127.22 0.78457 0.00260 0.00497 0.01085	0.03256
500000.00 2127.22 0.78469 0.00175 0.00340 0.01085	0.03256

B.L.E.V.E FIREBALL RESULTS:

METHANE BLEVE FIREBALL RESULTS:

MASS	TEMP	H/DC	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	1207.65	0.50000	0.07901	0.14555	0,10448	0 31343
5000.00	1207.65	0.50000	0.05326	0.09945	0,10448	0 31343
10000.00	1207.65	0.50000	0.04495	0.08440	0,10448	0 31343
25000.00	1207.65	0.50000	0.03591	0.06795	0,10448	0.31343
50000.00	1207.65	0.50000	0.03030	0.05767	0.10448	0.31343
75000.00	1207.65	0.50000	0.02744	0.05239	0.10448	0 31343
100000.00	1207.65	0.50000	0.02557	0.04895	0,10448	0 31343
500000.00	1207.65	0.50000	0.01724	0.03344	0.10448	0.31343

ETHANE BLEVE FIREBALL RESULTS:

MASS	TEMP	H/DC	C F21-HUMAN F21-ANIN		F21-HUMAN F21-ANIN		F21-BUILD	F21-PROCESS
1000.00	1219.64	0.50000	0.08859	0.16235	0,10043	0.30129		
5000.00	1219.64	0.50000	0.05972	0.11093	0.10043	0.30129		
10000.00	1219.64	0.50000	0.05040	0.09415	0.10043	0.30129		
25000.00	1219.64	0.50000	0.04026	0.07579	0.10043	0.30129		
50000.00	1219.64	0.50000	0.03398	0.06433	0.10043	0.30129		
75000.00	1219.64	0.50000	0.03076	0.05844	0.10043	0.30129		
100000.00	1219.64	0.50000	0.02867	0.05460	0,10043	0.30129		
500000.00	1219.64	0.50000	0.01933	0.03730	0.10043	0.30129		

PROPANE BLEVE FIREBALL RESULTS:

MASS	TEMP	H/DC	F21-HI MAN	F21_ANUM	E21 PULLE	F21 0000500
MASS		HILL	F21-HUWAN	F21-ANIM	F21-BUILD	F21-PROCESS

1000.00	1220.49	0.50000	0.09703	0.17726	0.10015	0.30045
5000.00	1220.49	0.50000	0.06542	0.12111	0.10015	0.30045
10000.00	1220.49	0.50000	0.05520	0.10279	0.10015	0.30045
25000.00	1220.49	0.50000	0.04410	0.08275	0.10015	0.30045
50000.00	1220.49	0.50000	0.03721	0.07023	0.10015	0.30045
75000.00	1220.49	0.50000	0.03370	0.06381	0.10015	0.30045
100000.00	1220.49	0.50000	0.03140	0.05961	0.10015	0.30045
500000.00	1220.49	0.50000	0.02117	0.04073	0.10015	0.30045

BUTANE BLEVE FIREBALL RESULTS:

MASS	ASS TEMP H/DC F21-HUMAN F21-ANIM		F21-ANIM	F21-BUILD	JILD F21-PROCESS	
1000.00	1220.02	0.50000	0.10399	0.18952	0.10030	0.30091
5000.00	1220.02	0.50000	0.07010	0.12949	0.10030	0.30091
10000.00	1220.02	0.50000	0.05916	0.10990	0,10030	0.30091
25000.00	1220.02	0.50000	0.04726	0.08848	0.10030	0.30091
50000.00	1220.02	0.50000	0.03988	0.07509	0.10030	0.30091
75000.00	1220.02	0.50000	0.03611	0.06822	0.10030	0.30091
100000.00	1220.02	0.50000	0.03365	0.06373	0.10030	0.30091
500000.00	1220.02	0.50000	0.02269	0.04355	0.10030	0.30091

PENTANE BLEVE FIREBALL RESULTS:

MASS	TEMP	H/DC	F21-HUMAN	F21-ANIM	F21-BUILD	F21-PROCESS
1000.00	1220.51	0.50000	0.10947	0.19915	0,10014	0.30043
5000.00	1220.51	0.50000	0.07380	0.13607	0.10014	0.30043
10000.00	1220.51	0.50000	0.06227	0.11549	0.10014	0.30043
25000.00	1220.51	0.50000	0.04975	0.09298	0.10014	0.30043
50000.00	1220.51	0.50000	0.04198	0.07891	0.10014	0.30043
75000.00	1220.51	0.50000	0.03801	0.07169	0.10014	0.30043
100000.00	1220.51	0.50000	0.03543	0.06697	0.10014	0.30043
500000.00	1220.51	0.50000	0.02388	0.04576	0.10014	0.30043

00000 PROGRAM FOR EVALUATION OF THE FIREBALL DIAMETER AND HAZARD RANGE C -****** C USING AUTHOR'S PARAMETERS, HOTSHELL-COLD CORE , ISOTHERMAL & BLEVE MODELS: С C С C C DIMENSION W(10) С С С READ *,N READ *, (W(1), I=1, N)C PRINT 777 777 FORMAT(//12X, 'FIREBALL DIMENSIONS AND HAZARD RANGES RESULSTS: ', @/12X,48('=')//) С С С PRINT 13 FORMAT(5X, 'THE PREDICATION OF FIREBALL DIMENSIONS & '. 13 @'HAZARD RANGES', /, 5X, 55('*') /5X, 'USING PREVIOUS WORK', @' EQUATIONS & PARAMETERS: ', /5X, 44('*') //) С С С C C CALCULATION OF FIREBALL DIMENSIONS BY USING PREVIOUS WORK. С 000 С С READ *, THETA, V, V2, V3, V4, V5, GN, GN2, GN3, GN4, GN5, FC, FC2, @FC3, FC4, FC5, WTM, WTM2, WTM3, WTM4, WTM5 C PRINT 19 FORMAT(10X, 'USING GAYLE PARAMETERS (A, AS, B & BS)', /, 10X, 36('=')//) 19 C READ *, SECMA, TEMS1, TEMS2, TEMS3, TEMS4, TEMS5 С C READ *, A, AS, B, BS, A2, AS2, B2, BS2, A3, AS3, B3, BS3, A4, AS4, B4, BS4 @, A5, AS5, B5, BS5 C CALL FIRBAL (N,W, THETA, V, V2, V3, GN, GN2, GN3, FC, FC2, FC3, WTM, WTM2 +,WTM3,A,A2,A3,AS,AS2,AS3,B,B2,B3,BS,BS2,BS3,A4,AS4,B4,BS4,

```
@A5, AS5, B5, BS5, SECMA, V4, GN4, FC4, WTM4, V5, GN5, FC5, WTM5,
      @TEMS1, TEMS2, TEMS3, TEMS4, TEMS5)
C
С
С
       PRINT 21
   21
        FORMAT(10X, 'USING BRASIE PARAMETERS (A, AS, B & BS)', /, 10X,
      +40('=')//)
С
          READ *, SECMA, TEMS1, TEMS2, TEMS3, TEMS4, TEMS5
C
C
       READ *, A, AS, B, BS, A2, AS2, B2, BS2, A3, AS3, B3, BS3, A4, AS4, B4, BS4
      @, A5, AS5, B5, BS5
       CALL FIRBAL (N,W, THETA, V, V2, V3, GN, GN2, GN3, FC, FC2, FC3, WTM, WTM2,
      @WTM3, A, A2, A3, AS, AS2, AS3, B, B2, B3, BS, BS2, BS3, A4, AS4, B4, BS4,
      @A5, AS5, B5, BS5, SECMA, V4, GN4, FC4, WTM4, V5, GN5, FC5, WTM5,
      @TEMS1, TEMS2, TEMS3, TEMS4, TEMS5)
С
С
С
       PRINT 22
       FORMAT(10X, 'USING MARSHALL PARAMETERS (A, AS, B & BS): ', /, 10X,
 22
      @40('=')//)
C
          READ *, SECMA, TEMS1, TEMS2, TEMS3, TEMS4, TEMS5
C
С
       READ *, A, AS, B, BS, A2, AS2, B2, BS2, A3, AS3, B3, BS3, A4, AS4, B4, BS4
       CALL FIRBAL (N,W, THETA, V, V2, V3, GN, GN2, GN3, FC, FC2, FC3, WTM, WTM2,
      @WTM3, A, A2, A3, AS, AS2, AS3, B, B2, B3, BS, BS2, BS3, A4, AS4, B4, BS4,
      @A5, AS5, B5, BS5, SECMA, V4, GN4, FC4, WTM4, V5, GN5, FC5, WTM5
      @, TEMS1, TEMS2, TEMS3, TEMS4, TEMS5)
С
С
С
       PRINT 23
       FORMAT(10X, 'USING ROBERTS PARAMETERS (A, AS, B & BS): ', /, 10X,
 23
      @40('=')//)
C
C
         READ *, SECMA, TEMS1, TEMS2, TEMS3, TEMS4, TEMS5
С
       READ *, A, AS, B, BS, A2, AS2, B2, BS2, A3, AS3, B3, BS3, A4, AS4, B4, BS4
      @, A5, AS5, B5, BS5
       CALL FIRBAL(N,W, THETA, V, V2, V3, GN, GN2, GN3, FC, FC2, FC3, WTM, WTM2,
      GWTM3, A, A2, A3, AS, AS2, AS3, B, B2, B3, BS, BS2, BS3, A4, AS4, B4, BS4,
      @A5, AS5, B5, BS5, SECMA, V4, GN4, FC4, WTM4, V5, GN5, FC5, WTM5,
      @TEMS1, TEMS2, TEMS3, TEMS4, TEMS5)
C
C
       PRINT 31
       FORMAT(10X, 'USING HOT SHELL-COLD CORE MODEL: ', /, 10X,
 31
```

```
@35('=')//)
С
C
      PRINT 201
 201 FORMAT (10X, ' METHANE RESULTS OF HOT SHELL-COLD CORE MODEL: ',
     @/,10X,46('*')/)
C
С
С
       CALL HOTMET (N,W, THETA)
С
C
      PRINT 203
 203 FORMAT(10X, ' BUTANE RESULTS OF HOT SHELL-COLD CORE MODEL: ',
     @/,10X,46('*')/)
С
      CALL HOTSHE (N,W, THETA, SECMA)
C
С
C
      PRINT 32
  32 FORMAT(10X, 'USING ISO-THERMAL MODEL: ', /, 10X, 23('=')//)
С
С
С
      PRINT 204
 204 FORMAT(10X, ' METANE RESULTS OF ISO-THERMAL MODEL: ',
     @/,10X,37('=')/)
С
      READ *, SECMA5, V5, GN5, WTM5, FC5, TEMS11
      CALL ISOTHR(N,W, THETA, SECMA5, V5, GN5, WTM5, FC5, TEMS11)
С
C
      PRINT 2001
 2001 FORMAT(10X, 'ETHANE RESULTS OF ISO-THERMAL MODEL: ',
     @/,10X,39('=')/)
С
      READ *, SECMA26, V26, GN26, WTM26, FC26, TEMS26
      CALL ISOTHR(N,W,THETA,SECMA26,V26,GN26,WTM26,FC26,TEMS26)
C
C
      PRINT 206
 206 FORMAT(10X, ' PROPANE RESULTS OF ISO-THERMAL MODEL: ',
     @/,10X,39('=')/)
C
      READ *, SECMA6, V6, GN6, WTM6, FC6, TEMS12
      CALL ISOTHR (N, W, THETA, SECMA6, V6, GN6, WTM6, FC6, TEMS12)
C
C
      PRINT 207
     FORMAT (10X, ' BUTANE RESULTS OF ISO-THERMAL MODEL: ',
 207
     @/,10X,38('=')/)
C
      READ *, SECMA7, V7, GN7, WTM7, FC7, TEMS13
```

```
336
```

CALL ISOTHR (N, W, THETA, SECMA7, V7, GN7, WTM7, FC7, TEMS13) С C PRINT 208 208 FORMAT(10X, ' PENTANE RESULTS OF ISO-THERMAL MODDEL: ', @/,10X,37('=')/) C READ *, SECMA8, V8, GN8, WTM8, FC8, TEMS14 CALL ISOTHR (N, W, THETA, SECMA8, V8, GN8, WTM8, FC8, TEMS14) С С С PRINT 901 901 FORMAT(10X, ' B.L.E.V.E FIREBALL RESULTS: ', /, 10X, 30('*')//) С С С PRINT 902 902 FORMAT(10X, ' METHANE BLEVE FIREBALL RESULTS: ', /, 10X, 35('=')/) С С READ *, SECMA, WTM, TEMS6 CALL BLEVE (N, W, THETA, SECMA, TEMS6, WTM) С С PRINT 926 FORMAT(10X, 'ETHANE BLEVE FIREBALL RESULTS: ',/,10X,35('=')/) 926 C READ *, SECMA, WTM, TEMS26 CALL BLEVE (N, W, THETA, SECMA, TEMS26, WTM) C С PRINT 903 903 FORMAT(10X, 'PROPANE BLEVE FIREBALL RESULTS: ', /, 10X, 35('=')/) C READ *, SECMA, WTM, TEMS7 CALL BLEVE (N, W, THETA, SECMA, TEMS7, WTM) C С PRINT 904 FORMAT(10X, ' BUTANE BLEVE FIREBALL RESULTS: ', /10X, 35('=')/) 904 C READ *, SECMA, WTM, TEMS8 CALL BLEVE (N, W, THETA, SECMA, TEMS8, WTM) С C PRINT 906 FORMAT(10X, ' PENTANE BLEVE FIREBALL RESULTS: ', /, 10X, 35('=')/) 906 C READ *, SECMA, WTM, TEMS9 CALL BLEVE (N, W, THETA, SECMA, TEMS9, WTM) 913 FORMAT(2X, 100('*')) С С



```
С
C
C
C
```

READ *, (RTAB(MM), MM=1, II)READ *, (RTAPE(KK), KK=1, 1)

```
DO 5 |=1, ||
        TC(1)=A1*(W(1)**AS1)
        DS(1)=B1*(W(1)**BS1)
        Y(1)=10*TC(1)
       RHUM(1)=DS(1)*RTAHUM(1)
      RPIG(1)=DS(1)*RTAPIG(1)
       RB(1)=DS(1)*RTAB(1)
       RPE(1)=DS(1)*RTAPE(1)
       Q1(1)=SECMA*THETA*(TEMS1**4.0)
       Q2HUM(1)=28.16/(TC(1)**0.735)
        Q2PIG(1)=50.0/(TC(1)**0.71)
        F21(1)=((Y(1)/DS(1))+0.5)/(4.0*(((Y(1)/DS(1)+0.5)**2.0)))
     @+(RHLM(1)/DS(1))**2.0)**1.5)
       X(1)=DS(1)*((((((((((((((()/DS(1)+0.5)**2.0)+(RHUM(1)/DS(1))**2.0))
     @**0.5)-0.5)
        TAW(1) = EXP(-0.0007 * X(1))
   5
           CONTINUE
С
       PRINT 1001, TEMS1
 1001 FORMAT(/10X, 'AT FIREBALL SURFACE TEMPERATUE =', F10.3, /10X,
     @30('-')/)
С
C
      PRINT 100
 100 FORMAT(1X,
     Q'
          MASS
                   TIME
                           DIAMETER HEIGHT HUMAN R. ANIMAL R.',
     @'BULID.R. PR.EQ.R.',
     @/1X,74('-'))
C
      DO 6 N=1,11
      PRINT 105, W(N), TC(N), DS(N), Y(N), RHUM(N), RPIG(N), RB(N), RPE(N)
 105
        FORMAT(8F9.2)
С
С
  6
        CONTINUE
C
C
C
С
С
   CALCULATION OF ETHANE FIREBALL DIMENSIONS & HAZARDS:
С
                         _____
C
        PRINT 3333
 3333
        FORMAT(10X,///)
C
С
      PRINT 261
 261 FORMAT(10X, ' ETHANE FIREBALL RESULTS: ', /, 10X, 26( '*')/)
```

```
C
С
C
```

```
READ *, (RTAHLM(JJ), JJ=1, II)
       READ *, (RTAPIG(KK), KK=1, 11)
       READ *, (RTAB(MM), MM=1, II)
        READ *, (RTAPE(NN), NN=1, 11)
С
С
С
        DO 51 J=1,11
        TC(J)=A2*(W(J)**AS2)
       DS(J)=B2*(W(J)**BS2)
       Y(J)=10*TC(J)
       RHUM(J)=DS(J)*RTAHUM(J)
       RPIG(J)=DS(J)*RTAPIG(J)
       RB(J)=DS(J)*RTAB(J)
       RPE(J)=DS(J)*RTAPE(J)
       Q1(J)=SECMA*THETA*(TEMS2**4.0)
       Q2HLM(J)=28.16/(TC(J)**0.735)
       Q2PIG(J)=50.0/(TC(J)**0.71)
       F21(J)=((Y(J)/DS(J))+0.5)/(4.0*(((Y(J)/DS(J)+0.5)**2.0)))
     @+(RHLM(J)/DS(J))**2.0)**1.5)
      X(J)=DS(J)*((((((Y(J)/DS(J)+0.5)**2.0)+(RHUM(J)/DS(J))**2.0)))
     @**0.5)-0.5)
C
       TAW(J) = EXP(-0.0007 * X(J))
C
  51
         CONTINUE
С
        PRINT 1002, TEMS2
 1002
        FORMAT(/10X, 'FIREBALL SURFACE TEMPERATUER =', F10.3, /10X,
     @30('-')/)
С
C
C
      PRINT 500
 500 FORMAT(1X,
                   TIME
     @'
          MASS
                          DIAMETER HEIGHT
                                              HUMAN R. ANIMAL R. ',
     @'BULID.R. PR.EQ.R',
     @/,1X,74('-'))
C
C
        DO 76 M=1,11
      PRINT 1051, W(M), TC(M), DS(M), Y(M), RHUM(M), RPIG(M), RB(M), RPE(M)
 1051 FORMAT(8F9.2)
C
  76
          CONTINUE
C
000
C
С
   CALCULATION OF PROPANE FIREBALL DIMENSIONS & HAZARDS:
```

```
C
   -
        PRINT 260
 260
        FORMAT(10X, ///)
С
        PRINT 150
 150
         FORMAT(10X, 'PROPANE FIREBALL RESULTS: ', /, 10X, 24('*'))
C
          READ *, (RTAHLM(MM), MM=1, 11)
        READ *, (RTAPIG(LL), LL=1, II)
        READ *, (RTAB(L), L=1, II)
         READ *, (RTAPE(K), K=1, 11)
С
C
       DO 10 M=1,11
        TC(M) = A3*(W(M) * * AS3)
       DS(M)=B3*(W(M)**BS3)
       Y(M) = 10*TC(M)
       RHLM(M)=DS(M)*RTAHLM(M)
       RPIG(M)=DS(M)*RTAPIG(M)
      RB(M) = DS(M) * RTAB(M)
      RPE(M)=DS(M)*RTAPE(M)
C
      Q1(M)=SECMA*THETA*(TEMS3**4.0)
       Q2HLM(M)=28.16/(TC(M)**0.735)
       Q2PIG(M)=50.0/(TC(M)**0.71)
      F21(M) = ((Y(M)/DS(M))+0.5)/(4.0*(((Y(M)/DS(M)+0.5)**2.0)))
     @+(RHLM(M)/DS(M))**2.0)**1.5)
      X(M) = DS(M) * (((((Y(M)/DS(M)+0.5)**2.0)+(RHUM(M)/DS(M))**2.0)))
     @**0.5)-0.5)
      TAW(M) = EXP(-0.0007*X(M))
 10
        CONTINUE
C
      PRINT 1003. TEMS3
 1003 FORMAT(/10X, 'FIREBALL SURFACE TEMPERATURE =', F10.3, /10X,
     @30('-')/)
С
C
С
C
       PRINT 200
 200 FORMAT(1X,
         MASS
     @'
                  TIME
                          DIAMETER
                                      HEIGHT HUMAN R. ANIMAL R. ',
     @'BULID.R. PR.EQ.R',
     @/1X,74('-'))
        DO 15 MM=1,11
       PRINT 210,W(MM),TC(MM),DS(MM),Y(MM),RHLM(MM),RPIG(MM),
     @RB(MM), RPE(MM)
 210
         FORMAT(8F9.2)
  15
         CONTINUE
С
С
C
C
    CALCULATION OF BUTANE FIREBALL DIMENSIONS & HAZARDS:
C
```

```
С
C
C
        PRINT 360
        FORMAT(2X, ///)
 360
С
         PRINT 255
         FORMAT(10X, 'BUTANE FIREBALL RESULTS: ', /, 10X, 25('*'))
 255
C
C
          READ *, (RTAHLM(JJ), JJ=1, II)
           READ *, (RTAPIG(MM), MM=1, 11)
        READ *, (RTAB(LL), LL=1, II)
        READ *, (RTAPE(KK), KK=1, 11)
C
000
C
        DO 20 J=1,11
        TC(J) = A4*(W(J)**AS4)
        DS(J) = B4*(W(J)**BS4)
        Y(J) = 10.0*TC(J)
      Q1(J)=SECMA*THETA*(TEMS4**4.0)
       RHUM(J)=DS(J)*RTAHUM(J)
       RPIG(J)=DS(J)*RTAPIG(J)
      RB(J)=DS(J)*RTAB(J)
      RPE(J)=DS(J)*RTAPE(J)
        Q2HUM(J)=28.16/(TC(J)**0.735)
       Q2PIG(J)=50.0/(TC(J)**0.71)
       F21(J)=((Y(J)/DS(J)+0.5)/(4.0*(((Y(J)/DS(J)+0.5)**2.0))))
     @+(RHLM(J)/DS(J)+0.5))**2.0)**1.5)
      X(J)=DS(J)*((((((Y(J)/DS(J)+0.5)**2.0)+(RHUM(J)/DS(J))**2.0)))
     @**0.5)-0.5)
      TAW(J) = EXP(-0.0007 * X(J))
 20
         CONTINUE
C
      PRINT 1004, TEMS4
 1004 FORMAT(/10X, 'FIREBALL SURFACE TEMPERATURE =', F10.3, /10X,
     @30('-')/)
С
C
C
C
       PRINT 300
 300 FORMAT(1X.
     @'
          MASS
                  TIME
                          DIAMETER
                                      HEIGHT HUMAN R. ANIMAL R.
     @'BULID.R. PR.EQ.R',
     @/1X,74('-'))
С
         DO 25 L=1,11
       PRINT 320,W(L),TC(L),DS(L),Y(L),RHUM(L),RPIG(L),RB(L),RPE(L)
  320
         FORMAT(8F9.2)
C
```

```
342
```

```
25
          CONTINUE
C
С
С
         PRINT 811
 811
         FORMAT(5X, ///)
C
С
C
C
   CALCULATION OF PENTANE FIREBALL DIMENSIONS & HAZARDS:
C
С
        READ *, (RTAHLM(1), 1=1, 11)
        READ *, (RTAPIG(KK), KK=1, 11)
        READ *, (RTAB(J), J=1, II)
        READ *, (RTAPE(K), K=1, 11)
C
C
      PRINT 301
      FORMAT(10X, 'PENTANE FIREBALL RESULTS: ', /, 10X, 26('*')/)
 301
С
C
      DO 77 L=1,11
      TC(L)=A5*(W(L)**AS5)
      DS(L)=B5*(W(L)**BS5)
      Y(L) = 10.0*TC(L)
       Q1(L)=SECMA*THETA*(TEMS5**4.0)
      RHUM(L)=DS(L)*RTAHUM(L)
      RPIG(L)=DS(L)*RTAPIG(L)
      RB(L)=DS(L)*RTAB(L)
      RPE(L)=DS(L)*RTAPE(L)
       Q2HUM(L)=28.16/(TC(L)**0.735)
      Q2PIG(L)=50.0/(TC(L)**0.71)
      F21(L)=((Y(L)/DS(L)+0.5)/(4.0*(((Y(L)/DS(L)+0.5)**2.0))))
     @+(RHUM(L)/DS(L)+0.5))**2.0)**1.5)
     @**0.5)-0.5)
      TAW(L) = EXP(-0.0007 * X(L))
  77
        CONTINUE
C
      PRINT 1005, TEMS5
 1005 FORMAT(/10X, 'FIREBALL SURFACE TEMPERATURE =', F10.3, /10X,
     @30('-')/)
C
С
C
C
     PRINT 303
 303 FORMAT(1X,
    @'
         MASS
                  TIME
                          DIAMETER HEIGHT
                                             HUMAN R. ANIMAL R. ',
    @'BULID.R. PR.EQ.R',
     @/1X,74('-'))
C
C
```

```
DO 27 K=1,11
       PRINT 306,W(K),TC(K),DS(K),Y(K),RHUM(K),RPIG(K),RB(K),RPE(K)
С
C
  306
         FORMAT(8F9.2)
C
       CONTINUE
  27
C
C
          PRINT 99
  99
        FORMAT(10X, ////)
C
           RETURN
            END
С
С
C
C SUBROUTINE FOR METHANE ONLY (USING HOT SHELL-COLD CORE MODEL):
C
С
С
      SUBROUTINE HOTMET (M, W, THETA)
     DIMENSION Q1(25), Y(25), DC(25), RTAPE(25), RTAB(25), RB(25), RPE(25),
     @TC(25),W(25),D0(25),Q2PIG(25),RHLM(25),F21(25),X(25),
     @Q2HUM(25), RPIG(25), RTAPIG(25), RTAHUM(25), TAW(25), DSH(25)
С
С
     SECMA=0.5
     QB=12.6
     QPE=37.8
С
           READ *, WTM, TEMS, ZH
C
C
     READ *, (RTAHLM(J), J=1,M)
     READ *, (RTAPIG(K), K=1, M)
     READ *, (RTAB(1), I=1,M)
     READ *, (RTAPE(N), N=1,M)
C
C
     DO 140 I=1,M
     Q1(1)=SECMA*THETA*(TEMS**4.0)
     DO(1)=1.42*(W(1)**0.3334)
     DC(1)=D0(1)*((((273.0*WTM*ZH)/(44.8*290*Q1(1)))+1.0)**0.3334)
     TC(1)=(DC(1)-DO(1))/1.0
     Y(1)=1.0*TC(1)
     RHUM(1)=DC(1)*RTAHUM(1)
     RPIG(1)=DC(1)*RTAPIG(1)
     RB(1)=DC(1)*RTAB(1)
     RPE(1)=DC(1)*RTAPE(1)
      Q2HUM(1)=28.16/(TC(1)**0.735)
     Q2PIG(1)=50.0/(TC(1)**0.71)
     F21(1)=((Y(1)/DC(1)+0.5)/(4.0*(((Y(1)/DC(1)+0.5)**2.0))))
```

```
@+(RHLM(1)/DC(1)+0.5))**2.0)**1.5)
      X(1)=DC(1)*(((((Y(1)/DC(1)+0.5)**2.0)+(RHUM(1)/DC(1)+0.5)**2.0)))
     @**0.5)-0.5)
      TAW(1) = EXP(-0.0007 * X(1))
      DSH(1)=Y(1)/DC(1)
 140 CONTINUE
C
      PRINT 1006, TEMS
 1006 FORMAT(/10X, 'FIREBALL SURFACE TEMPERATURE =', F10.3, /10X,
     @30('-')/)
С
С
C
      PRINT 1421
 1421 FORMAT(1X,
               TIME
    @'
       MASS
                       DIAMETER HEIGHT
                                         HUMAN R.
                                                  ANIMAL R. ',
     " BULID.R. PR.EQ.R',
     @/1X,74('-'))
С
C
     DO 145 J=1,M
      PRINT 1400,W(J),TC(J),DC(J),Y(J),
    @RHUM(J),RPIG(J),RB(J),RPE(J)
 1400 FORMAT(8F9.2)
  145
         CONTINUE
С
C
        PRINT 5600
 5600
        FORMAT(10X, ///)
С
C
        RETURN
        END
С
0000
C
    SUBROUTINE FOR CALCULATION OF FIREBALL
C
      С
    DIMENSIONS USING HOT SHELL-COLD CORE MODEL :
C
    C
C
     SUBROUTINE HOTSHE (MM, W, THETA, SECMA)
С
С
С
     DIMENSION Q1(25), Y(25), DC(25), RTAB(25), RTAPE(25), RB(25), RPE(25)
    @,F21(25),DSH(25),RTAHLM(25),TC(25),D0(25),Q2PIG(25),D1(25)
    @,W(25),Q2HLM(25),RHLM(25),RPIG(25),RTAPIG(25),X(25),TAW(25)
С
С
С
```

CC	CALCULATION OF FIREBALL BY USING HOT SHELL-COLD CORE MODEL:
c	
C	
C	QB=12.6
	QPE=37.8
C	
C	READ *, TEMS, WTM, C, RS, ZH
С	
C	
-	READ *, (RTAHLM(J), J=1,MM)
	READ *, (RTAPIG(JJ), JJ=1,MM)
	READ $*$, (RTAB(L), L=1, MM) READ $*$, (RTAPE(J) J=1 MM)
С	
C	
C	F1=((1.0-(29.0/WTM))*(93.05*SECMA*((TEMS/1000 0)**4 0)))/(74
-	@-RS*C*(TEMS-290.0)-93.05*SECMA*((TEMS/1000.0)**4.0)*RS)
c	
С	
~	PRINT *, ' THE VALUE OF F1=', F1
c	
	IF(F1.LT.O.O) THEN
122	PRINT 1222 22 FORMAT(// 5% 'THIS MODEL SHOLD NOT LISED IN TURSE CONDUCTIONS
	@/,5X, 'BECAUSE F1 VALUE IS NEGATIVE & IT MUST BE POSTIVE'. //)
C	
c	
	ELSE
c	
č	
	DO 40 K=1,MM
С	Q1(K)=SECMA*THETA*(TEMS**4.0)
-	D1(K)=(1.567*(1.0+(RS*F1))*W(K))*0.3334
	DO(K)=3.57*((W(K)/WTM)**0.3334)
	$((((273.0^{+}(1.0+1))/(448.0^{+}Q1(K)*(1.0+RS*F1))))$ @+1.0)**0.3334)*D1(K)
С	
~	TC(K) = (DC(K) - DO(K))/1.0
c	
С	
	HUM(K) = DC(K) * RTAHUM(K) $BPIG(K) = DC(K) * RTAPIG(K)$
	RB(K)=DC(K)*RTAB(K)

```
RPE(K) = DC(K) * RTAPE(K)
        Y(K)=10.0*TC(K)
        Q2HUM(K)=28.16/(TC(K)**0.735)
        Q2PIG(K)=50.0/(TC(K)**0.71)
      F21(K) = ((Y(K)/DC(K)+0.5)/(4.0*(((Y(K)/DC(K)+0.5)**2.0))))
     @+(RHLM(K)/DC(K)+0.5))**2.0)**1.5)
      X(K) = DC(K)*(((((Y(K)/DC(K)+0.5)**2.0)+(RHUM(K)/DC(K))**2.0)))
     @**0.5)-0.5)
       TAW(K) = EXP(-0.0007*X(K))
C
             DSH(K)=Y(K)/DC(K)
С
 40
        CONTINUE
С
      PRINT 1007, TEMS
 1007 FORMAT(/10X, 'FIREBALL SURFACE TEMPERATURE =', F10.3, /10X,
     @30('-')/)
C
С
C
С
      PRINT 421
 421 FORMAT(1X,
                  TIME DIAMETER HEIGHT HUMAN R. ANIMAL R.
     @'
          MASS
     @'BULID.R. PR.EQ.R',
     @/1X,74('-'))
C
         DO 45 KI=1,MM
        PRINT 493, W(KI), TC(KI), DC(KI), Y(KI),
     @RHUM(KI), RPIG(KI), RB(KI), RPE(KI)
 493
        FORMAT(8E9.2)
С
  45
        CONTINUE
С
C
С
      IF(DSH(1).GT.2.5) THEN
C
        PRINT 5601, DSH(1)
 5601
           FORMAT(/, 10X, 'THE VALUE OF H/DS =', F10.3, /)
С
С
C
      PRINT 1223
 1223 FORMAT(/, 5X, 'THE VALUE OF F21 BECOMES VERY SMALL AND THAT MEAN',
     @/,5X, 'THE VALUE OF F21 APPROACHES ZERO.',
     @/,5X, 'THEN THIS MODEL MUST NOT USED IN THESE CONDITIONS. ',///)
C
C
С
         ELSE
С
С
          PRINT 29
```

29	FORMAT(3X,76('*')///)
c	ENDIF
сc	RETURN END
0000	
0000	SUBROUTINE PROGRAM FOR CALCULATION OF FIREBALL USING ISO-THERMAL MODEL EQUATIONS:
C C C	SUBROUTINE ISOTHR(LL,W, THETA, SECMA, V, GN, WTM, FC, TEMS)
c	DIMENSION W(25), RTAHUM(25), TC(25), DC(25), DO(25), @RHUM(25), RB(25), RPE(25), Y(25), Q1(25), Q2PIG(25), RTAB(25), @RHUM(25), RTAB(25), 2014(25), Q1(25), Q2PIG(25), RTAB(25),
000	Chin (25), RTAP (G(25), Q2HUM(25), RTAPE(25), F21(25), TAW(25), X(25)
с	READ *, (RTAHUM(J), J=1,LL) READ *, (RTAPIG(KK), KK=1,LL) READ *, (RTAB(K), K=1,LL) READ *, (RTAPE(N), N=1,LL)
0000	CALCULATION OF FIREBALL USING ISOTHERMAL MODEL:
c	QB=12.6 QPE=37.8
000	
~	DO(L)=0.539*((290.0*W(L)/WTM)**0.3333) DC(L)=0.539*((V+(GN+1.0)*FC)**0.3333)*((TEMS*W(L)/WTM)**0.3333) TC(L)=0.45*(W(L)**0.3334)
C	RHUM(L) = DC(L) * RTAHUM(L) $RP IG(L) = DC(L) * RTAP IG(L)$ $RB(L) = RTAB(L) * DC(L)$ $RPE(L) = RTAPE(L) * DC(L)$ $Y(L) = 10.0 * TC(L)$

```
Q1(L)=SECMA*THETA*(TEMS**4.0)
        Q2HUM(L)=28.16/(TC(L)**0.735)
          Q2PIG(L)=50.0/(TC(L)**0.71)
      F21(L)=((Y(L)/DC(L)+0.5)/(4.0*(((Y(L)/DC(L)+0.5)**2.0))))
     @+(RHUM(L)/DC(L)+0.5))**2.00)**1.5)
      X(L)=DC(L)*(((((Y(L)/DC(L)+0.5)**2.0)+(RHUM(L)/DC(L))**2.0)))
     @**0.5)-0.5)
      TAW(L) = EXP(-0.0007 * X(L))
С
С
 50
          CONTINUE
C
      PRINT 1008, TEMS
  1008 FORMAT(/10X, 'FIREBALL SURFACE TEMPERATURE =', F10.3, /10X,
     @30('-')/)
С
С
C
      PRINT 752
 752 FORMAT(1X,
     @'
          MASS
                  TIME
                          DIAMETER HEIGHT HUMAN R. ANIMAL R.',
     @' BULID.R PR.EQ.R',
     @/1X,74('-'))
С
С
C
      DO 55 N=1,LL
      PRINT 500,W(N),TC(N),DC(N),Y(N),RHUM(N),RPIG(N),RB(N),RPE(N)
 500
        FORMAT(8F9.2)
 55
        CONTINUE
С
       PRINT 43
 43
       FORMAT(2X,65('*')////)
C
                RETURN
                 END
С
c
С
c
С
   SUBROUTINE TO CALCULATE B.L.E.V.E. FIREBALL
С
      С
   DIMENSIONS AND PROPERTIES:
C
       С
C
C
C
      SUBROUTINE BLEVE (M, W, THETA, SECMA, TEMS, WTM)
С
С
C
      DIMENSION W(25), RTAHUM(25), TC(25), DC(25), RB(25), RPE(25),
```

```
@Y(25),Q1BL(25),RTAB(25),RTAPE(25),F21BL(25),TAW(25),X(25)
     @, RHUM(25), RPIG(25), RTAPIG(25), Q2PIG(25), Q2HUM(25)
C
С
С
       QB=12.6
       QPE=37.8
C
C
С
       READ *, (RTAHLM(1), I=1, M)
       READ *, (RTAPIG(J), J=1,M)
      READ *, (RTAB(K), K=1,M)
      READ *, (RTAPE(J), J=1,M)
C
C
C
      DO 83 N=1,M
      TC(N)=0.31*((44.8*W(N)/WTM)**0.3333)
      DC(N)=3.44*((44.8*W(N)/WTM)**0.3333)
С
      Y(N)=0.5*DC(N)
      Q1BL(N)=SECMA*THETA*(TEMS**4.0)
        Q2HUM(N)=28.16/(TC(N)**0.735)
        Q2PIG(N)=50.0/(TC(N)**0.71)
         RHUM(N) = RTAHUM(N) * DC(N)
         RPIG(N) = RTAPIG(N) * DC(N)
         RB(N) = RTAB(N) * DC(N)
         RPE(N) = RTAPE(N) * DC(N)
      F21BL(N) = ((Y(N)/DC(N)+0.5)/(4.0*(((Y(N)/DC(N)+0.5)**2.0))))
     @+(RHLM(N)/DC(N)+0.5))**2.0)**1.5)
      X(N) = DC(N)*(((((Y(N)/DC(N)+0.5)**2.0)+(RHUM(N)/DC(N))**2.0)))
     @**0.5)-0.5)
      TAW(N) = EXP(-0.0007*X(N))
C
  83 CONTINUE
С
      PRINT 1009, TEMS
 1009 FORMAT(/10X, 'FIREBALL SURFACE TEMPERATURE =', F10.3, /10X,
     @30('-')/)
С
С
C
      PRINT 703
 703 FORMAT(1X,
     @'
          MASS
                   TIME
                           DIAMETER HEIGHT
                                              HUMAN R. ANIMAL R.',
         BULID.R. PR.EQ.R',
     @'
     @/1X,74('-'))
С
C
      DO 87 K=1,M
      PRINT 808, W(K), TC(K), DC(K), Y(K), RHUM(K), RPIG(K), RB(K), RPE(K)
 808
        FORMAT(8F9.2)
C
```

87	CONTINUE
С	
С	
	PRINT 819
819	FORMAT(2X,//)
С	
	RETURN
	END
С	
С	
С	

FIREBALL DIMENSIONS AND HAZARD RANGES RESULSTS:

USING GAYLE PARAMETERS (A, AS, B & BS)

METHANE FIREBALL RESULTS:

AT FIREBALL SURFACE TEMPERATUE = 2138.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN R	. ANIMAL	R.BULID.R.	PR.EQ.R.
1000.00	2.87	34.98	28.65	76.96	78.71	89.20	59 47
5000.00	5.08	59.12	50.82	156.66	153.70	150.75	100.50
10000.00	6.50	74.11	65.04	211.20	200.08	192.67	129.68
25000.00	9.01	99.90	90.12	309.70	294.71	259.75	174 83
50000.00	11.53	125.23	115.35	407.00	394.48	331.86	225 42
75000.00	13.33	142.93	133.26	478.81	464.52	378.76	257 27
100000.00	14.76	156.98	147.63	541.59	533.74	423.85	290 42
500000.00	26.18	265.28	261.82	941.76	955.02	716.27	490.78

ETHANE FIREBALL RESULTS:

FIREBALL SURFACE TEMPERATUER = 2182.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID	.R. PR.EQ.R
1000.00	2.87	34.98	28.65	94.45	78.71	94,45	62.97
5000.00	5.08	59.12	50.82	168.48	156.66	159.62	106.41
10000.00	6.50	74.11	65.04	218.61	211.20	203.79	137.09
25000.00	9.01	99.90	90.12	314.69	313.69	274.73	184.82

50000.00	11.53	125.23	115.35	413.26	419.52	350.65	237.94
75000.00	13.33	142.93	133.26	493.10	500.25	400.20	271.56
100000.00	14.76	156.98	147.63	557.28	572.98	447.40	306.11
500000.00	26.18	265.28	261.82	981.55	1008.08	756.06	517.30

PROPANE FIREBALL RESULTS:

FIREBALL SURFACE TEMPERATURE = 2188.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID.F	R. PR.EQ.R
1000.00	2.87	34.98	28.65	89.20	80.46	94.45	61 22
5000.00	5.08	59.12	50.82	162.57	156.66	159.62	103 45
10000.00	6.50	74.11	65.04	218.61	214.90	203.79	133 39
25000.00	9.01	99.90	90.12	314.69	309.70	274.73	179 82
50000.00	11.53	125.23	115.35	413.26	407.00	350.65	231 68
75000.00	13.33	142.93	133.26	493.10	485.96	400.20	264 42
100000.00	14.76	156.98	147.63	572.98	549.43	447.40	298 26
500000.00	26.18	265.28	261.82	994.82	955.02	756.06	504.04

BUTANE FIREBALL RESULTS:

FIREBALL SURFACE TEMPERATURE = 2188.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID	.R. PR.EQ.R
1000.00	2.87	34.98	28.65	85.71	82.21	96.20	61 22
5000.00	5.08	59.12	50.82	162.57	156.66	162 57	103 45
10000.00	6.50	74.11	65.04	218.61	214.90	207 49	133 30
25000.00	9.01	99.90	90.12	324.68	309.70	279 73	179 82
50000.00	11.53	125.23	115.35	425.79	407.00	356 91	231 69
75000.00	13.33	142.93	133.26	507.39	485.96	407 35	264 42
100000.00	14.76	156.98	147.63	572.98	549 43	455 25	209.92
500000.00	26.18	265.28	261.82	1021.35	955.02	769.33	504.04

FIREBALL SURFACE TEMPERATURE = 2190.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN	R. ANIMAL	R. BULI	D.R. PR.EQ.R
1000.00	2.87	34.98	28.65	87.46	80.46	96.20	61 22
5000.00	5.08	59.12	50.82	162.57	156.66	162.57	103 45
10000.00	6.50	74.11	65.04	218.61	214.90	207.49	133 39
25000.00	9.01	99.90	90.12	324.68	309.70	279.73	179 82
50000.00	11.53	125.23	115.35	432.05	407.00	356.91	231 68
75000.00	13.33	142.93	133.26	514.54	485.96	407.35	264 42
100000.00	14.76	156.98	147.63	596.53	549.43	455.25	298 26
500000.00	26.18	265.28	261.82	1034.61	955.02	769.33	504.04

USING BRASIE PARAMETERS (A, AS, B & BS)

AT FIREBALL SURFACE TEMPERATUE = 2086.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN	R. ANIMAL	R.BULID.R.	PR.EQ.R.
1000.00	2.99	37.91	29.93	104.26	83.41	104.26	60 66
5000.00	5.12	64.79	51.15	184.67	155.51	178,19	103 67
10000.00	6.44	81.62	64.43	248.93	224.45	228 53	130 59
25000.00	8.74	110.74	87.42	359.90	332.21	310 07	177 19
50000.00	11.01	139.49	110.12	481.24	439.39	397 54	230 16
75000.00	12.60	159.65	126.04	582.74	526.86	455 01	262 42
100000.00	13.87	175.70	138.71	676.46	597.40	509 54	203.43
500000.00	23.71	300.29	237.07	1186.14	1051.01	870.84	495.48

FIREBALL SURFACE TEMPERATUER = 2126.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID	R. PR.EQ.R
1000.00	2.99	37.91	29.93	104.26	83.41	104 26	64 AE
5000.00	5.12	64.79	51.15	191.14	158.75	178 19	110 15
10000.00	6.44	81.62	64.43	257.10	228.53	228 53	138 75
25000.00	8.74	110.74	87.42	370.97	332.21	310.07	188 26
50000.00	11.01	139.49	110.12	495.19	446.37	397.54	237.13

75000.00	12.60	159.65	126.04	598.70	534.84	455.01	271 41
100000.00	13.87	175.70	138.71	676.46	606.18	509.54	298 70
500000.00	23.71	300.29	237.07	1216.17	1066.03	870.84	510.49

FIREBALL SURFACE TEMPERATURE = 2132.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID.R	. PR.EQ.R
1000.00	2.99	37.91	29.93	106.16	85.30	102 36	64 45
5000.00	5.12	64.79	51.15	194.38	161.99	174.95	110 15
10000.00	6.44	81.62	64.43	261.18	228.53	220.37	138 75
25000.00	8.74	110.74	87.42	376.51	337.75	298.99	188 26
50000.00	11.01	139.49	110.12	502.16	453.34	376.62	237 13
75000.00	12.60	159.65	126.04	606.68	542.82	431.07	271 41
100000.00	13.87	175.70	138.71	685.25	614.97	474.40	298 70
500000.00	23.71	300.29	237.07	1231.19	1081.04	810.78	510.49

FIREBALL SURFACE TEMPERATURE = 2132.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL F	R. BULID	.R. PR.EQ.R
1000.00	2.99	37.91	29.93	106.16	85.30	102 36	64 AE
5000.00	5.12	64.79	51.15	194.38	161.99	174 95	110 15
10000.00	6.44	81.62	64.43	265.26	228.53	220 37	129 75
25000.00	8.74	110.74	87.42	382.05	337.75	298 99	199 26
50000.00	11.01	139.49	110.12	509.14	453.34	376 62	227 12
75000.00	12.60	159.65	126.04	614.67	542.82	431 07	237.13
100000.00	13.87	175.70	138.71	694.03	614.97	474 40	208 70
500000.00	23.71	300.29	237.07	1246.20	1081.04	810.78	510.49

FIREBALL SURFACE TEMPERATURE = 2134.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN	R. ANIMAL	R. BULI	D.R. PR.EQ.R
1000.00	2.99	37.91	29.93	106.16	85.30	102.36	64 45
5000.00	5.12	64.79	51.15	194.38	161.99	174.95	110 15
10000.00	6.44	81.62	64.43	269.34	228.53	220.37	138 75
25000.00	8.74	110.74	87.42	387.58	337.75	298.99	188 26
50000.00	11.01	139.49	110.12	516.11	453.34	376.62	237 13
75000.00	12.60	159.65	126.04	622.65	542.82	431.07	271 41
100000.00	13.87	175.70	138.71	702.82	614.97	474 40	298 70
500000.00	23.71	300.29	237.07	1261.22	1081.04	810.78	510.49

USING MARSHALL PARAMETERS (A, AS, B & BS):

METHANE FIREBALL RESULTS:

AT FIREBALL SURFACE TEMPERATUE = 2024.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN	R. ANIMAL	R.BULID.R.	PR.EQ.R.
1000.00	3.79	54.87	37.91	150.90	117.98	137.18	82 31
5000.00	6.48	93.78	64.79	276.66	239.14	234 46	140 67
10000.00	8.16	118.13	81.62	360.30	324.86	295.33	177 20
25000.00	11.07	160.28	110.74	512.89	472.82	400.70	240 42
50000.00	13.95	201.89	139.49	686.43	615.77	504 73	302 84
75000.00	15.97	231.08	159.65	843.43	762.56	577 69	346 62
100000.00	17.57	254.31	175.70	953.66	877.37	635 77	381 46
500000.00	30.03	434.63	300.29	1673.33	1564.67	1086.58	651.95

FIREBALL SURFACE TEMPERATUER = 2060.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID	.R. PR.EQ.F	2
1000.00	3.79	54.87	37.91	150.90	120.72	142 67	87 80	-
5000.00	6.48	93.78	64.79	276.66	243.83	243.83	150.05	
10000.00	8.16	118.13	81.62	366.21	330.77	307.14	189 01	
25000.00	11.07	160.28	110.74	520.91	480.84	416.73	256 45	
50000.00	13.95	201.89	139.49	696.53	625.87	524.92	323.03	

75000.00	15.97	231.08	159.65	854.99	762.56	600.80	369.72
100000.00	17.57	254.31	175.70	966.38	890.08	661.20	406.90
500000.00	30.03	434.63	300.29	1695.06	1586.40	1130.04	695.41

PROPANE FIREBALL RESULTS:

FIREBALL SURFACE TEMPERATURE = 2064.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID.F	R. PR.EQ.R
1000.00	3.79	54.87	37.91	153.65	126.21	142.67	87.80
5000.00	6.48	93.78	64.79	276.66	243.83	243.83	150.05
10000.00	8.16	118.13	81.62	372.11	330.77	307.14	189.01
25000.00	11.07	160.28	110.74	520.91	480.84	416.73	256.45
50000.00	13.95	201.89	139.49	706.62	635.96	524.92	323 03
75000.00	15.97	231.08	159.65	866.54	762.56	600.80	369 72
100000.00	17.57	254.31	175.70	979.09	877.37	661.20	406 90
500000.00	30.03	434.63	300.29	1716.79	1586.40	1130.04	695.41

FIREBALL SURFACE TEMPERATURE = 2064.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID	.R. PR.EQ.R
1000.00	3.79	54.87	37.91	156.39	126.21	142.67	87 80
5000.00	6.48	93.78	64.79	276.66	243.83	243.83	150.05
10000.00	8.16	118.13	81.62	378.02	330.77	307.14	189 01
25000.00	11.07	160.28	110.74	528.92	480.84	416.73	256 45
50000.00	13.95	201.89	139.49	726.81	635.96	524.92	323 03
75000.00	15.97	231.08	159.65	889.65	762.56	600.80	369 72
100000.00	17.57	254.31	175.70	1004.52	877.37	661.20	406 90
500000.00	30.03	434.63	300.29	1781.98	1586.40	1130.04	695.41

PENTANE FIREBALL RESULTS:

FIREBALL SURFACE TEMPERATURE = 2066.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN	R. ANIMAL	R. BULI	D.R. PR.EQ.R
1000.00	2.99	37.91	29.93	108.05	87.20	98.57	60,66
5000.00	5.12	64.79	51.15	194.38	168.47	168.47	103.67
10000.00	6.44	81.62	64.43	261.18	228.53	212.21	130.59
25000.00	8.74	110.74	87.42	370.97	332.21	287.92	177 18
50000.00	11.01	139.49	110.12	502.16	439.39	362.67	223 18
75000.00	12.60	159.65	126.04	614.67	526.86	415.10	255 45
100000.00	13.87	175.70	138.71	694.03	606.18	456.83	281 13
500000.00	23.71	300.29	237.07	1246.20	1096.06	780.75	480.46

USING ROBERTS PARAMETERS (A, AS, B & BS):

METHANE FIREBALL RESULTS:

AT FIREBALL SURFACE TEMPERATUE = 1980.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN	R. ANIMAL	R.BULID.R.	PR.EQ.R.
1000.00	4.49	57.87	44.90	150.45	144.67	138.88	81 01
5000.00	7.67	98.90	76.73	291.75	271.97	237.35	138 46
10000.00	9.67	124.57	96.65	386.18	367.49	298.98	174 40
25000.00	13.11	169.02	131.14	549.32	523.97	405.65	236 63
50000.00	16.52	212.90	165.18	723.88	681.29	510.97	298.07
75000.00	18.91	243.68	189.06	865.07	804.15	584.84	341 15
100000.00	20.81	268.18	208.07	1019.09	911.82	643.63	375 45
500000.00	35.56	458.34	355.61	1810.43	1627.10	1100.01	641.67

FIREBALL SURFACE TEMPERATUER = 2012.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID	.R. PR.EQ.F	2
1000.00	4.49	57.87	44.90	147.56	130.20	144.67	86 80	-
5000.00	7.67	98.90	76.73	291.75	247.24	247.24	148 35	
10000.00	9.67	124.57	96.65	392.41	348.81	311.44	186.86	
25000.00	13.11	169.02	131.14	549.32	490.16	422.55	253 53	
50000.00	16.52	212.90	165.18	734.52	660.00	532.26	319.36	

75000.00	18.91	243.68	189.06	877.26	791.97	609.20	365.52
100000.00	20.81	268.18	208.07	1019.09	911.82	670.45	402.27
500000.00	35.56	458.34	355.61	1833.35	1650.01	1145.84	687.51

PROPANE FIREBALL RESULTS:

FIREBALL SURFACE TEMPERATURE = 2016.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID.R	. PR.EQ.R
1000.00	4.49	57.87	44.90	150.45	133.09	150.45	89.69
5000.00	7.67	98.90	76.73	286.80	247.24	257.13	153.29
10000.00	9.67	124.57	96.65	398.64	348.81	323.89	193.09
25000.00	13.11	169.02	131.14	557.77	498.61	439.46	261 98
50000.00	16.52	212.90	165.18	745.17	670.65	553.55	330.00
75000.00	18.91	243.68	189.06	889.44	816.33	633.57	377 71
100000.00	20.81	268.18	208.07	1019.09	925.22	697.27	415 68
500000.00	35.56	458.34	355.61	1856.27	1672.93	1191.68	710.42

FIREBALL SURFACE TEMPERATURE = 2016.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID	.R. PR.EQ.R
1000.00	4.49	57.87	44.90	153.35	144.67	150.45	P3 P8
5000.00	7.67	98.90	76.73	291.75	276.91	257.13	153 29
10000.00	9.67	124.57	96.65	404.87	367.49	323.89	193 09
25000.00	13.11	169.02	131.14	574.67	532.42	439.46	261 98
50000.00	16.52	212.90	165.18	755.81	702.59	553.55	330.00
75000.00	18.91	243.68	189.06	913.81	840.70	633.57	377 71
100000.00	20.81	268.18	208.07	1032.50	952.04	697.27	415 68
500000.00	35.56	458.34	355.61	1879.18	1695.85	1191.68	710.42

FIREBALL SURFACE TEMPERATURE = 2016.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN	R. ANIMAL	R. BULI	D.R. PR.EQ.R
1000.00	4.49	57.87	44.90	156.24	150.45	150.45	86.80
5000.00	7.67	98.90	76.73	301.64	281.86	257.13	148.35
10000.00	9.67	124.57	96.65	411.09	373.72	323.89	186.86
25000.00	13.11	169.02	131.14	591.58	540.87	439.46	253 53
50000.00	16.52	212.90	165.18	787.75	713.23	553.55	319 36
75000.00	18.91	243.68	189.06	938.18	852.89	633.57	365 52
100000.00	20.81	268.18	208.07	1086.13	978.86	697.27	402 27
500000.00	35.56	458.34	355.61	1925.02	1764.60	1191.68	687.51

USING HOT SHELL-COLD CORE MODEL:

METHANE RESULTS OF HOT SHELL-COLD CORE MODEL:

FIREBALL SURFACE TEMPERATURE = 1500.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL R.	BULID.R.	PR.EQ.R	
1000.00	55.59	69.80	55.59	157.04	153.55	202.41	125.63	
5000.00	95.07	119.36	95.07	310.34	298.40	346.15	214.85	
10000.00	119.78	150.39	119.78	421.10	413.58	436.14	270.71	
25000.00	162.58	204.13	162.58	632.79	612.38	591.97	367.43	
50000.00	204.84	257.19	204.84	835.88	810.16	745.87	462.95	
75000.00	234.49	294.42	234.49	1001.04	956.87	853.83	529.96	
100000.00	258.10	324.06	258.10	1150.41	1085.60	939.77	583 31	
500000.00	441.39	554.19	441.39	2050.52	1939.68	1607.16	997 55	

BUTANE RESULTS OF HOT SHELL-COLD CORE MODEL:

THE VALUE OF F1= 7.14186E-03

FIREBALL SURFACE TEMPERATURE = 1300.000

MASS TIME DIAMETER HEIGHT HUMAN R. ANIMAL R. BULID.R. PR.EQ.R

0.10E+04 0.31E+04 0.31E+04 0.31E+05 0.00E+01 0.00E+01 0.00E+01 0.00E+01 0.50E+04 0.16E+05 0.16E+05 0.16E+06 0.00E+01 0.00E+01 0.00E+01 0.00E+01 0.10E+05 0.31E+05 0.31E+05 0.31E+06 0.00E+01 0.00E+01 0.00E+01 0.00E+01 0.25E+05 0.78E+05 0.78E+05 0.78E+06 0.00E+01 0.00E+01 0.00E+01 0.00E+01
0.50E+05 0.16E+06 0.16E+06 0.16E+07 0.00E+01 0.00E+01 0.00E+01 0.00E+01 0.00E+01 0.00E+01 0.00E+01 0.75E+05 0.23E+06 0.23E+06 0.23E+07 0.00E+01 0.0

THE VALUE OF H/DS = 9.971

THE VALUE OF F21 BECOMES VERY SMALL AND THAT MEAN THE VALUE OF F21 APPROACHES ZERO. THEN THIS MODEL MUST NOT USED IN THESE CONDITIONS.

USING ISO-THERMAL MODEL:

METANE RESULTS OF ISO-THERMAL MODEL:

FIREBALL SURFACE TEMPERATURE = 1980.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN	R. ANIMAL	R. BULID	.R PR.EQ.R	
1000.00	4.50	58.65	45.02	155.42	149.56	140.76	82 11	-
5000.00	7.70	100.29	76.99	280.80	270.77	240.69	140 40	
10000.00	9.70	126.35	97.01	379.05	379.05	303.24	176 89	
25000.00	13.17	171.48	131.67	548.73	548.73	411.55	240 07	
50000.00	16.59	216.04	165.90	723.74	723.74	518.50	302 46	
75000.00	18.99	247.30	189.91	877.93	853.20	593.53	346.23	
100000.00	20.90	272.19	209.03	1020.71	966.28	653.26	381 07	
500000.00	35.75	465.41	357.48	1791.84	1722.03	1116.99	651 58	
******	******	*****	******	*****	******	*******	***	

ETHANE RESULTS OF ISO-THERMAL MODEL:

FIREBALL SURFACE TEMPERATURE = 2012.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN F	R. ANIMAL	R. BULID	R PR.EQ.F	2
1000.00	4.50	57.44	45.02	149.34	134.98	143.60	86 16	
5000.00	7.70	98.21	76.99	270.09	270.09	245.54	147 32	
10000.00	9.70	123.74	97.01	365.03	377.40	309.35	185 61	
25000.00	13.17	167.93	131.67	529.00	537.39	419.84	251 90	
50000.00	16.59	211.58	165.90	687.63	708.79	528.95	317.37	

75000.00	18.99	242.20	189.91	835.57	847.68	605.49	363.29
100000.00	20.90	266.57	209.03	986.30	972.97	666.42	399.85
500000.00	35.75	455.80	357.48	1777.62	1709.25	1139.50	683.70
*******	******	******	*****	****	*****	*******	***

PROPANE RESULTS OF ISO-THERMAL MODEL:

FIREBALL SURFACE TEMPERATURE = 2016.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN	R. ANIMAL	R. BULID	R PR.EQ.F	R
1000.00	4.50	56.89	45.02	139.38	136.53	142.22	85.33	
5000.00	7.70	97.27	76.99	262.64	252.91	243.19	145.91	
10000.00	9.70	122.56	97.01	355.41	337.03	306.39	183.83	
25000.00	13.17	166.33	131.67	515.62	498.98	415.82	249 49	
50000.00	16.59	209.56	165.90	691.53	670.58	523.89	314 33	
75000.00	18.99	239.88	189.91	839.57	803.59	599.69	359 82	
100000.00	20.90	264.02	209.03	976.86	924.06	660.04	396 03	
500000.00	35.75	451.44	357.48	1738.04	1647.75	1128 60	677 16	
******	*****	*****	****	******	******	*******	k**	

BUTANE RESULTS OF ISO-THERMAL MODEL:

FIREBALL SURFACE TEMPERATURE = 2016.000

	 	 	LIMIO	1 1km -	- 20	10.0
14	 					

MASS	TIME	DIAMETER	HEIGHT	HUMAN	R. ANIMAL	R. BULID	.R PR.EQ.R
1000.00	4.50	56.58	45.02	144.29	135.80	141.46	84 87
5000.00	7.70	96.75	76.99	261.23	251.55	241.88	145.13
10000.00	9.70	121.90	97.01	347.40	341.31	304.74	182.84
25000.00	13.17	165.43	131.67	512.84	512.84	413.58	248.15
50000.00	16.59	208.43	165.90	687.81	687.81	521.07	312.64
75000.00	18.99	238.59	189.91	835.05	823.12	596.46	357.88
100000.00	20.90	262.59	209.03	958.47	945.34	656.49	393.89
500000.00	35.75	449.01	357.48	1706.22	1683.77	1122.52	673.51
*******	******	******	******	*****	*******	******	

PENTANE RESULTS OF ISO-THERMAL MODDEL:

FI	REBALL	SURFACE	TEMPERATURE =	= 2016.000
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MASS	TIME	DIAMETER	HEIGHT	HLMAN	R. ANIMAL	R. BULID	R PR.EQ.F	R
1000.00	4.50	56.39	45.02	140.98	132.52	140,98	84.59	
5000.00	7.70	96.42	76.99	260.34	250.70	241.05	144.63	
10000.00	9.70	121.48	97.01	346.22	334.07	303.70	182.22	
25000.00	13.17	164.87	131.67	494.61	511.10	412.18	247.31	
50000.00	16.59	207.72	165.90	664.70	675.09	519.30	311.58	
75000.00	18.99	237.78	189.91	808.44	808.44	594.44	356 66	
100000.00	20.90	261.70	209.03	929.05	915.96	654.26	392 55	
500000.00	35.75	447.48	357.48	1678.06	1633.31	1118.70	671.22	
******	******	*****	******	*****	*****	*******	***	

B.L.E.V.E FIREBALL RESULTS:

METHANE BLEVE FIREBALL RESULTS:

FIREBALL SURFACE TEMPERATURE = 1482.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID	R. PR.EQ.R
1000.00	4.37	48.47	24.24	60.59	60.59	67.86	33 93
5000.00	7.47	82.88	41.44	124.32	116.04	116.04	58 02
10000.00	9.41	104.42	52.21	177.52	167.08	146.19	73 10
25000.00	12.77	141.72	70.86	269.27	255.10	198.41	99.20
50000.00	16.09	178.55	89.28	366.03	357.10	249.97	124 99
75000.00	18.42	204.39	102.19	439.43	429.21	286.14	143 07
100000.00	20.27	224.96	112.48	562.39	506.15	314.94	157 47
500000.00	34.66	384.65	192.32	1019.32	961.62	538.51	269.25

ETHANE BLEVE FIREBALL RESULTS:

FIREBALL SURFACE TEMPERATURE = 1642.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL F	. BULID.	R. PR.EQ.R
1000.00	3.54	39.31	19.66	55.03	55.03	70.76	35.38
5000.00	6.06	67.22	33.61	107.55	114.27	120.99	60.49
10000.00	7.63	84.68	42.34	148.20	156.67	152.43	76.22

25000.00	10.36	114.93	57.47	229.86	241.36	206.88	103.44
50000.00	13.05	144.80	72.40	318.56	333.04	260.64	130.32
75000.00	14.94	165.75	82.88	397.81	414.38	298.36	149.18
100000.00	16.44	182.43	91.22	465.21	483.45	328.38	164 19
500000.00	28.11	311.94	155.97	873.43	873.43	561.49	280.75

PROPANE BLEVE FIREBALL RESULTS:

FIREBALL SURFACE TEMPERATURE = 1740.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID	.R. PR.EQ.R
1000.00	3.12	34.60	17.30	51.90	51.90	69.20	38.06
5000.00	5.33	59.16	29.58	100.57	103.53	118.32	65.08
10000.00	6.72	74.54	37.27	141.62	149.07	149.07	81 99
25000.00	9.12	101.16	50.58	212.43	222.55	202.32	111 27
50000.00	11.49	127.45	63.72	293.13	305.88	254.90	140 19
75000.00	13.15	145.89	72.94	364.72	372.02	291.78	160.48
100000.00	14.47	160.57	80.29	433.54	425.51	321.14	176 63
500000.00	24.74	274.56	137.28	782.49	782.49	549.12	302.01

BUTANE BLEVE FIREBALL RESULTS:

FIREBALL SURFACE TEMPERATURE = 1818.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID.	R. PR.EQ.R
1000.00	2.84	31.56	15.78	47.33	50,49	69.42	37 87
5000.00	4.86	53.96	26.98	94.42	99.82	118.70	64.75
10000.00	6.13	67.98	33.99	132.56	139.36	149.56	81.58
25000.00	8.31	92.26	46.13	193.75	202.97	202.97	110.71
50000.00	10.47	116.24	58.12	267.35	278.97	255.72	139.48
75000.00	11.99	133.06	66.53	332.64	345.95	292.72	159.67
100000.00	13.20	146.45	73.22	388.08	402.73	322.18	175 74
500000.00	22.57	250.41	125.20	713.66	726.18	550.89	300.49

PENTANE BLEVE FIREBALL RESULTS:

FIREBALL SURFACE TEMPERATURE = 1884.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID.	R. PR.EQ.R
1000.00	2.65	29.36	14.68	46.98	49.91	67.53	38.17
5000.00	4.52	50.21	25.10	90.37	102.92	115.47	65.27
10000.00	5.70	63.25	31.63	129.67	142.32	145.48	82.23
25000.00	7.74	85.84	42.92	188.86	210.32	197.44	111.60
50000.00	9.75	108.16	54.08	259.57	286.61	248.76	140.60
75000.00	11.16	123.81	61.90	321.89	352.84	284.75	160.95
100000.00	12.28	136.26	68.13	374.73	401.98	313.41	177.14
500000.00	21.00	233.00	116.50	675.69	722.29	535.89	302.89

FIREBALL DIMENSIONS AND HAZARD RANGES RESULSTS:

USING GAYLE PARAMETERS (A, AS, B & BS)

METHANE FIREBALL RESULTS:

AT FIREBALL SURFACE TEMPERATUE = 2014.468

MASS	TIME	DIAMETER	HEIGHT	HUMAN	R. ANIMAL	R.BULID.R.	PR.EQ.R.
1000.00	2.87	34.98	28.65	69.96	76.96	87.46	52 47
5000.00	5.08	59.12	50.82	133.01	144.84	147.79	88 68
10000.00	6.50	74.11	65.04	192.67	192.67	185.26	111 16
25000.00	9.01	99.90	90.12	274.73	289.72	249.76	149 85
50000.00	11.53	125.23	115.35	369.43	394.48	313.08	187 85
75000.00	13.33	142.93	133.26	450.22	471.66	357.32	214 39
100000.00	14.76	156.98	147.63	518.04	541.59	392 45	235 47
500000.00	26.18	265.28	261.82	915.23	941.76	663.21	397.93

FIREBALL SURFACE TEMPERATUER = 2092.953

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID.	R. PR.EQ.R
1000.00	2.87	34.98	28.65	90.95	69.96	94,45	55 97
5000.00	5.08	59.12	50.82	165.53	135.97	159.62	94.59
10000.00	6.50	74.11	65.04	218.61	188.97	200.08	118.57
25000.00	9.01	99.90	90.12	314.69	264.74	269.74	159.84

50000.00	11.53	125.23	115.35	419.52	375.69	338.12	200.37
75000.00	13.33	142.93	133.26	500.25	464.52	385.91	228.69
100000.00	14.76	156.98	147.63	565.13	533.74	423.85	251.17
500000.00	26.18	265.28	261.82	981.55	941.76	716.27	424.46

FIREBALL SURFACE TEMPERATURE = 2113.337

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID.F	R. PR.EQ.R
1000.00	2.87	34.98	28.65	83.96	80.46	94,45	59.47
5000.00	5.08	59.12	50.82	156.66	147.79	159.62	100.50
10000.00	6.50	74.11	65.04	218.61	200.08	200.08	125.98
25000.00	9.01	99.90	90.12	309.70	299.71	269.74	169.83
50000.00	11.53	125.23	115.35	407.00	394.48	338.12	212.89
75000.00	13.33	142.93	133.26	485.96	464.52	385.91	242 98
100000.00	14.76	156.98	147.63	557.28	525.89	423.85	266 87
500000.00	26.18	265.28	261.82	968.29	941.76	716.27	450.98

FIREBALL SURFACE TEMPERATURE = 2120.308

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID	.R. PR.EQ.R
1000.00	2.87	34.98	28.65	82.21	76.96	94.45	59 47
5000.00	5.08	59.12	50.82	156.66	147.79	159.62	100.50
10000.00	6.50	74.11	65.04	211.20	200.08	200.08	125.98
25000.00	9.01	99.90	90.12	314.69	299.71	269.74	169.83
50000.00	11.53	125.23	115.35	419.52	388.22	338.12	212.89
75000.00	13.33	142.93	133.26	493.10	471.66	385.91	242 98
100000.00	14.76	156.98	147.63	557.28	533.74	423.85	266 87
500000.00	26.18	265.28	261.82	968.29	941.76	716.27	450.98

PENTANE FIREBALL RESULTS:

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FIREBALL SURFACE TEMPERATURE = 2127.224

MASS	TIME	DIAMETER	HEIGHT	HUMAN R	R. ANIMAL	R. BULI	D.R. PR.EQ.R
1000.00	2.87	34.98	28.65	89.20	87.46	94.45	59 47
5000.00	5.08	59.12	50.82	162.57	159.62	159.62	100.50
10000.00	6.50	74.11	65.04	222.32	222.32	200.08	125.98
25000.00	9.01	99.90	90.12	314.69	314.69	269.74	169.83
50000.00	11.53	125.23	115.35	407.00	407.00	338.12	212 89
75000.00	13.33	142.93	133.26	500.25	493.10	385 91	242 98
100000.00	14.76	156.98	147.63	588.68	557.28	423.85	266 87
500000.00	26.18	265.28	261.82	1034.61	968.29	716.27	450.98

USING BRASIE PARAMETERS (A, AS, B & BS)

METHANE FIREBALL RESULTS:

AT FIREBALL SURFACE TEMPERATUE = 2014.468

MASS	TIME	DIAMETER	HEIGHT	HUMAN	R. ANIMAL	R.BULID.R.	PR.EQ.R.
1000.00	2.99	37.91	29.93	98.57	90.99	94.78	56 87
5000.00	5.12	64.79	51.15	178.19	165.23	161.99	97 19
10000.00	6.44	81.62	64.43	240.77	224.45	204.04	122 43
25000.00	8.74	110.74	87.42	337.75	326.68	276.85	166 11
50000.00	11.01	139.49	110.12	453.34	432.42	348.72	209 23
75000.00	12.60	159.65	126.04	550.81	510,89	399 13	239 48
100000.00	13.87	175.70	138.71	623.75	579.83	439 26	263 56
500000.00	23.71	300.29	237.07	1096.06	1036.00	750.72	450.43

FIREBALL SURFACE TEMPERATUER = 2092.953

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID	R. PR.EQ.	R
1000.00	2.99	37.91	29.93	98.57	92.89	102 36	64 45	-
5000.00	5.12	64.79	51.15	178.19	168.47	174 95	110 15	
10000.00	6.44	81.62	64.43	240.77	232.61	220.37	138 75	
25000.00	8.74	110.74	87.42	348.83	332.21	298.99	188 26	
50000.00	11.01	139.49	110.12	467.29	439.39	376.62	237.13	

75000.00	12.60	159.65	126.04	566.77	518.87	431.07	271.41
100000.00	13.87	175.70	138.71	641.32	597.40	474.40	298.70
500000.00	23.71	300.29	237.07	1141.10	1066.03	810.78	510.49

FIREBALL SURFACE TEMPERATURE = 2113.337

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID.R	. PR.EQ.R
1000.00	2.99	37.91	29.93	102.36	94.78	106.16	64.45
5000.00	5.12	64.79	51.15	187.91	174.95	181.43	110.15
10000.00	6.44	81.62	64.43	253.01	236.69	228.53	138.75
25000.00	8.74	110.74	87.42	365.44	343.29	310.07	188 26
50000.00	11.01	139.49	110.12	481.24	453.34	390.57	237 13
75000.00	12.60	159.65	126.04	566.77	534.84	447.03	271 41
100000.00	13.87	175.70	138.71	641.32	614.97	491.97	298 70
500000.00	23.71	300.29	237.07	1141.10	1111.07	840.81	510.49

FIREBALL SURFACE TEMPERATURE = 2120.308

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID	.R. PR.EQ.R
1000.00	2.99	37.91	29.93	89.09	94.78	106.16	64 45
5000.00	5.12	64.79	51.15	165.23	174.95	181 43	110 15
10000.00	6.44	81.62	64.43	228.53	236.69	228 53	138 75
25000.00	8.74	110.74	87.42	332.21	343.29	310 07	188 26
50000.00	11.01	139.49	110.12	439.39	453.34	390 57	237 12
75000.00	12.60	159.65	126.04	534.84	542.82	447 03	271 41
100000.00	13.87	175.70	138.71	606.18	623.75	491 97	208 70
500000.00	23.71	300.29	237.07	1096.06	1096.06	840.81	510.49

FIREBALL SURFACE TEMPERATURE = 2127.224

MASS	TIME	DIAMETER	HEIGHT	HUMAN	R. ANIMAL	R. BULI	D.R. PR.EQ.R
1000.00	2.99	37.91	29.93	102.36	94.78	106.16	64.45
5000.00	5.12	64.79	51.15	187.91	174.95	181.43	110, 15
10000.00	6.44	81.62	64.43	253.01	236.69	228.53	138 75
25000.00	8.74	110.74	87.42	365.44	343.29	310.07	188.26
50000.00	11.01	139.49	110.12	488.21	453.34	390.57	237 13
75000.00	12.60	159.65	126.04	590.72	534.84	447.03	271 41
100000.00	13.87	175.70	138.71	676.46	606.18	491.97	298 70
500000.00	23.71	300.29	237.07	1231.19	1081.04	840.81	510.49

USING MARSHALL PARAMETERS (A, AS, B & BS):

METHANE FIREBALL RESULTS:

AT FIREBALL SURFACE TEMPERATUE = 2014.468

MASS	TIME	DIAMETER	HEIGHT	HUMAN	R. ANIMAL	R.BULID.R.	PR.EQ.R.
1000.00	3.79	54.87	37.91	153.65	131.70	137.18	82 31
5000.00	6.48	93.78	64.79	281.35	253.21	234.46	140 67
10000.00	8.16	118.13	81.62	372.11	342.58	295.33	177 20
25000.00	11.07	160.28	110.74	520.91	488.85	400.70	240 42
50000.00	13.95	201.89	139.49	706.62	646.06	504.73	302 84
75000.00	15.97	231.08	159.65	854.99	762.56	577.69	346 62
100000.00	17.57	254.31	175.70	979.09	877.37	635 77	381 46
500000.00	30.03	434.63	300.29	1716.79	1586.40	1086.58	651.95

ETHANE FIREBALL RESULTS:

FIREBALL SURFACE TEMPERATUER = 2092.953

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID	.R. PR.EQ.F	2
1000.00	3.79	54.87	37.91	153.65	142.67	148.16	93 28	-
5000.00	6.48	93.78	64.79	276.66	257.90	253.21	159 43	
10000.00	8.16	118.13	81.62	372.11	348.49	318.95	200 82	
25000.00	11.07	160.28	110.74	528.92	496.87	432.75	272.47	
50000.00	13.95	201.89	139.49	706.62	646.06	545.11	343.22	

75000.00	15.97	231.08	159.65	854.99	774.11	623.91	392.83
100000.00	17.57	254.31	175.70	979.09	877.37	686.64	432.33
500000.00	30.03	434.63	300.29	1738.52	1564.67	1173.50	738.87

FIREBALL SURFACE TEMPERATURE = 2113.337

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID.R.	PR.EQ.R
1000.00	3.79	54.87	37.91	153.65	137.18	148,16	87 80
5000.00	6.48	93.78	64.79	276.66	253.21	253.21	150.05
10000.00	8.16	118.13	81.62	372.11	354.39	318.95	189 01
25000.00	11.07	160.28	110.74	520.91	504.88	432.75	256 45
50000.00	13.95	201.89	139.49	706.62	666.24	545.11	323 03
75000.00	15.97	231.08	159.65	866.54	785.66	623.91	369 72
100000.00	17.57	254.31	175.70	991.81	902.80	686.64	406 90
500000.00	30.03	434.63	300.29	1738.52	1586.40	1173.50	695.41

FIREBALL SURFACE TEMPERATURE = 2120.308

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID	.R. PR.EQ.R
1000.00	3.79	54.87	37.91	156.39	131.70	148,16	90 54
5000.00	6.48	93.78	64.79	281.35	253.21	253.21	154 74
10000.00	8.16	118.13	81.62	378.02	336.67	318.95	194 92
25000.00	11.07	160.28	110.74	528.92	488.85	432.75	264 46
50000.00	13.95	201.89	139.49	726.81	646.06	545.11	333 12
75000.00	15.97	231.08	159.65	889.65	774.11	623 91	381 28
100000.00	17.57	254.31	175.70	1004.52	890.08	686.64	419 61
500000.00	30.03	434.63	300.29	1803.71	1608.13	1173.50	717.14

FIREBALL SURFACE TEMPERATURE = 2127.224

MASS	TIME	DIAMETER	HEIGHT	HUMAN	R. ANIMAL	R. BULI	D.R. PR.EQ.R
1000.00	2.99	37.91	29.93	108.05	92.89	102.36	62.56
5000.00	5.12	64.79	51.15	197.62	178.19	174.95	106.91
10000.00	6.44	81.62	64.43	265.26	236.69	220.37	134 67
25000.00	8.74	110.74	87.42	376.51	343.29	298.99	182 72
50000.00	11.01	139.49	110.12	502.16	460.31	376.62	230 16
75000.00	12.60	159.65	126.04	614.67	558.79	431 07	263 43
100000.00	13.87	175.70	138.71	711.60	641.32	474.40	289 91
500000.00	23.71	300.29	237.07	1261.22	1141.10	810.78	495.48

USING ROBERTS PARAMETERS (A, AS, B & BS):

METHANE FIREBALL RESULTS:

AT FIREBALL SURFACE TEMPERATUE = 2014.468

MASS	TIME	DIAMETER	HEIGHT	HUMAN	R. ANIMAL	R.BULID.R.	PR.EQ.R.
1000.00	4.49	57.87	44.90	153.35	144.67	144.67	86 80
5000.00	7.67	98.90	76.73	291.75	276.91	247.24	148 35
10000.00	9.67	124.57	96.65	392.41	379.95	311.44	186 86
25000.00	13.11	169.02	131.14	549.32	540.87	422.55	253 53
50000.00	16.52	212.90	165.18	723.88	723.88	532.26	319 36
75000.00	18.91	243.68	189.06	877.26	877.26	609 20	365 52
100000.00	20.81	268.18	208.07	1019.09	1005.68	670.45	402 27
500000.00	35.56	458.34	355.61	1833.35	1764.60	1145.84	687.51

FIREBALL SURFACE TEMPERATUER = 2092.953

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID	.R. PR.EQ.R	2
1000.00	4.49	57.87	44.90	147.56	141.77	159,13	95 48	•
5000.00	7.67	98.90	76.73	291.75	262.08	271.97	163 18	
10000.00	9.67	124.57	96.65	392.41	367.49	342.58	205 55	
25000.00	13.11	169.02	131.14	557.77	532.42	464.81	278 89	
50000.00	16.52	212.90	165.18	745.17	702.59	585.49	351.29	

75000.00	18.91	243.68	189.06	889.44	840.70	670.13	402.08
100000.00	20.81	268.18	208.07	1019.09	965.45	737.50	442.50
500000.00	35.56	458.34	355.61	1856.27	1718.76	1260.43	756.26

PROPANE FIREBALL RESULTS:

FIREBALL SURFACE TEMPERATURE = 2113.337

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID.R.	PR.EQ.R
1000.00	4.49	57.87	44.90	153.35	144.67	162.03	98 37
5000.00	7.67	98.90	76.73	291.75	267.02	276.91	168 13
10000.00	9.67	124.57	96.65	398.64	373.72	348.81	211 78
25000.00	13.11	169.02	131.14	574.67	532.42	473.26	287 34
50000.00	16.52	212.90	165.18	766.46	702.59	596.13	361 94
75000.00	18.91	243.68	189.06	901.62	840.70	682.31	414 26
100000.00	20.81	268.18	208.07	1032.50	965.45	750.91	455 91
500000.00	35.56	458.34	355.61	1879.18	1695.85	1283.34	779.17

BUTANE FIREBALL RESULTS:

FIREBALL SURFACE TEMPERATURE = 2120.308

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID	.R. PR.EQ.R
1000.00	4.49	57.87	44.90	156.24	147.56	162.03	98 37
5000.00	7.67	98.90	76.73	291.75	276.91	276.91	168 13
10000.00	9.67	124.57	96.65	392.41	373.72	348 81	211 78
25000.00	13.11	169.02	131.14	566.22	540.87	473.26	287 34
50000.00	16.52	212.90	165.18	755.81	723.88	596 13	361 04
75000.00	18.91	243.68	189.06	913.81	852.89	682 31	414 26
100000.00	20.81	268.18	208.07	1045.91	992.27	750 91	414.20
500000.00	35.56	458.34	355.61	1879.18	1764.60	1283.34	779.17

FIREBALL SURFACE TEMPERATURE = 2127.224

MASS	TIME	DIAMETER	HEIGHT	HUMAN I	R. ANIMAL	R. BULIC	R. PR.EQ.R
1000.00	4.49	57.87	44.90	162.03	150.45	159,13	98.37
5000.00	7.67	98.90	76.73	296.69	271.97	271.97	168.13
10000.00	9.67	124.57	96.65	404.87	373.72	342.58	211.78
25000.00	13.11	169.02	131.14	591.58	540.87	464.81	287.34
50000.00	16.52	212.90	165.18	798.39	723.88	585.49	361.94
75000.00	18.91	243.68	189.06	950.36	852.89	670.13	414 26
100000.00	20.81	268.18	208.07	1099.54	965.45	737.50	455 91
500000.00	35.56	458.34	355.61	1947.93	1718.76	1260.43	779.17

USING HOT SHELL-COLD CORE MODEL:

METHANE RESULTS OF HOT SHELL-COLD CORE MODEL:

FIREBALL SURFACE TEMPERATURE = 1500.000

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL R.	BULID.R.	PR.EQ.R
1000.00	55.59	69.80	55.59	153.55	150.06	209.39	136,10
5000.00	95.07	119.36	95.07	298.40	298.40	358.08	232.75
10000.00	119.78	150.39	119.78	406.06	406.06	451.18	293.27
25000.00	162.58	3 204.13	162.58	591.97	591.97	612.38	398.05
50000.00	204.84	257.19	204.84	797.30	784.44	771.58	501.53
75000.00	234.49	294.42	234.49	956.87	927.43	883.27	574.12
100000.00	258.10	324.06	258.10	1085.60	1036.99	972.18	631.92
500000.00	441.39	554.19	441.39	1911.97	1856.55	1662.58	1080.68

BUTANE RESULTS OF HOT SHELL-COLD CORE MODEL:

THE VALUE OF F1= 7.14186E-03

FIREBALL SURFACE TEMPERATURE = 1300.000

MASS TIME DIAMETER HEIGHT HUMAN R. ANIMAL R. BULID.R. PR.EQ.R

0.10E+04 0.31E+04 0.31E+04 0.31E+05 0.00E+01 0.00E+01 0.00E+01 0.00E+01 0.00E+01 0.00E+01 0.00E+01 0.50E+04 0.16E+05 0.16E+05 0.16E+06 0.00E+01 0.0

0.50E+05 0.16E+06 0.16E+06 0.16E+07 0.00E+01 0.00E+01 0.00E+01 0.00E+01 0.00E+01 0.00E+01 0.00E+01 0.75E+05 0.23E+06 0.23E+06 0.23E+07 0.00E+01 0.0

THE VALUE OF H/DS = 9.971

THE VALUE OF F21 BECOMES VERY SMALL AND THAT MEAN THE VALUE OF F21 APPROACHES ZERO. THEN THIS MODEL MUST NOT USED IN THESE CONDITIONS.

USING ISO-THERMAL MODEL:

METANE RESULTS OF ISO-THERMAL MODEL:

FIREBALL SURFACE TEMPERATURE = 2014.468

MASS	TIME	DIAMETER	HEIGHT	HUMAN	R. ANIMAL	R. BULID	R PR.EQ.R	
1000.00	4.50	58.99	45.02	159.27	150.42	147.47	97 33	-
5000.00	7.70	100.86	76.99	287.46	287.46	252.16	166.43	
10000.00	9.70	127.08	97.01	393.94	387.59	317.70	209 68	
25000.00	13.17	172.47	131.67	560.52	543.27	431.17	284.57	
50000.00	16.59	217.29	165.90	749.65	717.05	543.22	358.53	
75000.00	18.99	248.73	189.91	895.43	858.12	621.83	410.41	
100000.00	20.90	273.76	209.03	1040.29	999.23	684.40	451.70	
500000.00	35.75	468.10	357.48	1848.99	1778.78	1170.25	772.36	
*******	*******	******	******	*****	*******	****		

ETHANE RESULTS OF ISO-THERMAL MODEL:

FIREBALL SURFACE TEMPERATURE = 2092.953

MASS	TIME	DIAMETER	HEIGHT	HUMAN F	R. ANIMAL	R. BULID	R PR.EQ.R	2
1000.00	4.50	58.20	45.02	154.23	148.41	157.14	98 94	
5000.00	7.70	99.51	76.99	278.64	283.61	268.69	169.17	
10000.00	9.70	125.38	97.01	376.13	382.40	338.52	213.14	
25000.00	13.17	170.16	131.67	544.50	553.01	459.43	289.27	
50000.00	16.59	214.38	165.90	718.17	718.17	578.83	364.45	

75000.00	18.99	245.40	189.91	871.17	846.63	662.58	417.18
100000.00	20.90	270.10	209.03	1012.86	958.84	729.26	459.16
500000.00	35.75	461.83	357.48	1824.24	1731.87	1246.95	785.11
******	*******	******	******	*****	*****	*****	***

PROPANE RESULTS OF ISO-THERMAL MODEL:

FIREBALL SURFACE TEMPERATURE = 2113.337

MASS	TIME	DIAMETER	HEIGHT	HUMAN	R. ANIMAL	R. BULID	R PR.EQ.P	2
1000.00	4.50	57.79	45.02	147.37	150.26	161.81	98.24	
5000.00	7.70	98.82	76.99	281.62	281.62	276.68	167.99	
10000.00	9.70	124.50	97.01	379.71	392.16	348.59	211.64	
25000.00	13.17	168.96	131.67	532.23	549.13	473.10	287 24	
50000.00	16.59	212.87	165.90	691.84	734.42	596.05	361 89	
75000.00	18.99	243.68	189.91	840.69	865.06	682.30	414 25	
100000.00	20.90	268.20	209.03	965.52	1005.75	750.96	455 94	
500000.00	35.75	458.59	357.48	1765.57	1765.57	1284.05	779 60	
******	*****	*****	****	*****	*****	********	***	

BUTANE RESULTS OF ISO-THERMAL MODEL:

FIREBALL SURFACE TEMPERATURE = 2120.308

MASS	TIME	DIAMETER	HEIGHT	HUMAN	R. ANIMAL	R. BULID	.R PR.EQ.R
1000.00	4.50	57.54	45.02	149.61	143.86	155.36	97 82
5000.00	7.70	98.39	76.99	275.49	260.74	265.66	167.26
10000.00	9.70	123.96	97.01	359.49	365.69	334.70	210.74
25000.00	13.17	168.24	131.67	529.95	521.53	454.24	286.00
50000.00	16.59	211.96	165.90	710.07	688.87	572.29	360.33
75000.00	18.99	242.63	189.91	861.34	824.94	655.10	412.47
100000.00	20.90	267.05	209.03	988.07	948.02	721.03	453.98
500000.00	35.75	456.62	357.48	1757.99	1666.66	1232.87	776.25
******	******	******	******	*****	*****	*******	***

PENTANE RESULTS OF ISO-THERMAL MODDEL:

FIREBALL SURFACE TEMPERATURE = 2127.224

MASS	TIME	DIAMETER	HEIGHT	HLMAN	R. ANIMAL	R. BULID	R PR.EQ.I	R
1000.00	4.50	57.41	45.02	149.26	149.26	155.00	97.60	
5000.00	7.70	98.16	76.99	279.77	274.86	265.04	166.88	
10000.00	9.70	123.68	97.01	383.39	371.03	333.92	210.25	
25000.00	13.17	167.85	131.67	537.11	528.72	453.19	285.34	
50000.00	16.59	211.47	165.90	708.42	687.28	570.97	359.50	
75000.00	18.99	242.07	189.91	835.14	823.04	653.59	411.52	
100000.00	20.90	266.43	209.03	945.82	932.50	719.36	452.93	
500000.00	35.75	455.56	357.48	1685.58	1662.81	1230.02	774.46	
******	*****	*****	******	******	******	******	***	

B.L.E.V.E FIREBALL RESULTS:

METHANE BLEVE FIREBALL RESULTS:

FIREBALL SURFACE TEMPERATURE = 1207.654

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID	.R. PR.EQ.R
1000.00	4.37	48.47	24.24	53.32	55.74	43.63	2 42
5000.00	7.47	82.88	41.44	103.60	107.75	74.59	4 14
10000.00	9.41	104.42	52.21	140.97	146.19	93.98	5 22
25000.00	12.77	141.72	70.86	205.49	212.58	127.55	7 09
50000.00	16.09	178.55	89.28	276.75	294.61	160.70	8 93
75000.00	18.42	204.39	102.19	337.24	357.68	183.95	10.22
100000.00	20.27	224.96	112.48	393.67	416.17	202.46	11 25
500000.00	34.66	384.65	192.32	711.60	769.30	346.18	19.23

ETHANE BLEVE FIREBALL RESULTS:

FIREBALL SURFACE TEMPERATURE = 1219.642

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL R.	BULID.R.	PR.EQ.R
1000.00	3.54	39.31	19.66	39.31	41.28	35.38	1.97
5000.00	6.06	67.22	33.61	77.30	80.66	60.49	3.36
10000.00	7.63	84.68	42.34	105.86	110.09	76.22	4.23

25000.00	10.36	114.93	57.47	155.16	160.90	103.44	5.75
50000.00	13.05	144.80	72.40	209.96	217.20	130.32	7.24
75000.00	14.94	165.75	82.88	256.92	265.21	149.18	8.29
100000.00	16.44	182.43	91.22	301.02	319.26	164.19	9.12
500000.00	28.11	311.94	155.97	577.09	592.69	280.75	15.60

PROPANE BLEVE FIREBALL RESULTS:

FIREBALL SURFACE TEMPERATURE = 1220.490

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL	R. BULID.R.	PR.EQ.R
1000.00	3.12	34.60	17.30	36.33	38.06	31.14	1.73
5000.00	5.33	59.16	29.58	68.03	70.99	53.24	2.96
10000.00	6.72	74.54	37.27	93.17	96.90	67.08	3 73
25000.00	9.12	101.16	50.58	136.56	141.62	91.04	5.06
50000.00	11.49	127.45	63.72	184.80	191.17	114.70	6.37
75000.00	13.15	145.89	72.94	226.13	233.42	131.30	7 29
100000.00	14.47	160.57	80.29	264.94	272.97	144.51	8 03
500000.00	24.74	274.56	137.28	480.48	494.20	247.10	13.73

BUTANE BLEVE FIREBALL RESULTS:

FIREBALL SURFACE TEMPERATURE = 1220.024

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL R	. BULID.R.	PR.EQ.R
1000.00	2.84	31.56	15.78	33.13	33.13	28.40	1 58
5000.00	4.86	53.96	26.98	62.05	62.05	48.56	2.70
10000.00	6.13	67.98	33.99	84.97	88.37	61.18	3.40
25000.00	8.31	92.26	46.13	124.55	129.16	83.03	4.61
50000.00	10.47	116.24	58.12	168.54	174.36	104.61	5.81
75000.00	11.99	133.06	66.53	206.24	212.89	119.75	6.65
100000.00	13.20	146.45	73.22	241.64	248.96	131.80	7.32
500000.00	22.57	250.41	125.20	425.69	438.21	225.37	12.52

PENTANE BLEVE FIREBALL RESULTS:

FIREBALL SURFACE TEMPERATURE = 1220.514

MASS	TIME	DIAMETER	HEIGHT	HUMAN R.	ANIMAL R.	BULID.R.	PR.EQ.R
1000.00	2.65	29.36	14.68	29.36	27.89	26.43	1.47
5000.00	4.52	50.21	25.10	55.23	52.72	45.18	2.51
10000.00	5.70	63.25	31.63	75.90	72.74	56.93	3.16
25000.00	7.74	85.84	42.92	111.60	107.31	77.26	4.29
50000.00	9.75	108.16	54.08	151.42	146.01	97.34	5.41
75000.00	11.16	123.81	61.90	185.71	179.52	111.42	6.19
100000.00	12.28	136.26	68.13	218.02	211.21	122.64	6.81
500000.00	21.00	233.00	116.50	384.44	384.44	209.70	11.65

APPENDIX H

Computer program for evaluation of BLEVE's fireball diameter and duration using different authors' parameters

```
C
C PROGRAM FOR EVALUATION OF BLEVE FIREBALL
C ------
C USING MAURER, TND, CRAWLEY, AND LIHOU EQUATIONS:
C ===
С
C
С
      DIMENSION W(30), DM(30), DT(30), DC(30), DL(30), TM(30), TT(30), TL(30)
C
    READ *, NN
C
      READ *, (W(11), 11=1, NN)
С
      DO 3 1=1.NN
       DM(1)=3.51*(W(1)**0.333)
       DT(1)=6.48*(W(1)**0.325)
       DC(1)=105.0*((W(1)/(1000.0*44.0))**0.333)
       DL(1)=3.44*((44.8*W(1)/44.0)**0.333)
       TM(1)=0.32*(W(1)**0.333)
       TT(1)=0.85*(W(1)**0.26)
       TL(1)=0.31*((44.8*W(1)/44.0)**0.333)
С
  3
         CONTINUE
C
C
      PRINT 100
 100 FORMAT(//10X, 'BLEVE PROPANE FIREBALL DIAMETER & DURATION: ',/10X,
     @43('-')//)
C
C
        PRINT 200
 200 FORMAT(11X, 'MASS
                           FIREBALL DIAMETER
                                                    FIREBALL DURATION ',
     @/,11X,'(KG) MAURER TNO CRAWLEY LIHOU MAURER TNO
                                                               LIHOU ',
     @/,10X,55('-'))
C
С
      DO 5 J=1,NN
      PRINT 300, W(J), DM(J), DT(J), DC(J), DL(J), TM(J), TT(J), TL(J)
 300 FORMAT(5X,F9.1,7(F6.1,1X))
  5
       CONTINUE
C
С
        STOP
        END
```

MAS (KG	S) MAUF	FIREBA	LL DIAN CRAW	ETER	FIRE HOU MAL	BALL DU	RATION	UCI
0.1	1.6	3.1	1.4	1.6	0.1	0.5	0.1	-
1.0	3.5	6.5	3.0	3.5	0.3	0.9	0.3	
5.0	6.0	10.9	5.1	5.9	0.5	1.3	0.5	
10.0	7.6	13.7	6.4	7.5	0.7	1.5	0.7	
20.0	9.5	17.2	8.1	9.4	0.9	1.9	0.8	
50.0	12.9	23.1	11.0	12.7	1.2	2.4	1.1	
75.0	14.8	26.4	12.6	14.6	1.3	2.6	1.3	
100.0	16.3	28.9	13.8	16.0	1.5	2.8	1.4	
250.0	22.1	39.0	18.8	21.8	2.0	3.6	2.0	
500.0	27.8	48.8	23.6	27.4	2.5	4.3	2.5	
1000.0	35.0	61.2	29.8	34.5	3.2	5.1	3.1	
5000.0	59.9	103.2	50.9	59.0	5.5	7.8	5.3	
10000.0	75.4	129.3	64.1	74.3	6.9	9.3	6.7	
25000.0	102.3	174.1	87.0	100.9	9.3	11.8	9.1	
50000.0	128.8	218.1	109.6	127.0	11.7	14.2	11.4	
75000.0	147.5	248.9	125.4	145.4	13.4	15.7	13.1	
100000.0	162.3	273.3	138.0	160.0	14.8	17.0	14.4	
250000.0	220.2	368.0	187.3	217.1	20.1	21.5	19.6	
500000.0	277.4	461.0	235.9	273.5	25.3	25.8	24.6	
1000000.0	349.4	577.5	297.1	344.5	31.9	30.9	31.0	

BLEVE PROPANE FIREBALL DIAMETER & DURATION:

APPENDIX I

Computer program for calculation of pool fires on vertical cylinders

```
С
C
C PROGRAM FOR CALCULATION OF POOL FIRES ON VERTICAL CYLINDERS:
C =
C
С
С
       DIMENSION RLD(99), WR(99), E(99), DS(99), Q1(99),
     @R(99), AREA(99), Q(99), Q2(99), DSH(99), DSB(99), DSPE(99)
C
C
       READ *, FC, SECMA, THETA
C
       PRINT 100
 100 FORMAT(//10X, 'POOL FIRE HAZARD DISTANCES RESULTS: ', /, 10X,
      @35('-')/)
C
C
       PRINT 110
 110 FORMAT(10X, 'PROPANE RESULTS: ', /, 10X, 16('-'))
С
       CALL VERT (FC, SECMA, THETA, E, DS, Q1, Q2)
C
C
       PRINT 120
 120
      FORMAT(10X, 'BUTANE RESULTS: ',/,10X,16('-')/)
C
      CALL VERT (FC, SECMA, THETA, E, DS, Q1, Q2)
C
C
С
         STOP
          END
С
C
C
С
C SUBROUTINE FOR CALCULATION OF POOL FIRE ON VERTICAL CYLINDER.
С
С
С
      SUBROUTINE VERT(F, SEOMA, THETA, E, DS, Q1, Q2)
C
      DIMENSION RLD(99), WR(99), RAT(99), DS(99), E(99), Q1(99)
     @, AREA(99), Q(99), R(99), Q2(99), DSH(99), DSB(99), DSPE(99)
С
C
           Q2H=4.7
          Q2B=12.6
          Q2PE=37.8
С
       READ *, J, N
       READ *, (DS(NN), NN=1, J)
       READ *, (RLD(||), ||=1,N)
```

```
READ *, TEMP, HC, TRW, CP
С
          PRINT 105, TEMP
  105
          FORMAT(//10X, 'AT FLAME TEMPERATURE =', F10.3, /10X, 22('-'))
С
С
С
      DO 5 M=1, J
      PRINT 220, M, DS(M)
      FORMAT(//, 10X, 'DIAMETER OF VERTICAL CYLINDER(', 12, ')=', F10.3, /,
 220
     @10X,34('-')//)
С
С
      PRINT 210
 210 FORMAT(1X,'
                            HUMAN DIST. BULID.DIST. ',
                   L/D
     @' PROCESS DSIT. WATER RATE',/3X,60('-'))
С
С
      DO 3 I=1,N
      E(1)=SECMA*THETA*(TEMP**4.0)
      R(1)=(E(1)/(F*HC))*(1.0+(4.0*RLD(1)))
      Q1(1)=F*HC*R(1)*(RLD(1)*(DS(M)**2.0))
      DSH(1)=(Q1(1)/(4.0*3.14159*Q2H))**0.5
      DSB(1)=(Q1(1)/(4.0*3.14159*Q2B))**0.5
      DSPE(1)=(Q1(1)/(4.0*3.14159*Q2PE))**0.5
      AREA(1)=(RLD(1)*(DS(M)**2.0)*3.14159)/2.0
      Q(1)=Q1(1)*AREA(1)
      WR(1)=Q(1)/(CP*TRW*60.0)
      PRINT 200, RLD(1), DSH(1), DSB(1), DSPE(1), WR(1)
 200
        FORMAT(F8.2, 1X, 3(F10.3, 2X), 6X, F10.2)
С
  3
       CONTINUE
  5
         CONTINUE
C
           PRINT 250
 250
          FORMAT(10X, //)
        RETURN
         END
```

POOL FIRE HAZARD DISTANCES RESULTS:

PROPANE RESULTS:

AT FLAME TEMPERATURE = 2113.000

DIAMETER OF VERTICAL CYLINDER(1)= 1.000

L/D	HUMAN DIST.	BULID.DIST.	PROCESS DSIT.	WATER RATE
1.00	9.782	5.974	3.449	0.50
1.20	11.541	7.049	4.070	0.84
1.40	13.298	8.122	4.689	1.30
1.60	15.053	9.193	5.308	1.91
1.80	16.807	10.265	5.926	2.68
2.00	18.560	11.335	6.544	3.63
2.20	20.312	12.406	7.162	4.78
2.40	22.065	13.476	7.780	6 16
2.60	23.816	14.546	8.398	7 77
2.80	25.568	15.616	9.016	9.65
3.00	27.319	16.685	9.633	11.80

DIAMETER OF VERTICAL CYLINDER(2)= 2.000

L/D	HUMAN DIST.	BULID.DIST.	PROCESS DSIT.	WATER RATE
1.00	19.564	11.949	6.899	8.07
1.20	23.082	14.097	8.139	13.48
1.40	26.595	16.243	9.378	20.88
1.60	30.105	18.387	10.616	30.57
1.80	33.613	20.529	11.853	42.88
2.00	37.120	22.671	13.089	58,10
2.20	40.625	24.812	14.325	76.55
2.40	44.129	26.952	15.561	98.53
2.60	47.633	29.092	16.796	124.37
2.80	51.136	31.231	18.031	154.36
3.00	54.639	33.371	19.266	188.82

DIAMETER OF VERTICAL CYLINDER(3)= 3.000

L/D	HUMAN DIST.	BULID.DIST.	PROCESS DSIT.	WATER RATE
1.00	29.346	17.923	10.348	40.85
1.20	34.623	21.146	12.209	68.24
1.40	39.893	24.365	14.067	105.69
1.60	45.158	27.580	15.923	154.77
1.80	50.420	30.794	17.779	217.06
2.00	55.679	34.006	19.633	294.12
2.20	60.937	37.217	21.487	387.52
2.40	66.194	40.428	23.341	498.82
2.60	71.449	43.638	25.194	629.61
2.80	76.704	46.847	27.047	781.44
3.00	81.958	50.056	28.900	955.88

DIAMETER OF VERTICAL CYLINDER(4)= 4.000

L/D	HUMAN DIST.	BULID.DIST.	PROCESS DSIT.	WATER RATE
1.00	39.127	23.897	13.797	129.11
1.20	46.164	28.195	16.278	215.66
1.40	53.190	32.486	18.756	334.02
1.60	60.211	36.774	21.231	489.15
1.80	67.226	41.059	23.705	686.01
2.00	74.239	45.342	26.178	929.56
2.20	81.250	49.623	28.650	1224.74
2.40	88.258	53.904	31.121	1576.53
2.60	95.266	58.183	33.592	1989.87
2.80	102.272	62.463	36.063	2469.73
3.00	109.277	66.741	38.533	3021.06

DIAMETER	OF	VERTICAL	CYLINDER(5)=	5.000
----------	----	----------	-----------	-----	-------

L/D	HUMAN DIST.	BULID.DIST.	PROCESS DSIT.	WATER RATE
1.00	48.909	29.871	17.246	315.20
1.20	57.705	35.243	20.348	526.51
1.40	66.488	40.608	23.445	815.48
1.60	75.263	45.967	26.539	1194.22
1.80	84.033	51.323	29.631	1674.84
2.00	92.799	56.677	32.722	2269.43
2.20	101.562	62.029	35.812	2990.10
2.40	110.323	67.380	38.902	3848.95
2.60	119.082	72.729	41.990	4858.09
2.80	127.840	78.078	45.078	6029.61
3.00	136.596	83.426	48.166	7375.64

DIAMETER OF VERTICAL CYLINDER(6)= 6.000

L/D	HUMAN DIST.	BULID.DIST.	PROCESS DSIT.	WATER RATE
1.00	58.691	35.846	20.696	653.59
1.20	69.246	42.292	24.417	1091.76
1.40	79.786	48.729	28.134	1690.98
1.60	90.316	55.160	31.847	2476.34
1.80	100.840	61.588	35.558	3472.94
2.00	111.359	68.012	39.267	4705.88
2.20	121.874	74.435	42.975	6200.26
2.40	132.387	80.856	46.682	7981.18
2.60	142.898	87.275	50.388	10073.73
2.80	153.408	93.694	54.094	12503.01
3.00	163.916	100.112	57.799	15294.12

DIAMETER OF VERTICAL CYLINDER(7)= 7.000

L/D	HUMAN DIST.	BULID.DIST.	PROCESS DSIT.	WATER RATE
1.00	68.473	41.820	24.145	1210.87
1.20	80.787	49.340	28.487	2022.63
1.40	93.083	56.851	32.823	3132.75
1.60	105.368	64.354	37.155	4587.73
1.80	117.646	71.852	41.484	6434.05
2.00	129.919	79.348	45.811	8718.23
2.20	142.187	86.841	50.137	11486.75
2.40	154.452	94.332	54.462	14786.12
2.60	166.715	101.821	58.786	18662 82
2.80	178.976	109.309	63.110	23163.37
3.00	191.235	116.797	67.433	28334.24

DIAMETER OF VERTICAL CYLINDER(8)= 8.000

L/D	HUMAN DIST.	BULID.DIST.	PROCESS DSIT.	WATER RATE
1.00	78.255	47.794	27.594	2065.68
1.20	92.328	56.389	32.556	3450.52
1.40	106.381	64.972	37.512	5344.33
1.60	120.421	73.547	42.463	7826.46
1.80	134.453	82.117	47.410	10976.21
2.00	148.478	90.683	52.356	14872.91
2.20	162.499	99.246	57.300	19595.89
2.40	176.517	107.808	62.243	25224.46
2.60	190.531	116.367	67.184	31837.95
2.80	204.544	124.925	72.125	39515 68
3.00	218.554	133.482	77.066	48336.97

DIAMETER OF VERTICAL CYLINDER(9)= 9.000

L/D	HUMAN DIST.	BULID.DIST.	PROCESS DSIT.	WATER RATE
1.00	88.037	53.769	31.043	3308.82
1.20	103.869	63.438	36.626	5527.06
1.40	119.678	73.094	42.201	8560.59
1.60	135.474	82.741	47.770	12536.47
1.80	151.259	92.382	53.337	17581.77
2.00	167.038	102.019	58.900	23823.53
2.20	182.812	111.652	64.462	31388.83
2.40	198.581	121.283	70.023	40404.71
2.60	214.348	130.913	75.583	50998.24
2.80	230.111	140.541	81.141	63296.48
3.00	245.873	150.167	86.699	77426.48

DIAMETER OF VERTICAL CYLINDER(10)= 10.000

L/D	HUMAN DIST.	BULID.DIST.	PROCESS DSIT.	WATER RATE
1.00	97.819	59.743	34.493	5043.17
1.20	115.409	70.486	40.695	8424.11
1.40	132.976	81.215	46.890	13047.69
1.60	150.526	91.934	53.078	19107.56
1.80	168.066	102.646	59.263	26797.39
2.00	185.598	113.354	65.445	36310.82
2.20	203.124	124.058	71.625	47841.53
2.40	220.646	134.759	77.803	61583.16
2.60	238.164	145.459	83.981	77729.37
2.80	255.679	156.156	90.157	96473.83
3.00	273.193	166.853	96.332	118010.18

BUTANE RESULTS:

AT FLAME TEMPERATURE = 2120.000

DIAMETER OF VERTICAL CYLINDER(1)= 1.000

L/D HUMAN DIST. BULID.DIST. PROCESS DSIT. WATER RATE

1.00	9.847	6.014	3.472	0.51
1.20	11.618	7.095	4.097	0.85
1.40	13.386	8.175	4.720	1.32
1.60	15.153	9.254	5.343	1.94
1.80	16.918	10.333	5.966	2.72
2.00	18.683	11.411	6.588	3.68
2.20	20.447	12.488	7.210	4.85
2.40	22.211	13.565	7.832	6.24
2.60	23.974	14.642	8.454	7.88
2.80	25.738	15.719	9.076	9.78
3.00	27.501	16.796	9.697	11.96

DIAMETER OF VERTICAL CYLINDER(2)= 2.000

L/D	HUMAN DIST.	BULID.DIST.	PROCESS DSIT.	WATER RATE
1.00	19.694	12.028	6.944	8,18
1.20	23.235	14.191	8.193	13.66
1.40	26.772	16.351	9.440	21.15
1.60	30.305	18.509	10.686	30.98
1.80	33.836	20.666	11.931	43.45
2.00	37.366	22.821	13.176	58.87
2.20	40.894	24.976	14.420	77.57
2.40	44.422	27.131	15.664	99.85
2.60	47.949	29.285	16.908	126.02
2.80	51.475	31.439	18.151	156.41
3.00	55.001	33.592	19.394	191.33

DIAMETER OF VERTICAL CYLINDER(3)= 3.000

L/D	HUMAN DIST.	BULID.DIST.	PROCESS DSIT.	WATER RATE
1.00	29.540	18.042	10.416	41.39
1.20	34.853	21.286	12.290	69.14
1.40	40.158	24.526	14.160	107.09
1.60	45.458	27.763	16.029	156.83
1.80	50.754	30.998	17.897	219.95
2.00	56.049	34.232	19.764	298.03
2.20	61.342	37.464	21.630	392.68
2.40	66.633	40.696	23.496	505.47
2.60	71.923	43.927	25.361	637.99
2.80	77.213	47.158	27.227	791.84
3.00	82.502	50.388	29.092	968.61

DIAMETER OF VERTICAL CYLINDER(4)= 4.000

390

270	HUMAN DIST.	BULID.DIST.	PROCESS DSIT.	WATER RATE
1.00	39.387	24.056	13.889	130.82
1.20	46.470	28.382	16.386	218.53
1.40	53.543	32.702	18.880	338.47
1.60	60.610	37.018	21.372	495.67
1.80	67.673	41.331	23.862	695.15
2.00	74.732	45.643	26.352	941.94
2.20	81.789	49.953	28.840	1241.05
2.40	88.844	54.261	31.328	1597.52
2.60	95.898	58.570	33.815	2016.37
2.80	102.951	62.877	36.302	2502.62
3.00	110.002	67.184	38.789	3061.29
	DIAMETER OF V	ERTICAL CYLIN	DER(5)= 5.	000

L/D	HUMAN DIST.	BULID.DIST.	PROCESS DSIT.	WATER RATE
1.00	49.234	30.070	17.361	319.40
1.20	58.088	35.477	20.483	533.52
1.40	66.929	40.877	23.600	826.34
1.60	75.763	46.272	26.715	1210.13
1.80	84.591	51.664	29.828	1697.14
2.00	93.415	57.053	32,940	2299.65
2.20	102.236	62.441	36.050	3029.92
2.40	111.055	67.827	39.160	3900.20
2.60	119.872	73.212	42.269	4922 78
2.80	128.688	78.596	45.378	6109 91
3.00	137.503	83.980	48.486	7473.86

L/D

DIAMETER C	F	VERTICAL	CYLINDER(6)=	6.000
------------	---	----------	-----------	-----	-------

L/D	HUMAN DIST.	BULID.DIST.	PROCESS DSIT.	WATER RATE	
1.00	59.081	36.084	20.833	662.30	
1.20	69.705	42.572	24.579	1106.30	
1.40	80.315	49.052	28.320	1713.50	
1.60	90.915	55.527	32.058	2509.32	
1.80	101.509	61.997	35.794	3519.19	
2.00	112.098	68.464	39.528	4768.55	
2.20	122.683	74.929	43.260	6282.83	
2.40	133.266	81.392	46.992	8087.46	
2.60	143.847	87.854	50.723	10207.88	
2.80	154.426	94.316	54.453	12669.51	
3.00	165.003	100.776	58.183	15497.80	

DIAMETER OF VERTICAL CYLINDER(7)= 7.000

391

L/D	HUMAN DIST.	BULID.DIST.	PROCESS DSIT.	WATER RATE
1.00	68.928	42.097	24.305	1226.99
1.20	81.323	49.668	28.676	2049.57
1.40	93.701	57.228	33.041	3174.47
1.60	106.068	64.781	37.401	4648.82
1.80	118.427	72.329	41.759	6519.74
2.00	130.781	79.874	46.115	8834.33
2.20	143.130	87.417	50.470	11639.72
2.40	155.477	94.958	54.824	14983.03
2.60	167.821	102.497	59.177	18911.36
2.80	180.163	110.035	63.529	23471.84
3.00	192.504	117.572	67.880	28711.58

DIAMETER OF VERTICAL CYLINDER(8)= 8.000

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L/D	HUMAN DIST.	BULID.DIST.	PROCESS DSIT.	WATER RATE
1.00	78.774	48.111	27.777	2093.19
1.20	92.940	56.763	32.772	3496.47
1.40	107.087	65.403	37.761	5415.51
1.60	121.220	74.035	42.744	7930.68
1.80	135.345	82.662	47.725	11122.38
2.00	149.464	91.285	52.703	15070.98
2.20	163.578	99.905	57.680	19856.85
2.40	177.688	108.523	62.656	25560.38
2.60	191.796	117.139	67.630	32261.95
2.80	205.901	125.754	72.604	40041.92
3.00	220.005	134.368	77.577	48980.69

DIAMETER OF VERTICAL CYLINDER(9)= 9.000

L/D	HUMAN DIST.	BULID.DIST.	PROCESS DSIT.	WATER RATE
1.00	88.621	54.125	31.249	3352.89
1.20	104.558	63.859	36.869	5600.66
1.40	120.473	73.579	42.481	8674.59
1.60	136.373	83.290	48.087	12703.42
1.80	152.263	92.995	53.691	17815.91
2.00	168.147	102.696	59.291	24140.80
2.20	184.025	112.393	64.890	31806.84
2.40	199.899	122.088	70.488	40942.79
2.60	215.770	131.782	76.084	51677.40
2.80	231.639	141.473	81.680	64139.41
3.00	247.505	151.164	87.275	78457.59

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DIAMETER OF VERTICAL CYLINDER(10)= 10.000

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L/D	HUMAN DIST.	BULID.DIST.	PROCESS DSIT.	WATER RATE
1.00	98.468	60.139	34.721	5110.33
1.20	116.175	70.954	40.965	8536.30
1.40	133.858	81.754	47.201	13221.45
1.60	151.525	92.544	53.430	19362.02
1.80	169.181	103.328	59.656	27154.26
2.00	186.830	114.106	65.879	36794.39
2.20	204.472	124.881	72.100	48478.65
2.40	222.110	135.654	78.320	62403.28
2.60	239.745	146.424	84.538	78764.52
2.80	257.376	157.193	90.755	97758.60
3.00	275.006	167.960	96.972	119581.76

APPENDIX J

Computer program for calculation of spacing distance using TNT equivalent model (Jarrett's equation)

```
С
C
C PROGRAM FOR CALCULATION OF SPACING DISTANCE
C =
C USING T.N.T. EQUIVALENT MODEL (JARRETT'S EQUATION)
         C ===
С
      PRINT 333
      FORMAT(//15X, 'JARRETT EQUATION RESULTS: ', /15X, 25( '*')//)
 333
С
C
      PRINT 20
 20
      FORMAT(10X, 'METHANE RESULTS: ', /, 10X, 15( '*') //)
С
      READ *, CH4
      CALL PR(CH4)
С
      PRINT 25
 25
      FORMAT(///, 10X, 'PROPANE RESULTS: ', /, 10X, 15('*')//)
C
      READ *, CH8
      CALL PR(CH8)
C
C
      PRINT 30
 30
      FORMAT(///, 10X, 'BUTANE RESULTS: ', /, 10X, 15( '*')//)
C
       READ *, CH10
       CALL PR(CH10)
С
С
         STOP
         END
С
С
C
       SUBROUTINE PR(C)
С
C
С
      DIMENSION W(99), CH(99), R(99), TNT(99), FK(99), Y(99)
С
C
      READ *, (CH(11), 11=1,6)
      READ *, (FK(NN), NN=1, 15)
      READ *, (Y(JJ), JJ=1,3)
С
C
       DO 5 J=1,3
С
      PRINT 100, J, Y(J)
 100 FORMAT(10X, 'EXPLOSION YIELD FACTOR(', 12, ')=', F10.3, '%',
     */,10X,26('-')///)
C
```

```
395
```

```
TNT(J) = (1100.0*100.0)/(C*Y(J))
С
       PRINT 150, J, TNT(J)
      FORMAT(10X, 'TNT EQUIVALENT VALUE (', 12, ')=', F10.4//)
 150
C
C
       DO 10 N=1,15
С
      PRINT 200, N, FK(N)
 200 FORMAT(10X, 'CONSTANT OF JARRETT EQUATION (', 12, ')=', F10.3,
     */,10X,33('-')/)
C
      PRINT 300
 300
      FORMAT(10X, 'W.OF H.C. W.OF TNT
                                           DISTANCE' . /
     @, 10X, ' TONS
                         TONS
                                     METERS', /, 10X, 32('='))
С
С
C
         DO 15 I=1.6
      W(1)=CH(1)/TNT(J)
      R(1)=0.3048*(FK(N)*((W(1)*2200.0)**0.333))/
     @((1.0+((6996.0/(W(1)*2200.0))**2.0))**0.1667)
C
      PRINT 400, CH(1), W(1), R(1)
 400
      FORMAT(8X,3F10.3)
C
  15
        CONTINUE
C
       PRINT 500
 500
       FORMAT(8X,35('*')///)
        CONTINUE
  10
С
   5
        CONTINUE
С
         RETURN
         END
С
```


METHANE RESULTS:

EXPLOSION YIELD FACTOR(1)= 2.000%

TNT EQUIVALENT VALUE (1)= 4.5891

CONSTANT OF JARRETT EQUATION (1)= 3.000

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	1.090	8.384
10.000	2.179	12.712
20.000	4.358	18.036
30.000	6.537	21.395
40.000	8.716	23.892
50.000	10.895	25.921
*****	*****	**********

CONSTANT OF JARRETT EQUATION (2)= 9.000

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	1.090	25.151
10.000	2.179	38.136
20.000	4.358	54.109
30.000	6.537	64.185
40.000	8.716	71.675
50.000	10.895	77.762
******	******	*****

**

CONSTANT OF JARRETT EQUATION (3)= 10.050

W.OF H.C. W.OF TNT DISTANCE

	TONS	TONS	METERS	
	5.000	1.090	28.086	-
	10.000	2.179	42.585	
	20.000	4.358	60.422	
	30.000	6.537	71.674	
	40.000	8.716	80.037	
	50.000	10.895	86.834	
* *	*******	*****	*****	*

CONSTANT OF JARRETT EQUATION (4)= 13.900

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	1.090	38.845
10.000	2.179	58.898
20.000	4.358	83.569
30.000	6.537	99.131
40.000	8.716	110.698
50.000	10.895	120.099
*****	******	*******

CONSTANT OF JARRETT EQUATION (5)= 16.300

	W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
	5.000	1.090	45.552
	10.000	2.179	69.068
	20.000	4.358	97.998
	30.000	6.537	116.247
	40.000	8.716	129.812
	50.000	10.895	140.836
1 3	*********	******	

CONSTANT OF JARRETT EQUATION (6)= 18.500

N	.OF H.C. TONS	W.OF TNT TONS	DISTANCE METERS
	5.000	1.090	51.700
	10.000	2.179	78.390
	20.000	4.358	111.224
	30.000	6.537	131.936

40.000	8.716	147.332
50.000	10.895	159.844
*****	*******	****

CONSTANT OF JARRETT EQUATION (7)= 19.800

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	1.090	55.333
20.000	4.358	119.040
30.000	6.537	141.208
40.000	8.716	157.686
50.000	10.895	171.077
*****	********	*********

CONSTANT OF JARRETT EQUATION (8)= 21.600

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	1.090	60.363
10.000	2.179	91.526
20.000	4.358	129.862
30.000	6.537	154.045
40.000	8.716	172.021
50.000	10.895	186.629
*****	*****	*****

CONSTANT OF JARRETT EQUATION (9)= 22.800

	W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
	5.000	1.090	63.716
	10.000	2.179	96.610
	20.000	4.358	137.077
	30.000	6.537	162.603
	40.000	8.716	181.577
	50.000	10.895	196.998
•	*******	*****	******

CONSTANT OF JARRETT EQUATION (10)= 25.500

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	1.090	71.262
10.000	2.179	108.051
20.000	4.358	153.309
30.000	6.537	181.858
40.000	8.716	203.080
50.000	10.895	220.326

CONSTANT OF JARRETT EQUATION (11)= 37.000

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	1.090	103.400
10.000	2.179	156.780
20.000	4.358	222.449
30.000	6.537	263.873
40.000	8.716	294.665
50.000	10.895	319.689
*******	******	*****

CONSTANT OF JARRETT	EQUATION	(12)=	51.500
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W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	1.090	143.921
10.000	2.179	218.221
20.000	4.358	309.625
30.000	6.537	367.282
40.000	8.716	410.142
50.000	10.895	444.972
******	*******	******

*

CONSTANT	OF	JARRETT	EQUATION	(13)=	61.400

W.OF H.C.	W.OF TNT	DISTANCE
TONS	TONS	METERS

	5.000	1.090	171.587	
	10.000	2.179	260.170	
	20.000	4.358	369.145	
	30.000	6.537	437.886	
	40.000	8.716	488.984	
	50.000	10.895	530.511	
**	*******	******	*****	*

CONSTANT OF JARRETT EQUATION (14)= 72.000

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	1.090	201.210
10.000	2.179	305.085
20.000	4.358	432.874
30.000	6.537	513.482
40.000	8.716	573.402
50.000	10.895	622.097
******	*****	*****

**

CONSTANT OF JARRETT EQUATION (15)= 99.000

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	1.090	276.664
10.000	2.179	419.492
20.000	4.358	595.201
30.000	6.537	706.038
40.000	8.716	788.428
50.000	10.895	855.384
*******	*****	*****

EXPLOSION YIELD FACTOR(2)= 5.000%

TNT EQUIVALENT VALUE (2)= 1.8356

CONSTANT OF JARRETT EQUATION (1)= 3.000

W.OF H.C. W.OF TNT DISTANCE

	TONS	TONS	METERS	
100	5.000	2.724	14.349	:
	10.000	5.448	19.865	
	20.000	10.895	25.921	
	30.000	16.343	29.889	
	40.000	21.791	32.982	
	50.000	27.239	35.571	
*	******	*****	*****	

CONSTANT OF JARRETT	EQUATION	(2)=	9.000
---------------------	----------	------	-------

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	2.724	43.047
10.000	5.448	59.596
20.000	10.895	77.762
30.000	16.343	89.667
40.000	21.791	98.947
50.000	27.239	106.714
******	*****	*****

CONSTANT OF JARRETT EQUATION (3)= 10.050

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	2.724	48.069
10.000	5.448	66.549
20.000	10.895	86.834
30.000	16.343	100.129
40.000	21.791	110.491
50.000	27.239	119.164
******	******	*****

CONSTANT OF JARRETT EQUATION (4)= 13.900

N.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	2.724	66.484
10.000	5.448	92.042
20.000	10.895	120.099
30.000	16.343	138.486

40.000	21.791	152.819
50.000	27.239	164.814
*****	******	

CONSTANT OF JARRETT EQUATION (5)= 16.300

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	2.724	77.963
10.000	5.448	107.935
20.000	10.895	140.836
30.000	16.343	162.398
40.000	21.791	179.205
50.000	27.239	193.271
******	******	****

18.500	CONSTANT	OF	JARRETT	EQUATION (6)=	18.500
--------	----------	----	---------	------------	-----	--------

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	2.724	88.486
10.000	5.448	122.502
20.000	10.895	159.844
30.000	16.343	184.317
40.000	21.791	203.392
50.000	27.239	219.356
*****	*****	*****

CONSTANT OF JARRETT EQUATION (7)= 19.800

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	2.724	94.704
10.000	5.448	131.111
20.000	10.895	171.077
30.000	16.343	197.268
40.000	21.791	217.684
50.000	27.239	234.771
*****	*****	*****

CONSTANT OF JARRETT EQUATION (8)= 21.600

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	2.724	103.313
10.000	5.448	143.030
20.000	10.895	186.629
30.000	16.343	215.202
40.000	21.791	237.474
50.000	27.239	256.113
********	********	*****

CONSTANT OF JARRETT EQUATION (9)= 22.800

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	2.724	109.053
10.000	5.448	150.976
20.000	10.895	196.998
30.000	16.343	227.158
40.000	21.791	250.667
50.000	27.239	270.342
*******	******	*****

CONSTANT OF JARRETT EQUATION (10)= 25.500

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	2.724	121.967
10.000	5.448	168.855
20.000	10.895	220.326
30.000	16.343	254.058
40.000	21.791	280.351
50.000	27.239	302.356
****	*******	*****

(11) = 37.00	CONSTANT	OF	JARRETT	EQUATION	(11)=.	37.00
--------------	----------	----	---------	----------	-----	-----	-------

W.OF H.C.	W.OF TNT	DISTANCE
TONS	TONS	METERS

5.000	2.724	176.972
10.000	5.448	245.005
20.000	10.895	319.689
30.000	16.343	368.633
40.000	21.791	406.784
50.000	27.239	438.713
**********	******	*****

CONSTANT OF JARRETT EQUATION (12)= 51.500

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	2.724	246.326
10.000	5.448	341.020
20.000	10.895	444.972
30.000	16.343	513.097
40.000	21.791	566.199
50.000	27.239	610.640
*******	******	*****

CONSTANT OF JARRETT EQUATION (13)= 61.400

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	2.724	293.678
10.000	5.448	406.576
20.000	10.895	530.511
30.000	16.343	611.732
40.000	21.791	675.041
50.000	27.239	728.026
*****	*****	*****

	CONSTANT	OF	JARRETT	EQUATION	(14) =	72.000
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W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	2.724	344.378
10.000	5.448	476.766
20.000	10.895	622.097
30.000	16.343	717.340
40.000	21.791	791.579
50.000	27.239	853.711

CONSTANT O	F JARRETT E	QUATION (15)=	99.000
W.OF H.C.	W.OF TNT	DISTANCE	

	1010	1015	IVIE I ENS	
	5.000	2.724	473.520	
	10.000	5.448	655.554	
	20.000	10.895	855.384	
	30.000	16.343	986.342	
	40.000	21.791	1088.421	
	50.000	27.239	1173.853	
*	*****	******	*****	*

EXPLOSION YIELD FACTOR(3)= 10.000%

TNT EQUIVALENT VALUE (3)= 0.9178

CONSTANT OF JARRETT EQUATION (1)= 3.000

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	5.448	19.865
10.000	10.895	25.921
20.000	21.791	32.982
30.000	32.686	37.824
40.000	43.582	41.655
50.000	54.477	44.882
******	*****	*****

CONSTANT OF JARRETT EQUATION (2)= 9.000

W.OF H.C. W.OF TNT DISTANCE TONS TONS METERS 5.000 5.448 59.596 10.000 10.895 77.762 20.000 21.791 98.947 30.000 32.686 113.471

40.000	43.582	124.965
50.000	54.477	134.647
*****	******	****

CONSTANT OF JARRETT EQUATION (3)= 10.050

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	5.448	66.549
10.000	10.895	86.834
20.000	21.791	110.491
30.000	32.686	126.710
40.000	43.582	139.544
50.000	54.477	150.356
******	*****	*****

CONSTANT OF	JARRETT	EQUATION	(4)=	13.900
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W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	5.448	92.042
10.000	10.895	120.099
20.000	21.791	152.819
30.000	32.686	175.250
40.000	43.582	193.002
50.000	54.477	207.955
******	*******	*****

CONSTANT OF JARRETT EQUATION (5)= 16.300

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	5.448	107.935
10.000	10.895	140.836
20.000	21.791	179.205
30.000	32.686	205.509
40.000	43.582	226.326
50.000	54.477	243.861
**********	********	*****

CONSTANT OF JARRETT EQUATION (6)= 18.500

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	5.448	122.502
10.000	10.895	159.844
20.000	21.791	203.392
30.000	32.686	233.247
40.000	43.582	256.873
50.000	54.477	276.775
********	******	*****

CONSTANT OF JARRETT EQUATION (7)= 19.800

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	5.448	131.111
10.000	10.895	171.077
20.000	21.791	217.684
30.000	32.686	249.637
40.000	43.582	274.923
50.000	54.477	296.224
*****	******	*****

CONSTANT OF JARRETT EQUATION (8)= 21.600

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	5.448	143.030
10.000	10.895	186.629
20.000	21.791	237.474
30.000	32.686	272.331
40.000	43.582	299.916
50.000	54.477	323.154
*****	*****	******

CONSTANT O	F JARRETT E	QUATION (9)=	22.800
W.OF H.C.	W.OF TNT	DISTANCE	
TONS	TONS	METEDO	

TONS TONS METERS

5.000	5.448	150.976
10.000	10.895	196.998
20.000	21.791	250.667
30.000	32.686	287.461
40.000	43.582	316.578
50.000	54.477	341.107
*****	*******	*****

CONSTANT OF JARRETT EQUATION (10)= 25.500

**

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	5.448	168.855
10.000	10.895	220.326
20.000	21.791	280.351
30.000	32.686	321.502
40.000	43.582	354.068
50.000	54.477	381.501
******	******	*****

CONSTANT OF JARRETT EQUATION (11)= 37.000

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	5.448	245.005
10.000	10.895	319.689
20.000	21.791	406.784
30.000	32.686	466.493
40.000	43.582	513.745
50.000	54.477	553.550
*****	*****	*****

CONSTANT OF JARRETT EQUATION (12)= 51.500

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE METERS
5.000	5.448	341.020
10.000	10.895	444.972
20.000	21.791	566.199
30.000	32.686	649.308
40.000	43.582	715.078
50.000	54.477	770.482

CONSTANT C)F	JARRETT	EQUATION	(13)=	61.400
------------	----	---------	----------	-------	--------

W.OF H.C.	W.OF TNT	DISTANCE
		METERS
5.000	5.448	406.576
10.000	10.895	530.511
20.000	21.791	675.041
30.000	32.686	774.127
40.000	43.582	852.539
50.000	54.477	918.594
******	*******	*****

CONSTANT	OF	JARRETT	EQUATION	(14) =	72.000
				· · · /	

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	5.448	476.766
10.000	10.895	622.097
20.000	21.791	791.579
30.000	32.686	907.771
40.000	43.582	999.720
50.000	54.477	1077.178
********	*****	******

CONSTANT OF	JARRETT	EQUATION	(15)) =
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99.000

W.OF H.C. TONS	W.OF TN TONS	T DISTANCE METERS
5.000	5.448	655.554
10.000	10.895	855.384
20.000	21.791	1088.421
30.000	32.686	1248.185
40.000	43.582	1374.615
50.000	54.477	1481.120
******	*****	*****

PROPANE RESULTS:

EXPLOSION YIELD FACTOR(1)= 2.000%

TNT EQUIVALENT VALUE (1)= 4.9541

CONSTANT OF JARRETT EQUATION (1)= 3.000

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	1.009	7.987
10.000	2.019	12.174
20.000	4.037	17.420
30.000	6.056	20.749
40.000	8.074	23.216
50.000	10.093	25.214

CONSTANT OF JARRETT EQUATION (2)= 9.000

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	1.009	23.961
10.000	2.019	36.521
20.000	4.037	52.259
30.000	6.056	62.246
40.000	8.074	69.648
50.000	10.093	75.643
*****	*****	*****

CONSTANT OF JARRETT EQUATION (3)= 10.050

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	1.009	26.756
10.000	2.019	40.782
20.000	4.037	58.355
30.000	6.056	69.508

40.000	8.074	77.774	
50.000	10.093	84.468	
****	****		

CONSTANT OF JARRETT EQUATION (4)= 13.900

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	1.009	37.006
10.000	2.019	56.405
20.000	4.037	80.711
30.000	6.056	96.136
40.000	8.074	107.567
50.000	10.093	116.826
******	******	*****

	CONSTANT	OF	JARRETT	EQUATION	(5)=	16.300
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W.OF H.C. TONS	W.OF TNT	DISTANCE
5.000	1.009	43.395
10.000	2.019	66.144
20.000	4.037	94.646
30.000	6.056	112.734
40.000	8.074	126.140
50.000	10.093	136.997
*****	*****	*****

CONSTANT OF JARRETT EQUATION (6)= 18.500

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	1.009	49.252
10.000	2.019	75.071
20.000	4.037	107.421
30.000	6.056	127.950
40.000	8.074	143.165
50.000	10.093	155.488
*********	*****	*****

CONSTANT OF JARRETT EQUATION (7)= 19.800

W.OF TNT TONS	DISTANCE
1.009	52.713
2.019	80.346
4.037	114.969
6.056	136.941
8.074	153.225
10.093	166.414
	W.OF TNT TONS 1.009 2.019 4.037 6.056 8.074 10.093

CONSTANT OF JARRETT EQUATION (8)= 21.600

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	1.009	57.505
10.000	2.019	87.650
20.000	4.037	125.421
30.000	6.056	149.390
40.000	8.074	167.155
50.000	10.093	181.542
******	******	*****

CONSTANT OF JARRETT EQUATION (9)= 22.800

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE	
5.000	1.009	60.700	
10.000	2.019	92.520	
20.000	4.037	132.389	
30.000	6.056	157.690	
40.000	8.074	176.441	
50.000	10.093	191.628	
******	******	******	

CONSTANT O	F JARRETT E	QUATION (10)=	25.500
W.OF H.C.	W.OF TNT	DISTANCE	

5.000	1.009	67.888	
10.000	2.019	103.476	
20.000	4.037	148.066	
30.000	6.056	176.364	
40.000	8.074	197.336	
50.000	10.093	214.321	
*******	******	*****	5

CONSTANT OF JARRETT EQUATION (11)= 37.000

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W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	1.009	98.505
10.000	2.019	150.142
20.000	4.037	214.841
30.000	6.056	255.900
40.000	8.074	286.330
50.000	10.093	310.975
******	******	*****

CONSTANT OF JARRETT EQUATION (12)= 51.500

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	1.009	137.108
10.000	2.019	208.981
20.000	4.037	299.036
30.000	6.056	356.186
40.000	8.074	398.541
50.000	10.093	432.844
******	*******	*****

CONSTANT OF JARRETT EQUATION (13)= 61.400

W.OF H.C. TONS		W.OF TNT TONS	DISTANCE
	5.000	1.009	163.465
	10.000	2.019	249.154
	20.000	4.037	356.520
	30.000	6.056	424.656
	40.000	8.074	475.154
	50.000	10.093	516.051

W.OF H.C.	W.OF TNT	DISTANCE
TONS	TONS	METERS
5.000	1.009	191.685
10.000	2.019	292.168
20.000	4.037	418.069
30.000	6.056	497.968
40.000	8.074	557.184
50.000	10.093	605.141

CONSTANT OF LAPPETT FOLIATION (14) 72.000

CONSTANT OF JARRETT EQUATION (15)= 99.000

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	1.009	263.567
10.000	2.019	401.731
20.000	4.037	574.845
30.000	6.056	684.706
40.000	8.074	766.127
50.000	10.093	832.069
******	******	*****

EXPLOSION YIELD FACTOR(2)= 5.000%

TNT EQUIVALENT VALUE (2)= 1.9816

CONSTANT OF JARRETT EQUATION (1)= 3.000

W.OF H.C. TONS		W.OF TNT TONS	DISTANCE
	5.000	2.523	13.777
	10.000	5.046	19.233
	20.000	10.093	25.214
	30.000	15.139	29.108

40.000	20.185	32.134	
50.000	25.232	34.663	
*****	*****	*****	

CONSTANT	OF	JARRETT	EQUATION	(2)=	9.000
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W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	2.523	41.331
10.000	5.046	57.699
20.000	10.093	75.643
30.000	15.139	87.324
40.000	20.185	96.402
50.000	25.232	103.990
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CONSTANT	OF	JARRETT	EQUATION	(3)=	10.050
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W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	2.523	46,153
10.000	5.046	64.430
20.000	10.093	84.468
30.000	15.139	97.511
40.000	20.185	107.649
50.000	25.232	116.122
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CONSTANT OF JARRETT EQUATION (4)= 13.900

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	2.523	63.834
10.000	5.046	89.113
20.000	10.093	116.826
30.000	15.139	134.866
40.000	20.185	148.888
50.000	25.232	160.607
*******	******	******

CONSTANT OF JARRETT EQUATION (5)= 16.300

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	2.523	74.856
10.000	5.046	104.499
20.000	10.093	136.997
30.000	15.139	158.153
40.000	20.185	174.595
50.000	25.232	188.338

CONSTANT OF JARRETT EQUATION (6)= 18.500

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	2 523	84 959
10.000	5.046	118.603
20.000	10.093	155.488
40.000	20.185	198.160
50.000	25.232	213.758

CONSTANT OF JARRETT	EQUATION	(7)=	19.800
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W.OF H.C.	W.OF TNT	DISTANCE
TONS	TONS	METERS
5.000	2.523	90.929
10.000	5.046	126.937
20.000	10.093	166.414
30.000	15.139	192.112
40.000	20.185	212.085
50.000	25.232	228.778
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CONSTANT OF	F JARRETT E	EQUATION (8)=	21.600
W.OF H.C. TONS	W.OF TNT TONS	DISTANCE	

5.000	2.523	99.196
10.000	5.046	138.477
20.000	10.093	181.542
30.000	15.139	209.577
40.000	20.185	231.365
50.000	25.232	249.576
*****	******	*****

CONSTANT OF JARRETT EQUATION (9)= 22.800

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	2.523	104.706
10.000	5.046	146.170
20.000	10.093	191.628
30.000	15.139	221.220
40.000	20.185	244.219
50.000	25.232	263.442
******	******	*****

CONSTANT OF JARRETT EQUATION (10)= 25.500

	W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
	5.000	2.523	117.106
	10.000	5.046	163.480
	20.000	10.093	214.321
	30.000	15.139	247.417
	40.000	20.185	273.140
	50.000	25.232	294.639
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CONSTANT OF JARRETT EQUATION (11)= 37.000

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	2.523	169.918
10.000	5.046	237.206
20.000	10.093	310.975
30.000	15.139	358.997
40.000	20.185	396.320
50.000	25.232	427.515

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CONSTANT OF	JARRETT	EQUATION	(12)=	51.500
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W.OF H.C. TONS	W.OF TNT	DISTANCE
5.000	2.523	236.508
10.000	5.046	330.165
20.000	10.093	432.844
30.000	15.139	499.685
40.000	20.185	551.635
50.000	25.232	595.055
*******	*******	*****

CONSTANT OF JARRETT EQUAT	ION (13)= 61.	.400
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W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	2.523	281.973
10.000	5.046	393.634
20.000	10.093	516.051
30.000	15.139	595.741
40.000	20.185	657.678
50.000	25.232	709.444
*****	******	******

CONSTANT OF JARRETT EQUATION (14)= 72.000

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	2.523	330.652
10.000	5.046	461.590
20.000	10.093	605.141
30.000	15.139	698.589
40.000	20.185	771.218
50.000	25.232	831.921
*****	*****	*****

CONSTANT OF JARRETT EQUATION (15)= 99.000

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	2.523	454.646
10.000	5.046	634.687
20.000	10.093	832.069
30.000	15.139	960.560
40.000	20.185	1060.425
50.000	25.232	1143.892
*****	*****	*****

EXPLOSION YIELD FACTOR(3)= 10.000%

TNT EQUIVALENT VALUE (3)= 0.9908

CONSTANT OF JARRETT EQUATION (1)= 3.000

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	5.046	19.233
10.000	10.093	25.214
20.000	20.185	32.134
30.000	30.278	36.863
40.000	40.371	40.601
50.000	50.464	43.749
******	*****	*****

CONSTANT OF JARRETT EQUATION (2)= 9.000

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	5.046	57.699
10.000	10.093	75.643
20.000	20.185	96.402
30.000	30.278	110.588
40.000	40.371	121.803
50.000	50.464	131.247
*******	********	*****

CONSTANT OF JARRETT EQUATION (3)= 10.050

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	5.046	64.430
10.000	10.093	84.468
20.000	20.185	107.649
30.000	30.278	123.489
40.000	40.371	136.013
50.000	50.464	146.559
*********	*****	*****

CONSTANT OF JARRETT EQUATION (4)= 13.900

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	5.046	89.113
10.000	10.093	116.826
20.000	20.185	148.888
30.000	30.278	170.796
40.000	40.371	188.118
50.000	50.464	202.704
 *****	******	*****

CONSTANT OF JARRETT	EQUATION (5)=	16.300		

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5 000		
10.000	10.093	104.499
20.000	20.185	174.595
30.000	30.278	200.286
40.000	40.371	220.598
50.000	50.464	237.703

CONSTANT	OF	JARRETT	EQUATION	(6)=	18.500

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	5.046	118.603

10.000	10.093	155.488
20.000	20.185	198.160
30.000	30.278	227.319
40.000	40.371	250.372
50.000	50.464	269.785
*****	******	*****

CONSTANT OF JARRETT EQUATION (7)= 19.800

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W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	5.046	126,937
10.000	10.093	166.414
20.000	20.185	212.085
30.000	30.278	243.293
40.000	40.371	267.966
50.000	50.464	288.743
*********	*******	******

CONSTANT OF JARRET	T EQUATION	(8)=	21.600
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W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	5.046	138.477
10.000	10.093	181.542
20.000	20.185	231.365
30.000	30.278	265.410
40.000	40.371	292.327
50.000	50.464	314.993
******	*******	*****

CONSTANT	OF	JARRETT	EQUATION	(9)=	22.800
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W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	5.046	146.170
10.000	10.093	191.628
20.000	20.185	244.219
30.000	30.278	280.155
40.000	40.371	308.567
50.000	50.464	332.492
******	******	*****

CONSTANT OF JARRETT EQUATION (10)= 25.500

W.OF TNT TONS	DISTANCE
5.046	163.480
10.093	214.321
20.185	273.140
30.278	313.331
40.371	345.108
50.464	371.866
	W.OF TNT TONS 5.046 10.093 20.185 30.278 40.371 50.464

CONSTANT OF JARRETT EQUATION (11)= 37.000

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	5.046	237.206
10.000	10.093	310.975
20.000	20.185	396.320
30.000	30.278	454.638
40.000	40.371	500.745
50.000	50.464	539.571
******	*****	*****

CONSTANT OF JARRETT EQUATION (12)= 51.500

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	5.046	330.165
10.000	10.093	432.844
20.000	20.185	551.635
30.000	30.278	632.807
40.000	40.371	696.983
50.000	50.464	751.024
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CONSTANT OF	JARRETT E	QUATION (13)=	61.400
W.OF H.C. TONS	W.OF TNT TONS	DISTANCE	

	5.000	5.046	393.634
	10.000	10.093	516.051
	20.000	20.185	657.678
	30.000	30.278	754.453
	40.000	40.371	830.966
	50.000	50.464	895.396
**	*****	*******	*****

CONSTANT OF JARRETT EQUATION (14)= 72.000

12.00

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE METERS
5.000	5.046	461.590
10.000	10.093	605.141
20.000	20.185	771.218
30.000	30.278	884.700
40.000	40.371	974.422
50.000	50.464	1049.976
*****	*****	******

CONSTANT OF JARRETT EQUATION (15)= 99.000

W.OF H.C. TONS	W.OF TN TONS	T DISTANCE METERS
5.000	5.046	634.687
10.000	10.093	832.069
20.000	20.185	1060.425
30.000	30.278	1216.463
40.000	40.371	1339.831
50.000	50.464	1443.716
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BUTANE RESULTS:

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EXPLOSION YIELD FACTOR(1)= 2.000%

TNT EQUIVALENT VALUE (1)= 5.0237

CONSTANT OF JARRETT EQUATION (1)= 3.000

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	0.995	7.916
10.000	1.991	12.077
20.000	3.981	17.308
30.000	5.972	20.631
40.000	7.962	23.094
50.000	9.953	25.087
******	*****	****

CONSTANT OF JARRETT EQUATION (2)= 9.000

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	0.995	23.749
10.000	1.991	36.231
20.000	3.981	51.923
30.000	5.972	61.894
40.000	7.962	69.281
50.000	9.953	75.260
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CONSTANT OF JA	RRETT EQU	ATION (3)=	10.050
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W.OF H.C. TONS	W.OF TNT TONS	DISTANCE METERS	
5.000	0.995	26.519	
10.000	1.991	40.457	
20.000	3.981	57.980	
30.000	5.972	69.115	
40.000	7.962	77.364	
50.000	9.953	84.040	
******	******	*****	

CONSTANT OF JARRETT EQUATION (4)= 13.900

	W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
	5.000	0.995	36.678
	10.000	1.991	55.956
	20.000	3.981	80.192
	30.000	5.972	95.592
	40.000	7.962	107.000
	50.000	9.953	116.234
k	******	*******	*****

CONSTANT OF JARRETT EQUATION (5)= 16.300

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE	
5.000	0.995	43.011	
10.000	1.991	65.618	
20.000	3.981	94.038	
30.000	5.972	112.097	
40.000	7.962	125.475	
50.000	9.953	136.303	
*******	*****	****	

CONSTANT OF JARRETT EQUATION (6)= 18.500

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	0.995	48.817
10.000	1.991	74.474
20.000	3.981	106.730
30.000	5.972	127.227
40.000	7.962	142.411
50.000	9.953	154.700
******	*****	*****

CONSTANT OF JAR	RETT EQUAT	ION (7)=	19.800
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W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	0.995	52.247
10.000	1.991	79.707
20.000	3.981	114.230

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30.000	5.972	136.167
40.000	7.962	152.418
50.000	9.953	165.571
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CONSTANT OF	JARRETT	EQUATION	(8)=	21.600
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W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	0.995	56.997
10.000	1.991	86.953
20.000	3.981	124.614
30.000	5.972	148.546
40.000	7.962	166.274
50.000	9.953	180.623
******	******	********

CONSTANT	OF	JARRETT	EQUATION	(9)=	22.800
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W.OF H.C. W.OF TNT DISTANCE TONS TONS METERS -----_____ 5.000 0.995 10.000 1.991 60.163 91.784 10.000 20.000 3.981 131.537 30.000 5.972 156.798 40.000 7.962 175.511 9.953 190.657 50.000

CONSTANT OF JARRETT EQUATION (10)= 25.500

W.OF H.C. TONS	W.OF TNT	DISTANCE
5.000	0.995	67.288
10.000	1.991	102.653
20.000	3.981	147.114
30.000	5.972	175.367
40.000	7.962	196.296
50.000	9.953	213.235
*****	******	******

CONSTANT OF JARRETT EQUATION (11)= 37.000

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	0.995	97.633
10.000	1.991	148.948
20.000	3.981	213.460
30.000	5.972	254.454
40.000	7.962	284.821
50.000	9.953	309.400
*********	********	

CONSTANT OF JARRETT EQUATION (12)= 51.500

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	0,995	135.895
10.000	1.991	207.319
20.000	3.981	297.113
30.000	5.972	354.172
40.000	7.962	396.440
50.000	9.953	430.652
*********	*****	******

CONSTANT OF JARRETT EQUATION (13)= 61.400

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	0.995	162.018
10.000	1.991	247.173
20.000	3.981	354.228
30.000	5.972	422.256
40.000	7.962	472.649
50.000	9.953	513.437
********	******	*****

CONSTANT OF	JARRETT E	QUATION (14)=	72.000
W.OF H.C. TONS	W.OF TNT TONS	DISTANCE	

5.000	0.995	189.989
10.000	1.991	289.844
20.000	3.981	415.381
30.000	5.972	495.153
40.000	7.962	554.247
50.000	9.953	602.076
******	******	*****

CONSTANT OF JARRETT EQUATION (15)= 99.000

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	0.995	261.235
10.000	1.991	398.536
20.000	3.981	571.149
30.000	5.972	680.835
40.000	7.962	762.089
50.000	9.953	827.855
*******	******	*****

EXPLOSION YIELD FACTOR(2)= 5.000%

TNT EQUIVALENT VALUE (2)= 2.0095

CONSTANT OF JARRETT EQUATION (1)= 3.000

W.OF H.C.	W.OF TNT	DISTANCE
TONS	TONS	METERS
5.000	2.488	13.674
10.000	4.976	19.118
20.000	9.953	25.087
30.000	14.929	28.967
40.000	19.905	31.981
50.000	24.882	34.500
*******	******	*******

CONSTANT OF JARRETT EQUATION (2)= 9.000

W.OF H.C. W.OF TNT DISTANCE

	TONS	TONS	METERS
-	5.000	2.488	41.022
	10.000	4.976	57.354
	20.000	9.953	75.260
	30.000	14.929	86.901
	40.000	19.905	95.944
	50.000	24.882	103.500
**	*****	*****	*****

CONSTANT OF JARRETT EQUATION (3)= 10.050

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	2.488	45.808
10.000	4.976	64.046
20.000	9.953	84.040
30.000	14.929	97.039
40.000	19.905	107.137
50.000	24.882	115.575

* 2 ************

CONSTANT OF JARRETT EQUATION (4)= 13.900

W.OF H.C. W.OF TNT DISTANCE TONS TONS METERS 5.0002.48863.35610.0004.97688.58120.0009.953116.23430.00014.929134.21440.00019.905148.18050.00024.882159.850 *******

CONSTANT OF JARRETT EQUATION (5)= 16.300

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	2.488	74.295
10.000	4.976	103.875
20.000	9.953	136.303
30.000	14.929	157.387

40.000	19.905	173.765
50.000	24.882	187.450
*****	*****	****

CONSTANT OF JARRETT EQUATION (6)= 18.500

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	2.488	84.322
10.000	4.976	117.895
20.000	9.953	154.700
30.000	14.929	178.630
40.000	19.905	197.218
50.000	24.882	212.750
*****	*****	*****

CONSTANT OF JARRETT EQUATION (7)= 19.800

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5 000	2 499	
10.000	4.976	126.179
20.000	9.953	165.571
30.000	14.929	191.182
40.000	19.905	211.077
50.000	24.882	227.700

CONSTANT OF JARRETT EQUATION (8)= 21.600

W.OF H.C. TONS	W.OF TNT TONS	DISTANCE
5.000	2.488	98.452
10.000	4.976	137.650
20.000	9.953	180.623
30.000	14.929	208.562
40.000	19.905	230.265
50.000	24.882	248.400
*****	******	***********

APPENDIX K

Computer program for calculation of spacing distance using TNT equivalent model (Hopkinson's Law)
```
C
C
C PROGRAM FOR CALCULATION OF SPACING DISTANCE USING
C T.N.T. EQUIVALENT MODEL (HOPKINSON'S LAW)
C ==
                      C
С
C
      DIMENSION W(100), Y(30), TNT(100), PR(100), DS(100), SD(100)
C
C
        READ *,M,N
         READ *, ER
        READ *, (Y(JI), JI=1,3)
        READ *, (W(||), ||=1,M)
        READ *, (PR(JJ), JJ=1,N)
        READ *, (SD(KK), KK=1, N)
С
С
       PRINT 111
 111
       FORMAT(//15X, 'HOPKINSON S LAW RESULTS: ',/,15X,22('*')//)
С
C
        PRINT 124
 124
        FORMAT(//, 10X, 'PROPANE RESULTS: ', /, 10X, 16( '*'))
C
        DO 5 K=1,3
C
       PRINT 127,K,Y(K)
  127 FORMAT(///10X, 'EXPLOSION YIELD FACTOR(', 11, ')=', F10.3,
     @/9X,27('-'))
С
С
С
       DO 2 J=1,M
C
     PRINT 25, J, W(J)
  25 FORMAT(///,7X, 'MASS OF LIQUEFIED PROPANE(',11,')=',F10.2,/,
     *7X,29('-'),//)
C
С
      TNT(J)=W(J)*ER*Y(K)
C
       PRINT 35, J, TNT(J)
       FORMAT(7X, 'TNT EQUIVALENT (', 11, ')=', F10.3, /, 7X, 19('-')//)
  35
C
C
       PRINT 55
 55
       FORMAT(6X, 'RESULTS OF TNT MODEL (HOPKINSON S LAW)', /,
    @6X,36('=')/)
С
C
     PRINT 105
```

```
433
```

	105 F	DRMAT(6X, 'PR.WAVE SCALE DIST.',	
	@'	HAZARD RANGE', /6X, ' (PSI)	(METERS)', /5X, 35('-'))
С			
		DO 3 I=1,N	
		DS(1)=SD(1)*((TNT(J)*1000.0)**0.3333)	
С			
		PRINT 200, PR(1), SD(1), DS(1)	
	200	FORMAT(2X,3(F10.3,2X))	
С			
С			
-	3	CONTINUE	
С			
	2	CONTINUE	
С			
_	5	CONTINUE	
C			
С			
		STOP	
		END	

HOPKINSON S LAW RESULTS:

PROPANE RESULTS: ********

EXPLOSION YIELD FACTOR(1)= 0.020

MASS OF LIQUEFIED PROPANE(1)= 1.00

TNT EQUIVALENT (1)= 0.200

RESULTS OF TNT MODEL (HOPKINSON S LAW)

PR.WAVE	SCALE DIST.	HAZARD RANGE
(PST)		(METERS)
10.000	4.000	23.388
7.250	4.800	28.066
7.000	5.000	29.235
6.080	5.500	32.159
5.000	6.000	35.082
3.470	7.500	43.853
3.000	8.500	49.700
2.500	9.400	54.962
2.300	9.900	57.885
2.000	11.000	64.317
1.500	14.000	81.858
1.000	18.000	105.246
0.690	24.000	140.328
0.400	35.000	204.645
0.290	46.000	268.962
0.140	78.000	456.066
0.100	100.000	584.700

MASS OF LIQUEFIED PROPANE(2)= 5.00

435

TNT EQUIVALENT (2)= 1.000

PR.WAVE (PSI)	SCALE DIST.	(METERS)
10.000	4.000	39.991
7.250	4.800	47.989
7.000	5.000	49.988
6.080	5.500	54.987
5.000	6.000	59.986
3.470	7.500	74.983
3.000	8.500	84.980
2.500	9.400	93.978
2.300	9.900	98.977
2.000	11.000	109.975
1.500	14.000	139.968
1.000	18.000	179.959
0.690	24.000	239.945
0.400	35.000	349.919
0.290	46.000	459.894
0.140	78.000	779.820
0.100	100.000	999.770

RESULTS OF TNT MODEL (HOPKINSON S LAW)

MASS OF LIQUEFIED PROPANE(3)= 10.00

TNT EQUIVALENT (3)= 2.000

PR.WAVE (PSI)	SCALE DIST.	HAZARD RANGE (METERS)
10.000	4.000	50.384
7.250	4.800	60.461
7.000	5.000	62.980
6.080	5.500	69.278
5.000	6.000	75.576
3.470	7.500	94.470
3.000	8.500	107.066
2.500	9.400	118.403
2.300	9.900	124.701
2.000	11.000	138.556

1.500	14.000	176.344
1.000	18.000	226.728
0.690	24.000	302.304
0.400	35.000	440.861
0.290	46.000	579.417
0.140	78.000	982.489
0.100	100.000	1259.602

MASS OF LIQUEFIED PROPANE(4)= 20.00

TNT EQUIVALENT (4)= 4.000

RESULTS OF TNT MODEL (HOPKINSON S LAW)

PR.WAVE (PSI)	SCALE DIST.	HAZARD RANGE (METERS)
10.000	4.000	63.478
7.250	4.800	76.174
7.000	5.000	79.348
6.080	5.500	87.283
5.000	6.000	95.218
3.470	7.500	119.022
3.000	8.500	134.892
2.500	9.400	149.174
2.300	9.900	157.109
2.000	11.000	174.566
1.500	14.000	222.175
1.000	18.000	285.653
0.690	24.000	380.871
0.400	35.000	555.437
0.290	46.000	730.003
0.140	78.000	1237.831
0.100	100.000	1586.962

MASS OF LIQUEFIED PROPANE(5)= 30.00

TNT EQUIVALENT (5)= 6.000

PR.WAVE (PSI)	SCALE DIST.	HAZARD RANGE (METERS)
10.000	4.000	72.664
7.250	4.800	87.197
7.000	5.000	90.830
6.080	5.500	99.913
5.000	6.000	108.996
3.470	7.500	136.245
3.000	8.500	154.410
2.500	9.400	170.760
2.300	9.900	179.843
2.000	11.000	199.825
1.500	14.000	254.323
1.000	18.000	326.987
0.690	24.000	435.983
0.400	35.000	635.808
0.290	46.000	835.633
0.140	78.000	1416.943
0.100	100.000	1816.594

MASS OF LIQUEFIED PROPANE(6)= 40.00

TNT EQUIVALENT (6)= 8.000

PR.WAVE (PSI)	SCALE DIST.	HAZARD RANGE (METERS)
10.000	4.000	79.976
7.250	4.800	95.971
7.000	5.000	99.970
6.080	5.500	109.967
5.000	6.000	119.964
3.470	7.500	149.955
3.000	8.500	169.949
2.500	9.400	187.944
2.300	9.900	197.941
2.000	11.000	219.934
1.500	14.000	279.916
1.000	18.000	359.892
0.690	24.000	479.856
0.400	35.000	699.790
0.290	46.000	919.724
0.140	78.000	1559.533
0.100	100.000	1999.401

MASS OF LIQUEFIED PROPANE(7)= 50.00

TNT EQUIVALENT (7)= 10.000

RESULTS OF TNT MODEL (HOPKINSON S LAW)

PR.WAVE (PSI)	SCALE DIST.	HAZARD RANGE (METERS)
10.000	4.000	86.151
7.250	4.800	103.381
7.000	5.000	107.689
6.080	5.500	118.458
5.000	6.000	129.226
3.470	7.500	161.533
3.000	8.500	183.071
2.500	9.400	202.455
2.300	9.900	213.224
2.000	11.000	236.915
1.500	14.000	301.528
1.000	18.000	387.679
0.690	24.000	516.906
0.400	35.000	753.821
0.290	46.000	990.736
0.140	78.000	1679.943
0.100	100.000	2153.774

EXPLOSION YIELD FACTOR(2)= 0.050

MASS OF LIQUEFIED PROPANE(1)= 1.00

TNT EQUIVALENT (1)= 0.500

(PSI)	SCALE	DIST.	HAZARD RANGE (METERS)	
10.000	4	.000	31.7	741

7.250	4.800	38.090
7.000	5.000	39.677
6.080	5.500	43.644
5.000	6.000	47.612
3.470	7.500	59.515
3.000	8.500	67.451
2.500	9.400	74.592
2.300	9.900	78.560
2.000	11.000	87.289
1.500	14.000	111.095
1.000	18.000	142.837
0.690	24.000	190.449
0.400	35.000	277.738
0.290	46.000	365.027
0.140	78.000	618.958
0.100	100.000	793.536

MASS OF LIQUEFIED PROPANE(2)= 5.00

TNT EQUIVALENT	(2)=	2.500
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RESULTS OF TNT MODEL (HOPKINSON S LAW)

PR.WAVE (PSI)	SCALE DIST.	HAZARD RANGE (METERS)
10.000	4.000	54.274
7.250	4.800	65.129
7.000	5.000	67.843
6.080	5.500	74.627
5.000	6.000	81.411
3.470	7.500	101.764
3.000	8.500	115.333
2.500	9.400	127.544
2.300	9.900	134.329
2.000	11.000	149.254
1.500	14.000	189.960
1.000	18.000	244.234
0.690	24.000	325.645
0.400	35.000	474.899
0.290	46.000	624.153
0.140	78.000	1058.347
0.100	100.000	1356.855

MASS OF LIQUEFIED PROPANE(3)= 10.00

TNT EQUIVALENT (3)= 5.000

RESULTS OF TNT MODEL (HOPKINSON S LAW)

PR.WAVE (PSI)	SCALE DIST.	HAZARD RANGE (METERS)
10.000	4.000	68.380
7.250	4.800	82.056
7.000	5.000	85.475
6.080	5.500	94.022
5.000	6.000	102.569
3.470	7.500	128.212
3.000	8.500	145.307
2.500	9.400	160.692
2.300	9.900	169.240
2.000	11.000	188.044
1.500	14.000	239.329
1.000	18.000	307.708
0.690	24.000	410.278
0.400	35.000	598.322
0.290	46.000	786.366
0.140	78.000	1333.403
0.100	100.000	1709.491

MASS OF LIQUEFIED PROPANE(4)= 20.00

TNT EQUIVALENT (4)= 10.000

PR.WAVE (PSI)	SCALE DIST.	HAZARD RANGE (METERS)
10.000	4.000	86.151
7.250	4.800	103.381
7.000	5.000	107.689
6.080	5.500	118.458
5.000	6.000	129.226
3.470	7.500	161.533
3.000	8.500	183.071

2.500	9.400	202.455
2.300	9.900	213.224
2.000	11.000	236.915
1.500	14.000	301.528
1.000	18.000	387.679
0.690	24.000	516.906
0.400	35.000	753.821
0.290	46.000	990.736
0.140	78.000	1679.943
0.100	100.000	2153.774

MASS OF LIQUEFIED PROPANE(5)= 30.00

TNT EQUIVALENT (5)= 15.000

RESULTS OF TNT MODEL (HOPKINSON S LAW)

PR.WAVE (PSI)	SCALE DIST.	HAZARD RANGE (METERS)
10.000	4.000	98.617
7.250	4.800	118.340
7.000	5.000	123.271
6.080	5.500	135.598
5.000	6.000	147.925
3.470	7.500	184.907
3.000	8.500	209.561
2.500	9.400	231.750
2.300	9.900	244.077
2.000	11.000	271.196
1.500	14.000	345.159
1.000	18.000	443.776
0.690	24.000	591.701
0.400	35.000	862.898
0.290	46.000	1134.094
0.140	78.000	1923.029
0.100	100.000	2465.422

MASS OF LIQUEFIED PROPANE(6)= 40.00

TNT EQUIVALENT (6)= 20.000

PR.WAVE (PSI)	SCALE DIST.	HAZARD RANGE (METERS)
10.000	4.000	108.541
7.250	4.800	130.249
7.000	5.000	135.676
6.080	5.500	149.244
5.000	6.000	162.811
3.470	7.500	203.514
3.000	8.500	230.649
2.500	9.400	255.071
2.300	9.900	268.639
2.000	11.000	298.487
1.500	14.000	379.893
1.000	18.000	488.434
0.690	24.000	651.245
0.400	35.000	949.733
0.290	46.000	1248.220
0.140	78.000	2116.547
0.100	100.000	2713.522

RESULTS OF TNT MODEL (HOPKINSON S LAW)

MASS OF LIQUEFIED PROPANE(7)= 50.00

TNT EQUIVALENT (7)= 25.000

PR.WAVE (PSI)	SCALE DIST.	HAZARD RANGE (METERS)
10.000	4.000	116.921
7.250	4.800	140.305
7.000	5.000	146.152
6.080	5.500	160.767
5.000	6.000	175.382
3.470	7.500	219.227
3.000	8.500	248.458
2.500	9.400	274.765
2.300	9.900	289.380
2.000	11.000	321.533
1.500	14.000	409.224
1.000	18.000	526.146
0.690	24.000	701.527
0.400	35.000	1023.061

0.290	46.000	1344.594
0.140	78.000	2279.964
0.100	100.000	2923.031

EXPLOSION YIELD FACTOR(3)= 0.100

MASS OF LIQUEFIED PROPANE(1)= 1.00 _____

TNT EQUIVALENT (1)= 1.000

RESULTS OF TNT MODEL (HOPKINSON S LAW)

PR.WAVE (PSI)	SCALE DIST.	HAZARD RANGE (METERS)
10.000	4.000	39.991
7.250	4.800	47.989
7.000	5.000	49.988
6.080	5.500	54.987
5.000	6.000	59.986
3.470	7.500	74.983
3.000	8.500	84.980
2.500	9.400	93.978
2.300	9.900	98.977
2.000	11.000	109.975
1.500	14.000	139.968
1.000	18.000	179.959
0.690	24.000	239.945
0.400	35.000	349.919
0.290	46.000	459.894
0.140	78.000	779.820
0.100	100.000	999.770

MASS OF LIQUEFIED PROPANE(2)= 5.00

TNT EQUIVALENT (2)= 5.000

PR.WAVE (PSI)	SCALE DIST.	HAZARD RANGE (METERS)
10.000	4.000	68.380
7.250	4.800	82.056
7.000	5.000	85.475
6.080	5.500	94.022
5.000	6.000	102.569
3.470	7.500	128.212
3.000	8.500	145.307
2.500	9.400	160.692
2.300	9.900	169.240
2.000	11.000	188.044
1.500	14.000	239.329
1.000	18.000	307.708
0.690	24.000	410.278
0.400	35.000	598.322
0.290	46.000	786.366
0.140	78.000	1333.403
0.100	100.000	1709.491

MASS OF LIQUEFIED PROPANE(3)= 10.00

TNT EQUIVALENT (3)= 10.000

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PR.WAVE (PSI)	SCALE DIST.	HAZARD RANGE (METERS)
10.000	4.000	86.151
7.250	4.800	103.381
7.000	5.000	107.689
6.080	5.500	118.458
5.000	6.000	129.226
3.470	7.500	161.533
3.000	8.500	183.071
2.500	9.400	202.455
2.300	9.900	213.224
2.000	11.000	236.915
1.500	14.000	301.528
1.000	18.000	387.679
0.690	24.000	516.906
0.400	35.000	753.821
0.290	46.000	990.736

0.140	78.000	1679.943
0.100	100.000	2153.774

MASS OF LIQUEFIED PROPANE(4)= 20.00

TNT EQUIVALENT (4)= 20.000

RESULTS OF TNT MODEL (HOPKINSON S LAW)

PR.WAVE (PSI)	SCALE DIST.	HAZARD RANGE (METERS)
10.000	4.000	108.541
7.250	4.800	130.249
7.000	5.000	135.676
6.080	5.500	149.244
5.000	6.000	162.811
3.470	7.500	203.514
3.000	8.500	230.649
2.500	9.400	255.071
2.300	9.900	268.639
2.000	11.000	298.487
1.500	14.000	379.893
1.000	18.000	488.434
0.690	24.000	651.245
0.400	35.000	949.733
0.290	46.000	1248.220
0.140	78.000	2116.547
0.100	100.000	2713.522

MASS OF LIQUEFIED PROPANE(5)= 30.00

TNT EQUIVALENT (5)= 30.000

PR.WAVE	SCALE	DIST. HAZARD R		RANGE
(PSI)		(METERS		RS)
10.000	4	.000	124.2	247

7.250	4.800	149.096
7.000	5.000	155.308
6.080	5.500	170.839
5.000	6.000	186.370
3.470	7.500	232.962
3.000	8.500	264.024
2.500	9.400	291.980
2.300	9.900	307.510
2.000	11.000	341.678
1.500	14.000	434.863
1.000	18.000	559.110
0.690	24.000	745.480
0.400	35.000	1087.158
0.290	46.000	1428.836
0.140	78.000	2422.809
0.100	100.000	3106.165

MASS OF LIQUEFIED PROPANE(6)= 40.00

TNT EQUIVALENT (6)= 40.000

RESULTS OF TNT MODEL (HOPKINSON S LAW)

PR.WAVE (PSI)	SCALE DIST.	HAZARD RANGE (METERS)
10.000	4.000	136.750
7.250	4.800	164.100
7.000	5.000	170.937
6.080	5.500	188.031
5.000	6.000	205.125
3.470	7.500	256.406
3.000	8.500	290.593
2.500	9.400	321.362
2.300	9.900	338.456
2.000	11.000	376.062
1.500	14.000	478.624
1.000	18.000	615.374
0.690	24.000	820.499
0.400	35.000	1196.561
0.290	46.000	1572.622
0.140	78.000	2666.621
0.100	100.000	3418.745

MASS OF LIQUEFIED PROPANE(7)= 50.00

TNT EQUIVALENT (7)= 50.000

RESULTS OF TNT MODEL (HOPKINSON S LAW)

PR.WAVE (PSI)	SCALE DIST.	IST. HAZARD RANGE (METERS)				
10.000	4.000	147.308				
7.250	4.800	176.770				
7.000	5.000	184.135				
6.080	5.500	202.549				
5.000	6.000	220.962				
3.470	7.500	276.203				
3.000	8.500	313.030				
2.500	9.400	346.174				
2.300	9.900	364.588				
2.000	11.000	405.097				
1.500	14.000	515.578				
1.000	18.000	662.887				
0.690	24.000	883.849				
0.400	35.000	1288.946				
0.290	46.000	1694.043				
0.140	78.000	2872.508				
0.100	100.000	3682.703				

APPENDIX L

Computer program for calculation of spacing distance using Gugan's Model

```
С
C
C
C PROGRAM FOR CALCULATION OF SPACING DISTANCE USING GUGAN MODEL.
C ==
С
С
С
      DIMENSION W(100), PS(100), EO(100), PC(100), D(100), R(100),
     @N(100),E(100),RS(100),T(100),WTM(100),AN(100),EOR(100),V(100)
C
C
       READ *.A
С
       READ *, (W(11), 11=1,7)
       READ *, (PS(JJ), JJ=1, 18)
          READ *, (N(KK), E(KK), AN(KK), WTM(KK), KK=1, 4)
С
С
      PRINT 333
 333
      FORMAT(//15X, 'GUGAN S MODEL RESULTS: ',/15X,21('*')//)
С
С
      DO 3 |=1,7
С
      PRINT 100, I, W(1)
 100
     FORMAT(10X, 'MASS OF FLAMMABLE GAS IN THE CLOUD(', 12, ')=', F9.1,
     @/10X,40('-')//)
С
      DO 5 K=1,18
C
      PRINT 110,K,PS(K)
 110 FORMAT(10X, 'OVER PRESSURE(', 12, ')=', F10.3, /10X, 15('-')//)
C
С
C
      PRINT 150
 150 FORMAT(3X, ' COMPONENT
                                  V
                                           D
                                                      TO
                                                               EO
                                                                     ۰,
     Q'
          PC
                            R ',/,5X,71('-'))
                   RS
C
C
C
      DO 8 J=1,4
С
      V(J) = (22.4*AN(J)*W(I))/WTM(J)
      D(J)=((12.0*V(J))/3.14159)**0.3333
      T(J)=D(J)/A
      EO(J)=E(J)*W(I)
      PC(J)=(EO(J)*A*1000000.0)/(0.57*(D(J)**4.0))
      G=4.95E+07
      RS(J)=G/PC(J)
      R(J)=(37.86*(EO(J)**0.5)/PS(K))**0.6667
      EOR(J) = (EO(J) **0.333)/R(J)
```

С

PRINT 200,N(J),V(J),D(J),T(J),EO(J),PC(J),RS(J),R(J)
FORMAT(7X, 'C', 11, 5X, 7(1PE8.2, 1X))
CONTINUE
PRINT 220
FORMAT(10X,///)
CONTINUE
PRINT 230
FORMAT(10X ///)
CONTINUE
0700
SIOP
END

GUGAN S MODEL RESULTS:

MASS OF FLAMMABLE GAS IN THE CLOUD(1)= 1000.0

OVER PRESSURE(1)= 69.000

COMPONENT	V	D	то	EO	PC	RS	R
C1	1.47E+04	3.83E+01	1.16E-01	5.01E+04	1.35E+07	3.67E+00	2 47E+01
C2	1.32E+04	3.69E+01	1.12E-01	4.76E+04	1.48E+07	3.34E+00	2 435+01
C3	1.26E+04	3.64E+01	1.10E-01	4.64E+04	1.53E+07	3.23E+00	2.43E+01
C4	1.23E+04	3.61E+01	1.09E-01	4.58E+04	1.56E+07	3.17E+00	2.40E+01

OVER PRESSURE(2)= 50.000

COMPONENT	v	D	то	EO	PC	RS	R
C1	1.47E+04	3.83E+01	1.16E-01	5.01E+04	1.35F+07	3 67E+00	3 065.01
C2	1.32E+04	3.69E+01	1.12E-01	4.76E+04	1.48E+07	3.34E+00	3.01E+01
C3	1.26E+04	3.64E+01	1.10E-01	4.64E+04	1.53E+07	3.23E+00	2.99F+01
C4	1.23E+04	3.61E+01	1.09E-01	4.58E+04	1.56E+07	3.17E+00	2.97E+01

OVER PRESSURE(3)= 48.300

COMPONENT	v	D	то	EO	PC	RS	R
C1	1.47E+04	3.83E+01	1.16E-01	5.01E+04	1.35E+07	3 67E+00	3 145.01
C2	1.32E+04	3.69E+01	1.12E-01	4.76E+04	1.48E+07	3.34E+00	3.08E+01
C3	1.26E+04	3.64E+01	1.10E-01	4.64E+04	1.53E+07	3.23E+00	3.06E+01
C4	1.23E+04	3.61E+01	1.09E-01	4.58E+04	1.56E+07	3.17E+00	3.04E+01

OVER PRESSURE(4)= 42.000

COMPONENT	v	D	то	EO	PC	RS	R
C1	1.47E+04	3.83E+01	1.16E-01	5.01E+04	1.35E+07	3.67E+00	3 44F+01
C2	1.32E+04	3.69E+01	1.12E-01	4.76E+04	1.48E+07	3.34E+00	3.38E+01
C3	1.26E+04	3.64E+01	1.10E-01	4.64E+04	1.53E+07	3.23E+00	3.35E+01
C4	1.23E+04	3.61E+01	1.09E-01	4.58E+04	1.56E+07	3.17E+00	3.34E+01

OVER PRESSURE(5)= 34.500

COMPONENT	V	D	то	EO	PC	RS	R
C1	1.47E+04	3.83E+01	1.16E-01	5.01E+04	1.35E+07	3.67E+00	3 92F+01
C2	1.32E+04	3.69E+01	1.12E-01	4.76E+04	1.48E+07	3.34E+00	3.86F+01
C3	1.26E+04	3.64E+01	1.10E-01	4.64E+04	1.53E+07	3.23E+00	3.83E+01
C4	1.23E+04	3.61E+01	1.09E-01	4.58E+04	1.56E+07	3.17E+00	3.81E+01

OVER PRESSURE(6)= 24.000

COMPONENT	V	D	то	EO	PC	RS	R
C1	1.47E+04	3.83E+01	1.16E-01	5.01F+04	1 355+07	3 675.00	E 005.01
C2	1.32E+04	3.69E+01	1.12E-01	4.76E+04	1.48E+07	3.34E+00	4.91F+01
C3	1.26E+04	3.64E+01	1.10E-01	4.64E+04	1.53E+07	3.23E+00	4.87E+01
C4	1.23E+04	3.61E+01	1.09E-01	4.58E+04	1.56E+07	3.17E+00	4.85E+01

OVER PRESSURE(7)= 20.700

COMPONENT	v	·D	то	EO	PC	RS	R
C1	1.47E+04	3.83E+01	1.16E-01	5.01E+04	1.35E+07	3 67E+00	5 525.01
C2	1.32E+04	3.69E+01	1.12E-01	4.76E+04	1.48E+07	3.34E+00	5.42E+01
C3	1.26E+04	3.64E+01	1.10E-01	4.64E+04	1.53E+07	3.23E+00	5.38E+01
C4	1.23E+04	3.61E+01	1.09E-01	4.58E+04	1.56E+07	3.17E+00	5.35E+01

OVER PRESSURE(8)= 17.300

COMPONENT	V	D	то	EO	PC	RS	R
C1	1.47E+04	3.83E+01	1.16E-01	5.01E+04	1.35E+07	3.67E+00	6.22F+01
C2	1.32E+04	3.69E+01	1.12E-01	4.76E+04	1.48E+07	3.34E+00	6.11E+01
C3	1.26E+04	3.64E+01	1.10E-01	4.64E+04	1.53E+07	3.23E+00	6.06E+01
C4	1.23E+04	3.61E+01	1.09E-01	4.58E+04	1.56E+07	3.17E+00	6 03E+01

OVER PRESSURE(9)= 15.900

COMPONENT	V	D	то	EO	PC	RS	R
C1	1.47E+04	3.83E+01	1.16E-01	5.01E+04	1.35E+07	3.67E+00	6 58E+01
C2	1.32E+04	3.69E+01	1.12E-01	4.76E+04	1.48E+07	3.34E+00	6.46F+01
C3	1.26E+04	3.64E+01	1.10E-01	4.64E+04	1.53E+07	3.23E+00	6.41E+01
C4	1.23E+04	3.61E+01	1.09E-01	4.58E+04	1.56E+07	3.17E+00	6.38E+01

OVER PRESSURE(10)= 13.800

COMPONENT	v	D	то	EO	PC	RS	R
C1	1.47F+04	3 83E+01	1 165 01	5 015.04	1 055.07		
~	1.472404	5.05E+01	1.10E-01	5.01E+04	1.35E+07	3.67E+00	7.23E+01
62	1.32E+04	3.69E+01	1.12E-01	4.76E+04	1.48E+07	3.34E+00	7.10E+01
C3	1.26E+04	3.64E+01	1.10E-01	4.64E+04	1.53E+07	3.23E+00	7 05E+01
C4	1.23E+04	3.61E+01	1.09E-01	4.58E+04	1.56E+07	3.17E+00	7.01E+01

OVER PRESSURE(11)= 10.350

COMPONENT	v	D	то	EO	PC	RS	R
C1	1.47E+04	3.83E+01	1.16E-01	5.01E+04	1 35F+07	3 675+00	9 765.01
C2	1.32E+04	3.69E+01	1.12E-01	4.76E+04	1.48E+07	3.34E+00	8.61F+01
C3	1.26E+04	3.64E+01	1.10E-01	4.64E+04	1.53E+07	3.23E+00	8.54E+01
C4	1.23E+04	3.61E+01	1.09E-01	4.58E+04	1.56E+07	3.17E+00	8.50E+01

OVER PRESSURE(12)= 6.900

COMPONENT	V	D	то	EO	PC	RS	R
C1	1.47E+04	3.83E+01	1.16E-01	5.01E+04	1.35E+07	3.67E+00	1 15E+02
C2	1.32E+04	3.69E+01	1.12E-01	4.76E+04	1.48E+07	3.34E+00	1.13E+02
C3	1.26E+04	3.64E+01	1.10E-01	4.64E+04	1.53E+07	3.23E+00	1.12E+02
C4	1.23E+04	3.61E+01	1.09E-01	4.58E+04	1.56E+07	3.17E+00	1 11E+02

OVER PRESSURE(13)= 4.800

COMPONENT	V	D	то	EO	PC	RS	R
C1	1.47E+04	3.83E+01	1.16E-01	5.01E+04	1.35E+07	3.67E+00	1 46F+02
C2	1.32E+04	3.69E+01	1.12E-01	4.76E+04	1.48E+07	3.34E+00	1 44F+02
C3	1.26E+04	3.64E+01	1.10E-01	4.64E+04	1.53E+07	3.23E+00	1.42E+02
C4	1.23E+04	3.61E+01	1.09E-01	4.58E+04	1.56E+07	3.17E+00	1.42E+02

OVER PRESSURE(14)= 2.760

COMPONENT	v	D	то	EO	PC	RS	R
C1	1.47E+04	3.83E+01	1.16E-01	5.01E+04	1 355+07	2 675.00	0.115.00
C2	1.32E+04	3.69E+01	1.12E-01	4.76E+04	1.48E+07	3.34E+00	2.11E+02 2.08E+02
C3	1.26E+04	3.64E+01	1.10E-01	4.64E+04	1.53E+07	3.23E+00	2.06E+02
C4	1.23E+04	3.61E+01	1.09E-01	4.58E+04	1.56E+07	3.17E+00	2.05E+02

OVER PRESSURE(15)= 2.000

COMPONENT	v	D	то	EO	PC	RS	R
C1	1.47E+04	3.83E+01	1.16E-01	5.01E+04	1 35F+07	3.675.00	2 625.00
C2	1.32E+04	3.69E+01	1.12E-01	4.76E+04	1.48E+07	3.34E+00	2.57E+02
C3	1.26E+04	3.64E+01	1:10E-01	4.64E+04	1.53E+07	3.23E+00	2.55E+02
C4	1.23E+04	3.61E+01	1.09E-01	4.58E+04	1.56E+07	3.17E+00	2.54E+02

OVER PRESSURE(16)= 1.000

455

COMPONENT	v	D	то	EO	PC	RS	R
C1	1.47E+04	3.83E+01	1.16E-01	5.01E+04	1.35E+07	3.67E+00	4 16E+02
C2	1.32E+04	3.69E+01	1.12E-01	4.76E+04	1.48E+07	3.34E+00	4.09E+02
C3	1.26E+04	3.64E+01	1.10E-01	4.64E+04	1.53E+07	3.23E+00	4.05E+02
C4	1.23E+04	3.61E+01	1.09E-01	4.58E+04	1.56E+07	3.17E+00	4.04E+02

OVER PRESSURE(17)= 0.700

COMPONENT	V	D	то	EO	PC	RS	R
C1	1.47E+04	3.83E+01	1.16E-01	5.01E+04	1.35E+07	3.67E+00	5 285+02
C2	1.32E+04	3.69E+01	1.12E-01	4.76E+04	1.48E+07	3.34E+00	5 18E+02
C3	1.26E+04	3.64E+01	1.10E-01	4.64E+04	1.53E+07	3.23E+00	5 14F+02
C4	1.23E+04	3.61E+01	1.09E-01	4.58E+04	1.56E+07	3.17E+00	5.12E+02

OVER PRESSURE(18)= 0.200

COMPONENT	v	D	то	EO	PC	RS	R
C1	1.47E+04	3.83E+01	1.16E-01	5.01E+04	1 35F+07	3 675.00	1 225.02
C2	1.32E+04	3.69E+01	1.12E-01	4.76E+04	1.48E+07	3.34E+00	1.20E+03
C3	1.26E+04	3.64E+01	1.10E-01	4.64E+04	1.53E+07	3.23E+00	1.19E+03
C4	1.23E+04	3.61E+01	1.09E-01	4.58E+04	1.56E+07	3.17E+00	1 18F+03

MASS OF FLAMMABLE GAS IN THE CLOUD(2)= 5000.0

OVER PRESSURE(1)= 69.000

COMPONENT	v	D	то	EO	PC	RS	R
C1	7.36E+04	6.55E+01	1.98E-01	2.51E+05	7.89E+06	6.27E+00	4 235+01
C2	6.59E+04	6.31E+01	1.91E-01	2.38E+05	8.68E+06	5.70E+00	4 15E+01

C3 6.31E+04 6.22E+01 1.89E-01 2.32E+05 8.97E+06 5.52E+00 4.12E+01 C4 6.17E+04 6.17E+01 1.87E-01 2.29E+05 9.12E+06 5.43E+00 4.10E+01

OVER PRESSURE(2)= 50.000

COMPONENT	v	D	то	EO	PC	RS	R
C1	7.36E+04	6.55E+01	1.98E-01	2.51E+05	7.89E+06	6.27E+00	5 24E+01
C2	6.59E+04	6.31E+01	1.91E-01	2.38E+05	8.68E+06	5.70E+00	5 15E+01
C3	6.31E+04	6.22E+01	1.89E-01	2.32E+05	8.97E+06	5.52E+00	5.11E+01
C4	6.17E+04	6.17E+01	1.87E-01	2.29E+05	9.12E+06	5.43E+00	5.08E+01

OVER PRESSURE(3)= 48.300

COMPONENT	v	D	то	EO	PC	RS	R
C1	7.36E+04	6.55E+01	1.98E-01	2.51E+05	7.89E+06	6 27E+00	5 365.01
C2	6.59E+04	6.31E+01	1.91E-01	2.38E+05	8.68E+06	5 70E+00	5.30E+01
C3	6.31E+04	6.22E+01	1.89E-01	2.32E+05	8.97E+06	5.52E+00	5 235+01
C4	6.17E+04	6.17E+01	1.87E-01	2.29E+05	9.12E+06	5.43E+00	5.20E+01

OVER PRESSURE(4)= 42.000

COMPONENT	v	D	то	EO	PC	RS	R
ដ ដ ដ ដ ដ	7.36E+04 6.59E+04 6.31E+04 6.17E+04	6.55E+01 6.31E+01 6.22E+01 6.17E+01	1.98E-01 1.91E-01 1.89E-01 1.87E-01	2.51E+05 2.38E+05 2.32E+05 2.29E+05	7.89E+06 8.68E+06 8.97E+06 9.12E+06	6.27E+00 5.70E+00 5.52E+00 5.43E+00	5.89E+01 5.78E+01 5.74E+01 5.71E+01
OVE	R PRESSURE	E(5)=	34.500				

COMPONENT	v	D	то	EO	PC	RS	R

C1	7.36E+04	6.55E+01	1.98E-01	2.51E+05	7.89E+06	6.27E+00	6.71E+01
C2	6.59E+04	6.31E+01	1.91E-01	2.38E+05	8.68E+06	5.70E+00	6.59E+01
C3	6.31E+04	6.22E+01	1.89E-01	2.32E+05	8.97E+06	5.52E+00	6.54E+01
C4	6.17E+04	6.17E+01	1.87E-01	2.29E+05	9.12E+06	5.43E+00	6.51E+01

OVER PRESSURE(6)= 24.000

COMPONENT	v	D	то	EO	PC	RS	R
C1	7.36E+04	6.55E+01	1.98E-01	2.51E+05	7.89E+06	6 27E+00	8 555+01
C2	6.59E+04	6.31E+01	1.91E-01	2.38E+05	8.68E+06	5.70E+00	8 40E+01
C3	6.31E+04	6.22E+01	1.89E-01	2.32E+05	8.97E+06	5.52E+00	8 33F+01
C4	6.17E+04	6.17E+01	1.87E-01	2.29E+05	9.12E+06	5.43E+00	8.29E+01

OVER PRESSURE(7)= 20.700

		·					
COMPONENT	v	D	то	EO	PC	RS	R
C1	7.36E+04	6.55E+01	1.98E-01	2.51E+05	7.89E+06	6.27E+00	9.43E+01
C2	6.59E+04	6.31E+01	1.91E-01	2.38E+05	8.68E+06	5.70E+00	9 27E+01
C3	6.31E+04	6.22E+01	1.89E-01	2.32E+05	8.97E+06	5.52E+00	9 205+01
C4	6.17E+04	6.17E+01	1.87E-01	2.29E+05	9.12E+06	5.43E+00	9.15E+01

OVER PRESSURE(8)= 17.300

COMPONENT	V	D	то	EO	PC	RS	R
C1	7.36E+04	6.55E+01	1.98E-01	2.51E+05	7.89E+06	6.27E+00	1.06F+02
C2	6.59E+04	6.31E+01	1.91E-01	2.38E+05	8.68E+06	5.70E+00	1.04E+02
C3	6.31E+04	6.22E+01	1.89E-01	2.32E+05	8.97E+06	5.52E+00	1.04E+02
C4	6.17E+04	6.17E+01	1.87E-01	2.29E+05	9.12E+06	5.43E+00	1.03E+02

OVER PRESSURE(9)= 15.900

COMPONENT	v	D	то	EO	PC	RS	R
C1	7.36E+04	6.55E+01	1.98E-01	2.51E+05	7.89E+06	6.27E+00	1,12E+02
C2	6.59E+04	6.31E+01	1.91E-01	2.38E+05	8.68E+06	5.70E+00	1.11E+02
C3	6.31E+04	6.22E+01	1.89E-01	2.32E+05	8.97E+06	5.52E+00	1.10E+02
C4	6.17E+04	6.17E+01	1.87E-01	2.29E+05	9.12E+06	5.43E+00	1.09E+02

OVER PRESSURE(10)= 13.800

COMPONENT	v	D	то	EO	PC	RS	R
C1	7.36E+04	6.55E+01	1.98E-01	2.51E+05	7.89E+06	6.27E+00	1.24F+02
C2	6.59E+04	6.31E+01	1.91E-01	2.38E+05	8.68E+06	5.70E+00	1.21E+02
C3	6.31E+04	6.22E+01	1.89E-01	2.32E+05	8.97E+06	5.52E+00	1.20E+02
C4	6.17E+04	6.17E+01	1.87E-01	2.29E+05	9.12E+06	5.43E+00	1.20E+02

OVER PRESSURE(11)= 10.350

COMPONENT	V	D	TO.	EO	PC	RS	R
C1	7.36E+04	6.55E+01	1.98E-01	2.51E+05	7.89E+06	6.27E+00	1 50E+02
C2	6.59E+04	6.31E+01	1.91E-01	2.38E+05	8.68E+06	5.70E+00	1 475+02
C3	6.31E+04	6.22E+01	1.89E-01	2.32E+05	8.97E+06	5.52E+00	1 465+02
C4	6.17E+04	6.17E+01	1.87E-01	2.29E+05	9.12E+06	5.43E+00	1.45E+02

OVER PRESSURE(12)= 6.900

COMPONENT	v	D	то	EO	PC	RS	R
C1	7.36E+04	6.55E+01	1.98E-01	2.51E+05	7.89E+06	6.27E+00	1 965+02
C2	6.59E+04	6.31E+01	1.91E-01	2.38E+05	8.68E+06	5.70E+00	1.93E+02
C3	6.31E+04	6.22E+01	1.89E-01	2.32E+05	8.97E+06	5.52E+00	1.91E+02
C4	6.17E+04	6.17E+01	1.87E-01	2.29E+05	9.12E+06	5.43E+00	1.90E+02

OVER PRESSURE(13)= 4.800

COMPONENT	V	D	то	EO	PC	RS	R
C1	7.36E+04	6.55E+01	1.98E-01	2.51E+05	7.89E+06	6.27E+00	2 50E+02
C2	6.59E+04	6.31E+01	1.91E-01	2.38E+05	8.68E+06	5.70E+00	2.46F+02
C3	6.31E+04	6.22E+01	1.89E-01	2.32E+05	8.97E+06	5.52E+00	2.44E+02
C4	6.17E+04	6.17E+01	1.87E-01	2.29E+05	9.12E+06	5.43E+00	2.42E+02

OVER PRESSURE(14)= 2.760

COMPONENT	V	D	то	EO	PC	RS	R
C1	7.36E+04	6.55E+01	1.98E-01	2.51E+05	7.89E+06	6.27E+00	3 61E+02
C2	6.59E+04	6.31E+01	1.91E-01	2.38E+05	8.68E+06	5.70E+00	3 555+02
C3	6.31E+04	6.22E+01	1.89E-01	2.32E+05	8.97E+06	5.52E+00	3 52E+02
C4	6.17E+04	6.17E+01	1.87E-01	2.29E+05	9.12E+06	5.43E+00	3.51E+02

OVER PRESSURE(15)= 2.000

COMPONENT	v	D	то	EO	PC	RS	R
C1	7.36E+04	6.55E+01	1.98E-01	2.51E+05	7 895+06	6 27E.00	4 495.00
C2	6.59E+04	6.31E+01	1.91E-01	2.38E+05	8.68E+06	5.70E+00	4.48E+02
C3	6.31E+04	6.22E+01	1.89E-01	2.32E+05	8.97E+06	5.52E+00	4 37E+02
C4	6.17E+04	6.17E+01	1.87E-01	2.29E+05	9.12E+06	5.43E+00	4.35E+02

OVER PRESSURE(16)= 1.000

COMPONENT	v	D	то	EO	PC	RS	R
C1	7.36E+04	6.55E+01	1.98E-01	2.51E+05	7.89E+06	6.27E+00	7 115+02
C2	6.59E+04	6.31E+01	1.91E-01	2.38E+05	8.68E+06	5.70E+00	6 99F+02
C3	6.31E+04	6.22E+01	1.89E-01	2.32E+05	8.97E+06	5.52E+00	6.93E+02
C4	6.17E+04	6.17E+01	1.87E-01	2.29E+05	9.12E+06	5.43E+00	6.90E+02

OVER PRESSURE(17)= 0.700

COMPONENT	v	D	то	EO	PC	RS	R
C1	7.36E+04	6.55E+01	1.98E-01	2.51E+05	7.89E+06	6.27E+00	9.02E+02
C2	6.59E+04	6.31E+01	1.91E-01	2.38E+05	8.68E+06	5.70E+00	8.87E+02
C3	6.31E+04	6.22E+01	1.89E-01	2.32E+05	8.97E+06	5.52E+00	8.79E+02
C4	6.17E+04	6.17E+01	1.87E-01	2.29E+05	9.12E+06	5.43E+00	8.75E+02

OVER PRESSURE(18)= 0.200

COMPONENT	v	D	то	EO	PC	RS	R
C1	7.36E+04	6.55E+01	1.98E-01	2.51E+05	7.89E+06	6.27E+00	2 08F+03
C2	6.59E+04	6.31E+01	1.91E-01	2.38E+05	8.68E+06	5.70E+00	2.04F+03
C3	6.31E+04	6.22E+01	1.89E-01	2.32E+05	8.97E+06	5.52E+00	2.03E+03
C4	6.17E+04	6.17E+01	1.87E-01	2.29E+05	9.12E+06	5.43E+00	2.02E+03

MASS OF FLAMMABLE GAS IN THE CLOUD(3)= 10000.0

OVER PRESSURE(1)= 69.000

COMPONENT	v	D	то	EO	PC	RS	R
C1	1.47E+05	8.25E+01	2.50E-01	5.01E+05	6.26F+06	7 90F+00	5 335.01
C2	1.32E+05	7.95E+01	2.41E-01	4.76E+05	6.89E+06	7.19E+00	5 23E+01
C3	1.26E+05	7.84E+01	2.38E-01	4.64E+05	7.12E+06	6.95E+00	5 19F+01
C4	1.23E+05	7.78E+01	2.36E-01	4.58E+05	7.24E+06	6.83E+00	5.17E+01

OVER PRESSURE(2)= 50.000

COMPONENT	v	D	то	EO	PC	RS	R
C1	1.47E+05	8.25E+01	2.50E-01	5.01E+05	6.26E+06	7.90F+00	6 60E+01
C2	1.32E+05	7.95E+01	2.41E-01	4.76E+05	6.89E+06	7.19E+00	6.49F+01

C3 1.26E+05 7.84E+01 2.38E-01 4.64E+05 7.12E+06 6.95E+00 6.44E+01 C4 1.23E+05 7.78E+01 2.36E-01 4.58E+05 7.24E+06 6.83E+00 6.41E+01

OVER PRESSURE(3)= 48.300

COMPONENT	v	D	то	EO	PC	RS	R
C1	1.47E+05	8.25E+01	2.50E-01	5.01E+05	6.26F+06	7.90E+00	6 76E+01
C2	1.32E+05	7.95E+01	2.41E-01	4.76E+05	6.89E+06	7.19E+00	6.64E+01
C3	1.26E+05	7.84E+01	2.38E-01	4.64E+05	7.12E+06	6.95E+00	6.59E+01
C4	1.23E+05	7.78E+01	2.36E-01	4.58E+05	7.24E+06	6.83E+00	6.55E+01

OVER PRESSURE(4)= 42.000

COMPONENT	V	D	то	EO	PC	RS	R
C1	1.47E+05	8.25E+01	2.50E-01	5.01E+05	6.26E+06	7.90F+00	7 425+01
C2	1.32E+05	7.95E+01	2.41E-01	4.76E+05	6.89E+06	7.195+00	7 205.01
C3	1.26E+05	7.84E+01	2.38E-01	4.64E+05	7.12E+06	6.95E+00	7 235+01
C4	1.23E+05	7.78E+01	2.36E-01	4.58E+05	7.24E+06	6.83E+00	7.19E+01

OVER PRESSURE(5)= 34.500

COMPONENT	v	D	то	EO	PC	RS	R
C1	1.47E+05	8.25E+01	2.50E-01	5.01E+05	6.26E+06	7.90E+00	8.45E+01
C2	1.32E+05	7.95E+01	2.41E-01	4.76E+05	6.89E+06	7.19E+00	8.31E+01
C3	1.26E+05	7.84E+01	2.38E-01	4.64E+05	7.12E+06	6.95E+00	8.24E+01
C4	1.23E+05	7.78E+01	2.36E-01	4.58E+05	7.24E+06	6.83E+00	8.20E+01

OVER PRESSURE(6)= 24.000

COMPONENT	V	D	то	EO	PC	RS	R	

C1	1.47E+05 8.25E+01	2.50E-01	5.01E+05	6.26E+06	7.90E+00	1.08E+02
C2	1.32E+05 7.95E+01	2.41E-01	4.76E+05	6.89E+06	7.19E+00	1.06E+02
C3	1.26E+05 7.84E+01	2.38E-01	4.64E+05	7.12E+06	6.95E+00	1.05E+02
C4	1.23E+05 7.78E+01	2.36E-01	4.58E+05	7.24E+06	6.83E+00	1.04E+02

OVER PRESSURE(7)= 20.700

COMPONENT	V	D	то	EO	PC	RS	R
C1	1.47E+05	8.25E+01	2.50E-01	5.01E+05	6.26E+06	7.90F+00	1 195+02
C2	1.32E+05	7.95E+01	2.41E-01	4.76E+05	6.89E+06	7.19E+00	1 175+02
C3	1.26E+05	7.84E+01	2.38E-01	4.64E+05	7.12E+06	6.95E+00	1 16E+02
C4	1.23E+05	7.78E+01	2.36E-01	4.58E+05	7.24E+06	6.83E+00	1.15E+02

OVER PRESSURE(8)= 17.300

COMPONENT	v	D	то	EO	PC	RS	R
C1	1.47E+05	8.25E+01	2.50E-01	5.01E+05	6.26E+06	7.90E+00	1.34E+02
C2	1.32E+05	7.95E+01	2.41E-01	4.76E+05	6.89E+06	7.19E+00	1.32E+02
C3	1.26E+05	7.84E+01	2.38E-01	4.64E+05	7.12E+06	6.95E+00	1.31E+02
C4	1.23E+05	7.78E+01	2.36E-01	4.58E+05	7.24E+06	6.83E+00	1.30E+02

OVER PRESSURE(9)= 15.900

COMPONENT	V	D	то	EO	PC	RS	R
C1	1.47E+05	8.25E+01	2.50E-01	5.01E+05	6.26E+06	7.90E+00	1.42F±02
C2	1.32E+05	7.95E+01	2.41E-01	4.76E+05	6.89E+06	7.19E+00	1.39F+02
C3	1.26E+05	7.84E+01	2.38E-01	4.64E+05	7.12E+06	6.95E+00	1.38E+02
C4	1.23E+05	7.78E+01	2.36E-01	4.58E+05	7.24E+06	6.83E+00	1.37E+02

OVER PRESSURE(10)= 13.800

COMPONENT	V	D	то	EO	PC	RS	R
C1	1.47E+05	8.25E+01	2.50E-01	5.01E+05	6.26E+06	7.90E+00	1.56E+02
C2	1.32E+05	7.95E+01	2.41E-01	4.76E+05	6.89E+06	7.19E+00	1.53E+02
C3	1.26E+05	7.84E+01	2.38E-01	4.64E+05	7.12E+06	6.95E+00	1.52E+02
C4	1.23E+05	7.78E+01	2.36E-01	4.58E+05	7.24E+06	6.83E+00	1.51E+02

OVER PRESSURE(11)= 10.350

COMPONENT	V	D	то	EO	PC	RS	R
C1	1.47E+05	8.25E+01	2.50E-01	5.01E+05	6.26E+06	7.90F+00	1 89F+02
C2	1.32E+05	7.95E+01	2.41E-01	4.76E+05	6.89E+06	7.19E+00	1.85E+02
C3	1.26E+05	7.84E+01	2.38E-01	4.64E+05	7.12E+06	6.95E+00	1.84F+02
C4	1.23E+05	7.78E+01	2.36E-01	4.58E+05	7.24E+06	6.83E+00	1.83E+02

OVER PRESSURE(12)= 6.900

COMPONENT	v	D	то	EO	PC	RS	R
C1	1.47E+05	8.25E+01	2.50E-01	5.01E+05	6.26E+06	7.90E+00	2.47F+02
C2	1.32E+05	7.95E+01	2.41E-01	4.76E+05	6.89E+06	7.19E+00	2.43E+02
C3	1.26E+05	7.84E+01	2.38E-01	4.64E+05	7.12E+06	6.95E+00	2.41E+02
C4	1.23E+05	7.78E+01	2.36E-01	4.58E+05	7.24E+06	6.83E+00	2.40E+02

OVER PRESSURE(13)= 4.800

COMPONENT	v	D	то	EO	PC	RS	R
C1	1.47E+05	8.25E+01	2.50E-01	5.01E+05	6.26F+06	7 90E+00	3 155.02
C2	1.32E+05	7.95E+01	2.41E-01	4.76E+05	6.89E+06	7.19E+00	3.09E+02
C3	1.26E+05	7.84E+01	2.38E-01	4.64E+05	7.12E+06	6.95E+00	3.07E+02
C4	1.23E+05	7.78E+01	2.36E-01	4.58E+05	7.24E+06	6.83E+00	3.06E+02

OVER PRESSURE(14)= 2.760

COMPONENT	V	D	то	EO	PC	RS	R
C1	1.47E+05	8.25E+01	2.50E-01	5.01E+05	6 26F+06	7 90E+00	4 555.02
C2	1.32E+05	7.95E+01	2.41E-01	4.76E+05	6.89E+06	7.19E+00	4.48E+02
C3	1.26E+05	7.84E+01	2.38E-01	4.64E+05	7.12E+06	6.95E+00	4.44E+02
C4	1.23E+05	7.78E+01	2.36E-01	4.58E+05	7.24E+06	6.83E+00	4.42F+02

OVER PRESSURE(15)= 2.000

COMPONENT	V	D	то	EO	PC	RS	R
C1	1.47E+05	8.25E+01	2.50E-01	5.01E+05	6.26E+06	7.90E+00	5.64F+02
C2	1.32E+05	7.95E+01	2.41E-01	4.76E+05	6.89E+06	7.19E+00	5.55E+02
C3	1.26E+05	7.84E+01	2.38E-01	4.64E+05	7.12E+06	6.95E+00	5.50E+02
C4	1.23E+05	7.78E+01	2.36E-01	4.58E+05	7.24E+06	6.83E+00	5.48E+02

OVER PRESSURE(16)= 1.000

COMPONENT	v	D	то	EO	PC	RS	R
C1	1.47E+05	8.25E+01	2.50E-01	5.01E+05	6.26E+06	7 905+00	8 965.02
C2	1.32E+05	7.95E+01	2.41E-01	4.76E+05	6.89E+06	7.19E+00	8.81F+02
C3	1.26E+05	7.84E+01	2.38E-01	4.64E+05	7.12E+06	6.95E+00	8.73E+02
C4	1.23E+05	7.78E+01	2.36E-01	4.58E+05	7.24E+06	6.83E+00	8.69F+02

OVER PRESSURE(17)= 0.700

COMPONENT	V	D	то	EO	PC	RS	R
C1	1.47E+05	8.25E+01	2.50E-01	5.01E+05	6.26F+06	7 90E+00	1 145.02
C2	1.32E+05	7.95E+01	2.41E-01	4.76E+05	6.89E+06	7.19E+00	1 125+03
C3	1.26E+05	7.84E+01	2.38E-01	4.64E+05	7.12E+06	6.95E+00	1.11E+03
C4	1.23E+05	7.78E+01	2.36E-01	4.58E+05	7.24E+06	6.83E+00	1.10E+03

OVER PRESSURE(18)= 0.200

COMPONENT	v	D	то	EO	PC	RS	R
C1	1.47E+05	8.25E+01	2.50E-01	5.01E+05	6.26E+06	7.90E+00	2.62F+03
C2	1.32E+05	7.95E+01	2.41E-01	4.76E+05	6.89E+06	7.19E+00	2.58E+03
C3	1.26E+05	7.84E+01	2.38E-01	4.64E+05	7.12E+06	6.95E+00	2.55E+03
C4	1.23E+05	7.78E+01	2.36E-01	4.58E+05	7.24E+06	6.83E+00	2.54E+03

MASS OF FLAMMABLE GAS IN THE CLOUD(4)= 20000.0

OVER PRESSURE(1)= 69.000

COMPONENT	v	D	то	EO	PC	RS	R
C1	2.95E+05	1.04E+02	3.15E-01	1.00E+06	4 97F+06	9 965+00	6 715.01
C2	2.64E+05	1.00E+02	3.04E-01	9.52E+05	5.47E+06	9.05E+00	6.59E+01
C3	2.53E+05	9.88E+01	2.99E-01	9.29E+05	5.65E+06	8.75E+00	6.54E+01
C4	2.47E+05	9.80E+01	2.97E-01	9.16E+05	5.75E+06	8.61E+00	6.51E+01

OVER PRESSURE(2)= 50.000

COMPONENT	V	D	то	EO	PC	RS	R
C1	2.95E+05	1.04E+02	3.15E-01	1.00E+06	4.97E+06	9.96F+00	8 32E+01
C2	2.64E+05	1.00E+02	3.04E-01	9.52E+05	5.47E+06	9.05E+00	8 17E+01
C3	2.53E+05	9.88E+01	2.99E-01	9.29E+05	5.65E+06	8.75E+00	8 11E+01
C4	2.47E+05	9.80E+01	2.97E-01	9.16E+05	5.75E+06	8.61E+00	8.07E+01

OVER PRESSURE(3)= 48.300

COMPONENT	v	D	то	EO	PC	RS	R
C1	2.95E+05	1.04E+02	3.15E-01	1.00E+06	4.97E+06	9.96F+00	8 515:01
C2	2.64E+05	1.00E+02	3.04E-01	9.52E+05	5.47E+06	9.05E+00	8.36E+01

C3 2.53E+05 9.88E+01 2.99E-01 9.29E+05 5.65E+06 8.75E+00 8.30E+01 C4 2.47E+05 9.80E+01 2.97E-01 9.16E+05 5.75E+06 8.61E+00 8.26E+01

OVER PRESSURE(4)= 42.000

COMPONENT	v	D	то	EO	PC	RS	R
C1	2.95E+05	1.04E+02	3.15E-01	1.00E+06	4.97E+06	9,96E+00	9.34F+01
C2	2.64E+05	1.00E+02	3.04E-01	9.52E+05	5.47E+06	9.05E+00	9 18E+01
C3	2.53E+05	9.88E+01	2.99E-01	9.29E+05	5.65E+06	8.75E+00	9.11E+01
C4	2.47E+05	9.80E+01	2.97E-01	9.16E+05	5.75E+06	8.61E+00	9.07E+01

OVER PRESSURE(5)= 34.500

COMPONENT	v	D	то	EO	PC	RS	R
C1	2.95E+05	1.04E+02	3.15E-01	1.00E+06	4.97E+06	9 96F+00	1.075.02
C2	2.64E+05	1.00E+02	3.04E-01	9.52E+05	5.47E+06	9.05E+00	1.072+02
C3	2.53E+05	9.88E+01	2.99E-01	9.29E+05	5.65E+06	8.75E+00	1.04F+02
C4	2.47E+05	9.80E+01	2.97E-01	9.16E+05	5.75E+06	8.61E+00	1.03E+02

OVER PRESSURE(6)= 24.000

COMPONENT	v	D	то	EO	PC	RS	R
C1	2.95E+05	1.04E+02	3.15E-01	1.00E+06	4.97E+06	9.96E+00	1.36E+02
C2	2.64E+05	1.00E+02	3.04E-01	9.52E+05	5.47E+06	9.05E+00	1.33E+02
C3	2.53E+05	9.88E+01	2.99E-01	9.29E+05	5.65E+06	8.75E+00	1.32E+02
C4	2.47E+05	9.80E+01	2.97E-01	9.16E+05	5.75E+06	8.61E+00	1.32E+02
OVEF	PRESSURE	:(7)=	20.700				
COMPONENT	v	D	то	FO	PC	DS	

C1	2.95E+05	1.04E+02	3.15E-01	1.00E+06	4.97E+06	9.96E+00	1.50E+02
C2	2.64E+05	1.00E+02	3.04E-01	9.52E+05	5.47E+06	9.05E+00	1.47E+02
C3	2.53E+05	9.88E+01	2.99E-01	9.29E+05	5.65E+06	8.75E+00	1.46E+02
C4	2.47E+05	9.80E+01	2.97E-01	9.16E+05	5.75E+06	8.61E+00	1.45E+02

OVER PRESSURE(8)= 17.300

COMPONENT	v	D	то	EO	PC	RS	R
C1	2.95E+05	1.04E+02	3.15E-01	1.00E+06	4.97E+06	9.96F+00	1 695+02
C2	2.64E+05	1.00E+02	3.04E-01	9.52E+05	5.47E+06	9.05E+00	1.66F+02
C3	2.53E+05	9.88E+01	2.99E-01	9.29E+05	5.65E+06	8.75E+00	1.65E+02
C4	2.47E+05	9.80E+01	2.97E-01	9.16E+05	5.75E+06	8.61E+00	1.64E+02

OVER PRESSURE(9)= 15.900

COMPONENT	V	D	то	EO	PC	RS	R
C1	2.95E+05	1.04E+02	3.15E-01	1.00E+06	4.97E+06	9.96F+00	1 795+02
C2	2.64E+05	1.00E+02	3.04E-01	9.52E+05	5.47E+06	9 05E+00	1.755.02
C3	2.53E+05	9.88E+01	2.99E-01	9.29E+05	5.65E+06	8.75E+00	1 745+02
C4	2.47E+05	9.80E+01	2.97E-01	9.16E+05	5.75E+06	8.61E+00	1.73E+02

OVER PRESSURE(10)= 13.800

 COMPONENT
 V
 D
 TO
 EO
 PC
 RS
 R

 C1
 2.95E+05
 1.04E+02
 3.15E-01
 1.00E+06
 4.97E+06
 9.96E+00
 1.96E+02

 C2
 2.64E+05
 1.00E+02
 3.04E-01
 9.52E+05
 5.47E+06
 9.05E+00
 1.93E+02

 C3
 2.53E+05
 9.88E+01
 2.99E-01
 9.29E+05
 5.65E+06
 8.75E+00
 1.91E+02

 C4
 2.47E+05
 9.80E+01
 2.97E-01
 9.16E+05
 5.75E+06
 8.61E+00
 1.90E+02

OVER PRESSURE(11)= 10.350

.
COMPONENT	v	D	то	EO	PC	RS	R
C1	2.95E+05	1.04E+02	3.15E-01	1.00E+06	4.97E+06	9.96E+00	2.38E+02
C2	2.64E+05	1.00E+02	3.04E-01	9.52E+05	5.47E+06	9.05E+00	2.34E+02
C3	2.53E+05	9.88E+01	2.99E-01	9.29E+05	5.65E+06	8.75E+00	2.32E+02
C4	2.47E+05	9.80E+01	2.97E-01	9.16E+05	5.75E+06	8.61E+00	2.31E+02

OVER PRESSURE(12)= 6.900

COMPONENT	V	D	то	EO	PC	RS	R
C1	2.95E+05	1.04E+02	3.15E-01	1.00E+06	4.97E+06	9.96E+00	3.11E+02
C2	2.64E+05	1.00E+02	3.04E-01	9.52E+05	5.47E+06	9.05E+00	3.06E+02
C3	2.53E+05	9.88E+01	2.99E-01	9.29E+05	5.65E+06	8.75E+00	3 04F+02
C4	2.47E+05	9.80E+01	2.97E-01	9.16E+05	5.75E+06	8.61E+00	3.02E+02

OVER PRESSURE(13)= 4.800

COMPONENT	v	D	то	EO	PC	RS	R
C1 C2	2.95E+05 2.64E+05	1.04E+02 1.00E+02	3.15E-01 3.04E-01	1.00E+06	4.97E+06	9.96E+00	3.97E+02
C3 C4	2.53E+05 2.47E+05	9.88E+01 9.80E+01	2.99E-01 2.97E-01	9.29E+05 9.16E+05	5.65E+06 5.75E+06	8.75E+00 8.61E+00	3.87E+02 3.85E+02

OVER PRESSURE(14)= 2.760

.

COMPONENT	V	D	то	EO	PC	RS	R
C1	2.95E+05	1.04E+02	3.15E-01	1.00E+06	4.97E+06	9.96F+00	5 74E+02
C2	2.64E+05	1.00E+02	3.04E-01	9.52E+05	5.47E+06	9.05E+00	5.64E+02
C3	2.53E+05	9.88E+01	2.99E-01	9.29E+05	5.65E+06	8.75E+00	5.59E+02
C4	2.47E+05	9.80E+01	2.97E-01	9.16E+05	5.75E+06	8.61E+00	5.57E+02

OVER PRESSURE(15)= 2.000

COMPONENT	V	D	то	EO	PC	RS	R
C1	2.95E+05	1.04E+02	3.15E-01	1.00E+06	4.97E+06	9.96E+00	7.11F+02
C2	2.64E+05	1.00E+02	3.04E-01	9.52E+05	5.47E+06	9.05E+00	6.99E+02
C3	2.53E+05	9.88E+01	2.99E-01	9.29E+05	5.65E+06	8.75E+00	6.93E+02
C4	2.47E+05	9.80E+01	2.97E-01	9.16E+05	5.75E+06	8.61E+00	6.90E+02

OVER PRESSURE(16)= 1.000

COMPONENT	V	D	то	EO	PC	RS	R
C1	2.95E+05	1.04E+02	3.15E-01	1.00E+06	4.97E+06	9.96E+00	1 13E+03
C2	2.64E+05	1.00E+02	3.04E-01	9.52E+05	5.47E+06	9.05E+00	1 11E+03
C3	2.53E+05	9.88E+01	2.99E-01	9.29E+05	5.65E+06	8.75E+00	1 105+03
C4	2.47E+05	9.80E+01	2.97E-01	9.16E+05	5.75E+06	8.61E+00	1.10E+03

OVER PRESSURE(17)= 0.700

COMPONENT	V	D	то	EO	PC	RS	R
C1	2.95E+05	1.04E+02	3.15E-01	1.00E+06	4.97E+06	9 965+00	1 425.02
C2	2.64E+05	1.00E+02	3.04E-01	9.52E+05	5.47E+06	9.05E+00	1.41E+03
C3	2.53E+05	9.88E+01	2.99E-01	9.29E+05	5.65E+06	8.75E+00	1.40E+03
C4	2.47E+05	9.80E+01	2.97E-01	9.16E+05	5.75E+06	8.61E+00	1.39E+03

OVER PRESSURE(18)= 0.200

COMPONENT	v	D	то	EO	PC	RS	R
C1	2.95E+05	1.04E+02	3.15E-01	1.00E+06	4.97F+06	9 965+00	3 305.03
C2	2.64E+05	1.00E+02	3.04E-01	9.52E+05	5.47E+06	9.05E+00	3.24F+03
C3	2.53E+05	9.88E+01	2.99E-01	9.29E+05	5.65E+06	8.75E+00	3.22E+03
C4	2.47E+05	9.80E+01	2.97E-01	9.16E+05	5.75E+06	8.61E+00	3.20E+03

MASS OF FLAMMABLE GAS IN THE CLOUD(5)= 30000.0

OVER PRESSURE(1)= 69.000

COMPONENT	v	D	то	EO	PC	RS	R
C1	4.42E+05	1.19E+02	3.61E-01	1.50E+06	4.34E+06	1.14E+01	7 68E+01
C2	3.96E+05	1.15E+02	3.48E-01	1.43E+06	4.78E+06	1.04E+01	7.55E+01
C3	3.79E+05	1.13E+02	3.43E-01	1.39E+06	4.94E+06	1.00E+01	7.49E+01
C4	3.70E+05	1.12E+02	3.40E-01	1.37E+06	5.02E+06	9.86E+00	7.45E+01

OVER PRESSURE(2)= 50.000

COMPONENT	v	D	то	EO	PC	RS	R
C1	4.42E+05	1.19E+02	3.61E-01	1.50F+06	4 34E+06	1 145.01	0.505.01
C2	3.96E+05	1.15E+02	3.48E-01	1.43E+06	4.78E+06	1.04E+01	9.52E+01
C3	3.79E+05	1.13E+02	3.43E-01	1.39E+06	4.94E+06	1.00E+01	9.28F+01
C4	3.70E+05	1.12E+02	3.40E-01	1.37E+06	5.02E+06	9.86E+00	9.24E+01

OVER PRESSURE(3)= 48.300

COMPONENT	V	D	то	EO	PC	RS	R
C1 C2 C3 C4	4.42E+05 3.96E+05 3.79E+05 3.70E+05	1.19E+02 1.15E+02 1.13E+02 1.12E+02	3.61E-01 3.48E-01 3.43E-01 3.40E-01	1.50E+06 1.43E+06 1.39E+06 1.37E+06	4.34E+06 4.78E+06 4.94E+06 5.02E+06	1.14E+01 1.04E+01 1.00E+01 9.86E+00	9.74E+01 9.58E+01 9.50E+01 9.45E+01

OVER PRESSURE(4)= 42.000

COMPONENT	v	D	то	EO	PC	RS	R
C1	4.42E+05	1.19E+02	3.61E-01	1.50E+06	4.34E+06	1.14F+01	1 07E+02
C2	3.96E+05	1.15E+02	3.48E-01	1.43E+06	4.78E+06	1.04E+01	1.05E+02

C3 3.79E+05 1.13E+02 3.43E-01 1.39E+06 4.94E+06 1.00E+01 1.04E+02 C4 3.70E+05 1.12E+02 3.40E-01 1.37E+06 5.02E+06 9.86E+00 1.04E+02

OVER PRESSURE(5)= 34.500

COMPONENT	V	D	то	EO	PC	RS	R
C1	4.42E+05	1.19E+02	3.61E-01	1.50E+06	4.34E+06	1.14E+01	1.22F+02
C2	3.96E+05	1.15E+02	3.48E-01	1.43E+06	4.78E+06	1.04E+01	1 20E+02
C3	3.79E+05	1.13E+02	3.43E-01	1.39E+06	4.94E+06	1.00E+01	1.19E+02
C4	3.70E+05	1.12E+02	3.40E-01	1.37E+06	5.02E+06	9.86E+00	1.18E+02

OVER PRESSURE(6)= 24.000

COMPONENT	V	D	то	EO	PC	RS	R
C1	4.42E+05	1.19E+02	3.61E-01	1.50E+06	4.34E+06	1.14F+01	1 555+02
C2	3.96E+05	1.15E+02	3.48E-01	1.43E+06	4.78E+06	1.04E+01	1.535+02
C3	3.79E+05	1.13E+02	3.43E-01	1.39E+06	4.94E+06	1.00E+01	1 515+02
C4	3.70E+05	1.12E+02	3.40E-01	1.37E+06	5.02E+06	9.86E+00	1.51E+02

OVER PRESSURE(7)= 20.700

COMPONENT	V	D	то	EO	PC	RS	R
C1	4.42E+05	1.19E+02	3.61E-01	1.50E+06	4.34E+06	1.14E+01	1.71E+02
C2	3.96E+05	1.15E+02	3.48E-01	1.43E+06	4.78E+06	1.04E+01	1.68E+02
C3	3.79E+05	1.13E+02	3.43E-01	1.39E+06	4.94E+06	1.00E+01	1.67E+02
C4	3.70E+05	1.12E+02	3.40E-01	1.37E+06	5.02E+06	9.86E+00	1.66E+02
						•	
OVER	R PRESSURE	E(8)=	17.300				
COMPONENT	v	D	то	EO	PC	RS	R

C1	4.42E+05	1.19E+02	3.61E-01	1.50E+06	4.34E+06	1.14E+01	1.93E+02
C2	3.96E+05	1.15E+02	3.48E-01	1.43E+06	4.78E+06	1.04E+01	1.90E+02
C3	3.79E+05	1.13E+02	3.43E-01	1.39E+06	4.94E+06	1.00E+01	1.88E+02
C4	3.70E+05	1.12E+02	3.40E-01	1.37E+06	5.02E+06	9.86E+00	1.87E+02

OVER PRESSURE(9)= 15.900

COMPONENT	v	D	то	EO	PC	RS	R
C1	4.42E+05	1.19E+02	3.61E-01	1.50E+06	4.34E+06	1.14F+01	2 04F+02
C2	3.96E+05	1.15E+02	3.48E-01	1.43E+06	4.78E+06	1.04E+01	2.01E+02
C3	3.79E+05	1.13E+02	3.43E-01	1.39E+06	4.94E+06	1.00E+01	1.99F+02
C4	3.70E+05	1.12E+02	3.40E-01	1.37E+06	5.02E+06	9.86E+00	1.98E+02

OVER PRESSURE(10)= 13.800

COMPONENT	V	D	то	EO	PC	RS	R
C1	4.42E+05	1.19E+02	3.61E-01	1.50E+06	4.34E+06	1.14F+01	2 255.02
C2	3.96E+05	1.15E+02	3.48E-01	1.43E+06	4.78E+06	1.04E+01	2 21E+02
C3	3.79E+05	1.13E+02	3.43E-01	1.39E+06	4.94E+06	1.00E+01	2 195+02
C4	3.70E+05	1.12E+02	3.40E-01	1.37E+06	5.02E+06	9.86E+00	2.18E+02

OVER PRESSURE(11)= 10.350

COMPONENT	V	D	то	EO	PC	RS	R
C1	4.42E+05	1.19E+02	3.61E-01	1.50E+06	4.34E+06	1.14F+01	2 72E+02
C2	3.96E+05	1.15E+02	3.48E-01	1.43E+06	4.78E+06	1.04E+01	2.72L+02
C3	3.79E+05	1.13E+02	3.43E-01	1.39E+06	4.94E+06	1.00E+01	2 655+02
C4	3.70E+05	1.12E+02	3.40E-01	1.37E+06	5.02E+06	9.86E+00	2.64E+02

OVER PRESSURE(12)= 6.900

COMPONENT	v	D	то	EO	PC	RS	R
C1	4.42E+05	1.19E+02	3.61E-01	1.50E+06	4.34E+06	1.14E+01	3.57E+02
C2	3.96E+05	1.15E+02	3.48E-01	1.43E+06	4.78E+06	1.04E+01	3.50E+02
C3	3.79E+05	1.13E+02	3.43E-01	1.39E+06	4.94E+06	1.00E+01	3.48E+02
C4	3.70E+05	1.12E+02	3.40E-01	1.37E+06	5.02E+06	9.86E+00	3.46E+02

OVER PRESSURE(13)= 4.800

COMPONENT	V	D	то	EO	PC	RS	R
C1	4.42E+05	1.19E+02	3.61E-01	1.50E+06	4.34E+06	1.14E+01	4.54F+02
C2	3.96E+05	1.15E+02	3.48E-01	1.43E+06	4.78E+06	1.04E+01	4.46E+02
C3	3.79E+05	1.13E+02	3.43E-01	1.39E+06	4.94E+06	1.00E+01	4.43E+02
C4	3.70E+05	1.12E+02	3.40E-01	1.37E+06	5.02E+06	9.86E+00	4.41E+02

OVER PRESSURE(14)= 2.760

COMPONENT	v	D	то	EO	PC	RS	R
C1	4.42E+05	1.19E+02	3.61E-01	1.50E+06	4.34F+06	1 14F+01	6 57E+02
C2	3.96E+05	1.15E+02	3.48E-01	1.43E+06	4.78E+06	1.04E+01	6 45E+02
C3	3.79E+05	1.13E+02	3.43E-01	1.39E+06	4.94E+06	1.00E+01	6 40F+02
C4	3.70E+05	1.12E+02	3.40E-01	1.37E+06	5.02E+06	9.86E+00	6.37E+02

OVER PRESSURE(15)= 2.000

COMPONENT	v	D	то	EO	PC	RS	R
C1	4.42E+05	1.19E+02	3.61E-01	1.50E+06	4.34E+06	1.14F+01	8 14F+02
C2	3.96E+05	1.15E+02	3.48E-01	1.43E+06	4.78E+06	1.04E+01	8.00E+02
C3	3.79E+05	1.13E+02	3.43E-01	1.39E+06	4.94E+06	1.00E+01	7.94E+02
C4	3.70E+05	1.12E+02	3.40E-01	1.37E+06	5.02E+06	9.86E+00	7.90E+02

OVER PRESSURE(16)= 1.000

COMPONENT	V	D	то	EO	PC	RS	R
C1	4.42E+05	1.19E+02	3.61E-01	1.50E+06	4.34E+06	1.14E+01	1.29F+03
C2	3.96E+05	1.15E+02	3.48E-01	1.43E+06	4.78E+06	1.04E+01	1.27E+03
C3	3.79E+05	1.13E+02	3.43E-01	1.39E+06	4.94E+06	1.00E+01	1.26E+03
C4	3.70E+05	1.12E+02	3.40E-01	1.37E+06	5.02E+06	9.86E+00	1.25E+03

OVER PRESSURE(17)= 0.700

COMPONENT	V	D	то	EO	PC	RS	R
C1	4.42E+05	1.19E+02	3.61E-01	1.50E+06	4.34E+06	1.14F+01	1 645+02
C2	3.96E+05	1.15E+02	3.48E-01	1.43E+06	4.78E+06	1.04E+01	1 615+03
C3	3.79E+05	1.13E+02	3.43E-01	1.39E+06	4.94E+06	1.00E+01	1 60F+03
C4	3.70E+05	1.12E+02	3.40E-01	1.37E+06	5.02E+06	9.86E+00	1.59E+03

OVER PRESSURE(18)= 0.200

COMPONENT	v	D	то	EO	PC	RS	R
C1 C2 C3 C4	4.42E+05 3.96E+05 3.79E+05 3.70E+05	1.19E+02 1.15E+02 1.13E+02 1.13E+02 1.12E+02	3.61E-01 3.48E-01 3.43E-01 3.40E-01	1.50E+06 1.43E+06 1.39E+06 1.37E+06	4.34E+06 4.78E+06 4.94E+06 5.02E+06	1.14E+01 1.04E+01 1.00E+01 9.86E+00	3.78E+03 3.71E+03 3.68E+03 3.67E+03

MASS OF FLAMMABLE GAS IN THE CLOUD(6)= 40000.0

OVER PRESSURE(1)= 69.000

COMPONENT	v	D	то	EO	PC	RS	R
C1	5.89E+05	1.31E+02	3.97E-01	2.01E+06	3.95E+06	1 25E+01	8 455.01
C2	5.27E+05	1.26E+02	3.83E-01	1.90E+06	4.34E+06	1.14F+01	8 31E+01

C3 5.05E+05 1.24E+02 3.77E-01 1.86E+06 4.49E+06 1.10E+01 8.24E+01 C4 4.94E+05 1.23E+02 3.74E-01 1.83E+06 4.56E+06 1.08E+01 8.20E+01

OVER PRESSURE(2)= 50.000

COMPONENT	v	D	то	EO	PC	RS	R
C1	5.89E+05	1.31E+02	3.97E-01	2.01E+06	3.95E+06	1.25E+01	1.05F+02
C2	5.27E+05	1.26E+02	3.83E-01	1.90E+06	4.34E+06	1.14E+01	1.03E+02
C3	5.05E+05	1.24E+02	3.77E-01	1.86E+06	4.49E+06	1.10E+01	1.02E+02
C4	4.94E+05	1.23E+02	3.74E-01	1.83E+06	4.56E+06	1.08E+01	1.02E+02

OVER PRESSURE(3)= 48.300

COMPONENT	V	D	то	EO	PC	RS	R
C1	5.89E+05	1.31E+02	3.97E-01	2.01E+06	3.95E+06	1.25E+01	1.07F+02
C2	5.27E+05	1.26E+02	3.83E-01	1.90E+06	4.34E+06	1.14E+01	1.05E+02
C3	5.05E+05	1.24E+02	3.77E-01	1.86E+06	4.49E+06	1.10E+01	1.05E+02
C4	4.94E+05	1.23E+02	3.74E-01	1.83E+06	4.56E+06	1.08E+01	1.04E+02

OVER PRESSURE(4)= 42.000

COMPONEN	л	v	D	то	EO	PC	RS	R
C1	5.89	E+05	1.31E+02	3.97E-01	2.01E+06	3.95E+06	1.25E+01	1,18E+02
C2	5.27	7E+05	1.26E+02	3.83E-01	1.90E+06	4.34E+06	1.14E+01	1.16E+02
C3	5.05	5E+05	1.24E+02	3.77E-01	1.86E+06	4.49E+06	1.10E+01	1.15E+02
C4	4.94	1E+05	1.23E+02	3.74E-01	1.83E+06	4.56E+06	1.08E+01	1.14E+02
						: *		
0	ER PRE	SSURE	E(5)=	34.500				
COMPONEN	п	v	D	то	EO	PC	RS	R

C1	5.89E+05	1.31E+02	3.97E-01	2.01E+06	3.95E+06	1.25E+01	1.34E+02
C2	5.27E+05	1.26E+02	3.83E-01	1.90E+06	4.34E+06	1.14E+01	1 32E+02
C3	5.05E+05	1.24E+02	3.77E-01	1.86E+06	4.49E+06	1.10E+01	1.31E+02
C4	4.94E+05	1.23E+02	3.74E-01	1.83E+06	4.56E+06	1.08E+01	1.30E+02

OVER PRESSURE(6)= 24.000

COMPONENT	v	D	то	EO	PC	RS	R
C1	5.89E+05	1.31E+02	3.97E-01	2.01E+06	3.95E+06	1.25E+01	1 71E+02
C2	5.27E+05	1.26E+02	3.83E-01	1.90E+06	4.34E+06	1.14E+01	1 68F+02
C3	5.05E+05	1.24E+02	3.77E-01	1.86E+06	4.49E+06	1.10E+01	1.67E+02
C4	4.94E+05	1.23E+02	3.74E-01	1.83E+06	4.56E+06	1.08E+01	1.66E+02

OVER PRESSURE(7)= 20.700

COMPONENT	v	D	то	EO	PC	RS	R
C1	5.89E+05	1.31E+02	3.97E-01	2.01E+06	3.95F+06	1 255+01	1 905.00
C2	5.27E+05	1.26E+02	3.83E-01	1.90E+06	4.34F+06	1 145+01	1.092+02
C3	5.05E+05	1.24E+02	3.77E-01	1.86E+06	4.49E+06	1 105+01	1.802+02
C4	4.94E+05	1.23E+02	3.74E-01	1.83E+06	4.56E+06	1.08E+01	1.83E+02

OVER PRESSURE(8)= 17.300

COMPONENT	V	D	то	EO	PC	RS	R
C1	5.89E+05	1.31E+02	3.97E-01	2.01E+06	3.95E+06	1 25E+01	2 135.02
C2	5.27E+05	1.26E+02	3.83E-01	1.90E+06	4.34E+06	1.14E+01	2.13E+02
C3	5.05E+05	1.24E+02	3.77E-01	1.86E+06	4.49E+06	1.10E+01	2.07E+02
C4	4.94E+05	1.23E+02	3.74E-01	1.83E+06	4.56E+06	1.08E+01	2.06E+02

OVER PRESSURE(9)= 15.900

COMPONENT	V	D	то	EO	PC	RS	R
C1	5.89E+05	1.31E+02	3.97E-01	2.01E+06	3.95E+06	1.25E+01	2 25F+02
C2	5.27E+05	1.26E+02	3.83E-01	1.90E+06	4.34E+06	1.14E+01	2.21E+02
C3	5.05E+05	1.24E+02	3.77E-01	1.86E+06	4.49E+06	1.10E+01	2.19E+02
C4	4.94E+05	1.23E+02	3.74E-01	1.83E+06	4.56E+06	1.08E+01	2.18E+02

OVER PRESSURE(10)= 13.800

COMPONENT	V	D	то	EO	PC	RS	R
C1	5.89E+05	1.31E+02	3.97E-01	2.01E+06	3.95E+06	1.25E+01	2.47F+02
C2	5.27E+05	1.26E+02	3.83E-01	1.90E+06	4.34E+06	1.14E+01	2.43E+02
C3	5.05E+05	1.24E+02	3.77E-01	1.86E+06	4.49E+06	1.10E+01	2 41F+02
C4	4.94E+05	1.23E+02	3.74E-01	1.83E+06	4.56E+06	1.08E+01	2.40E+02

OVER PRESSURE(11)= 10.350

COMPONENT	v	D	то	EO	PC	RS	R
C1	5.89E+05	1.31E+02	3.97E-01	2.01E+06	3.95E+06	1.25E+01	2.99E+02
C2	5.27E+05	1.26E+02	3.83E-01	1.90E+06	4.34E+06	1.14E+01	2.94E+02
C3	5.05E+05	1.24E+02	3.77E-01	1.86E+06	4.49E+06	1.10E+01	2.92E+02
C4	4.94E+05	1.23E+02	3.74E-01	1.83E+06	4.56E+06	1.08E+01	2.91E+02

OVER PRESSURE(12)= 6.900

COMPONENT	v	D	то	EO	PC	RS	R
C1	5.89E+05	1.31E+02	3.97E-01	2.01F+06	3 955+06	1 255.01	2.025.00
C2	5.27E+05	1.26E+02	3.83E-01	1.90E+06	4.34E+06	1.14E+01	3.86F+02
C3	5.05E+05	1.24E+02	3.77E-01	1.86E+06	4.49E+06	1.10E+01	3.83E+02
C4	4.94E+05	1.23E+02	3.74E-01	1.83E+06	4.56E+06	1.08E+01	3.81E+02

OVER PRESSURE(13)= 4.800

COMPONENT	V	D	то	EO	PC	RS	R
C1	5.89E+05	1.31E+02	3.97E-01	2.01E+06	3.95E+06	1.25E+01	5.00F+02
C2	5.27E+05	1.26E+02	3.83E-01	1.90E+06	4.34E+06	1.14E+01	4.91E+02
C3	5.05E+05	1.24E+02	3.77E-01	1.86E+06	4.49E+06	1.10E+01	4.87E+02
C4	4.94E+05	1.23E+02	3.74E-01	1.83E+06	4.56E+06	1.08E+01	4.85E+02

OVER PRESSURE(14)= 2.760

COMPONENT	V	D	то	EO	PC	RS	R
C1	5.89E+05	1.31E+02	3.97E-01	2.01E+06	3.95E+06	1.25E+01	7 23F+02
C2	5.27E+05	1.26E+02	3.83E-01	1.90E+06	4.34E+06	1.14E+01	7 10F+02
C3	5.05E+05	1.24E+02	3.77E-01	1.86E+06	4.49E+06	1.10E+01	7 055+02
C4	4.94E+05	1.23E+02	3.74E-01	1.83E+06	4.56E+06	1.08E+01	7.01E+02

OVER PRESSURE(15)= 2.000

COMPONENT	V	D	то	EO	PC	RS	R
C1	5.89E+05	1.31E+02	3.97E-01	2.01E+06	3 95F+06	1 255+01	9 065.00
C2	5.27E+05	1.26E+02	3.83E-01	1.90E+06	4.34F+06	1 145+01	8 815.02
C3	5.05E+05	1.24E+02	3.77E-01	1.86E+06	4.49E+06	1.10E+01	8 735+02
C4	4.94E+05	1.23E+02	3.74E-01	1.83E+06	4.56E+06	1.08E+01	8.69E+02

OVER PRESSURE(16)= 1.000

COMPONENT	V	D	то	EO	PC	RS	R
C1	5.89E+05	1.31E+02	3.97E-01	2.01E+06	3.95E+06	1.25E+01	1 425+03
C2	5.27E+05	1.26E+02	3.83E-01	1.90E+06	4.34E+06	1.14E+01	1.40F+03
C3	5.05E+05	1.24E+02	3.77E-01	1.86E+06	4.49E+06	1.10E+01	1.39E+03
C4	4.94E+05	1.23E+02	3.74E-01	1.83E+06	4.56E+06	1.08E+01	1.38E+03

OVER PRESSURE(17)= 0.700

COMPONENT	v	D	то	EO	PC	RS	R
C1	5.89E+05	1.31E+02	3.97E-01	2.01E+06	3.95E+06	1,25E+01	1.80F+03
C2	5.27E+05	1.26E+02	3.83E-01	1.90E+06	4.34E+06	1.14E+01	1.77E+03
C3	5.05E+05	1.24E+02	3.77E-01	1.86E+06	4.49E+06	1.10E+01	1.76E+03
C4	4.94E+05	1.23E+02	3.74E-01	1.83E+06	4.56E+06	1.08E+01	1.75E+03

OVER PRESSURE(18)= 0.200

COMPONENT	V	D	то	EO	PC	RS	R
C1	5.89E+05	1.31E+02	3.97E-01	2.01E+06	3.95E+06	1.25E+01	4 165.02
C2	5.27E+05	1.26E+02	3.83E-01	1.90E+06	4.34E+06	1.14F+01	4 09E+03
C3	5.05E+05	1.24E+02	3.77E-01	1.86E+06	4.49E+06	1.10E+01	4.05E+03
C4	4.94E+05	1.23E+02	3.74E-01	1.83E+06	4.56E+06	1.08E+01	4.04E+03

MASS OF FLAMMABLE GAS IN THE CLOUD(7)= 50000.0

OVER PRESSURE(1)= 69.000

COMPONENT	V	D	то	EO	PC	RS	R
C1	7.36E+05	1.41E+02	4.28E-01	2.51E+06	3.66F+06	1 35F+01	9 115.01
C2	6.59E+05	1.36E+02	4.12E-01	2.38E+06	4.03E+06	1.23E+01	9.11E+01
C3	6.31E+05	1.34E+02	4.06E-01	2.32E+06	4.17E+06	1.19E+01	8 88E+01
C4	6.17E+05	1.33E+02	4.03E-01	2.29E+06	4.24E+06	1.17E+01	8.84F+01

OVER PRESSURE(2)= 50.000

COMPONENT	v	D	то	EO	PC	RS	R
C1	7.36E+05	1.41E+02	4.28E-01	2.51E+06	3.66E+06	1 35F+01	1 135.02
C2	6.59E+05	1.36E+02	4.12E-01	2.38E+06	4.03E+06	1.23E+01	1.11E+02

C3 6.31E+05 1.34E+02 4.06E-01 2.32E+06 4.17E+06 1.19E+01 1.10E+02 C4 6.17E+05 1.33E+02 4.03E-01 2.29E+06 4.24E+06 1.17E+01 1.10E+02

OVER PRESSURE(3)= 48.300

COMPONENT	v	D	то	EO	PC	RS	R
C1	7.36E+05	1.41E+02	4.28E-01	2.51E+06	3.66E+06	1.35E+01	1,16F+02
C2	6.59E+05	1.36E+02	4.12E-01	2.38E+06	4.03E+06	1.23E+01	1 14F+02
C3	6.31E+05	1.34E+02	4.06E-01	2.32E+06	4.17E+06	1.19E+01	1.13E+02
C4	6.17E+05	1.33E+02	4.03E-01	2.29E+06	4.24E+06	1.17E+01	1.12E+02

OVER PRESSURE(4)= 42.000

COMPONENT	V	D	то	EO	PC	RS	R
C1	7.36E+05	1.41E+02	4.28E-01	2.51E+06	3.66E+06	1.35E+01	1 275+02
C2	6.59E+05	1.36E+02	4.12E-01	2.38E+06	4.03E+06	1.23E+01	1 255+02
C3	6.31E+05	1.34E+02	4.06E-01	2.32E+06	4.17E+06	1.19E+01	1.24E+02
C4	6.17E+05	1.33E+02	4.03E-01	2.29E+06	4.24E+06	1.17E+01	1.23E+02

OVER PRESSURE(5)= 34.500

COMPONENT	V	D	то	EO	PC	RS	R
C1 .	7.36E+05	1.41E+02	4.28E-01	2.51E+06	3.66E+06	1.35E+01	1 45F+02
C2	6.59E+05	1.36E+02	4.12E-01	2.38E+06	4.03E+06	1.23E+01	1 425+02
C3	6.31E+05	1.34E+02	4.06E-01	2.32E+06	4.17E+06	1.19E+01	1 41E+02
C4	6.17E+05	1.33E+02	4.03E-01	2.29E+06	4.24E+06	1.17E+01	1.40E+02

OVER PRESSURE(6)= 24.000

COMPONENT	V	D	то	EO	PC	RS	R	
-----------	---	---	----	----	----	----	---	--

C1	7.36E+05	1.41E+02	4.28E-01	2.51E+06	3.66E+06	1.35E+01	1.84E+02
C2	6.59E+05	1.36E+02	4.12E-01	2.38E+06	4.03E+06	1.23E+01	1.81E+02
C3	6.31E+05	1.34E+02	4.06E-01	2.32E+06	4.17E+06	1.19E+01	1.80E+02
C4	6.17E+05	1.33E+02	4.03E-01	2.29E+06	4.24E+06	1.17E+01	1.79E+02

OVER PRESSURE(7)= 20.700

COMPONENT	V	D	то	EO	PC	RS	R
C1	7.36E+05	1.41E+02	4.28E-01	2.51E+06	3.66E+06	1.35E+01	2.03E+02
C2	6.59E+05	1.36E+02	4.12E-01	2.38E+06	4.03E+06	1.23E+01	2.00E+02
C3	6.31E+05	1.34E+02	4.06E-01	2.32E+06	4.17E+06	1.19E+01	1.98E+02
C4	6.17E+05	1.33E+02	4.03E-01	2.29E+06	4.24E+06	1.17E+01	1.97E+02

OVER PRESSURE(8)= 17.300

COMPONENT	V	D	то	EO	PC	RS	R
C1	7.36E+05	1.41E+02	4.28E-01	2.51E+06	3.66E+06	1.35E+01	2.29F+02
C2	6.59E+05	1.36E+02	4.12E-01	2.38E+06	4.03E+06	1.23E+01	2.25E+02
C3	6.31E+05	1.34E+02	4.06E-01	2.32E+06	4.17E+06	1.19E+01	2.23E+02
C4	6.17E+05	1.33E+02	4.03E-01	2.29E+06	4.24E+06	1.17E+01	2.22E+02

OVER PRESSURE(9)= 15.900

COMPONENT	V	D	то	EO	PC	RS	R
C1	7.36E+05	1.41E+02	4.28E-01	2.51E+06	3.66E+06	1.35E+01	2.42F+02
C2	6.59E+05	1.36E+02	4.12E-01	2.38E+06	4.03E+06	1.23E+01	2.38E+02
C3	6.31E+05	1.34E+02	4.06E-01	2.32E+06	4.17E+06	1.19E+01	2.36E+02
C4	6.17E+05	1.33E+02	4.03E-01	2.29E+06	4.24E+06	1.17E+01	2.35E+02

OVER PRESSURE(10)= 13.800

COMPONENT	V	D	то	EO	PC	RS	R
C1	7.36E+05	1.41E+02	4.28E-01	2.51E+06	3.66E+06	1.35E+01	2.66F+02
C2	6.59E+05	1.36E+02	4.12E-01	2.38E+06	4.03E+06	1.23E+01	2.62E+02
C3	6.31E+05	1.34E+02	4.06E-01	2.32E+06	4.17E+06	1.19E+01	2.60E+02
C4	6.17E+05	1.33E+02	4.03E-01	2.29E+06	4.24E+06	1.17E+01	2.58E+02

OVER PRESSURE(11)= 10.350

COMPONENT	V	D	то	EO	PC	RS	R
C1	7.36E+05	1.41E+02	4.28E-01	2.51E+06	3.66E+06	1.35E+01	3 23E+02
C2	6.59E+05	1.36E+02	4.12E-01	2.38E+06	4.03E+06	1.23E+01	3.17F+02
C3	6.31E+05	1.34E+02	4.06E-01	2.32E+06	4.17E+06	1.19E+01	3.14E+02
C4	6.17E+05	1.33E+02	4.03E-01	2.29E+06	4.24E+06	1.17E+01	3.13E+02

OVER PRESSURE(12)= 6.900

COMPONENT	V	D	то	EO	PC	RS	R
C1	7.36E+05	1.41E+02	4.28E-01	2.51E+06	3.66E+06	1 355+01	4 225.02
C2	6.59E+05	1.36E+02	4.12E-01	2.38E+06	4.03E+06	1 23E+01	4.155+02
C3	6.31E+05	1.34E+02	4.06E-01	2.32E+06	4.17E+06	1.19E+01	4 125+02
C4	6.17E+05	1.33E+02	4.03E-01	2.29E+06	4.24E+06	1.17E+01	4.10F+02

OVER PRESSURE(13)= 4.800

COMPONENT	v	D	то	EO	PC	RS	R
C1	7.36E+05	1.41E+02	4.28E-01	2.51E+06	3.66F+06	1 35E+01	5 385.02
C2	6.59E+05	1.36E+02	4.12E-01	2.38E+06	4.03E+06	1.23E+01	5.30E+02
C3	6.31E+05	1.34E+02	4.06E-01	2.32E+06	4.17E+06	1.19E+01	5.25E+02
C4	6.17E+05	1.33E+02	4.03E-01	2.29E+06	4.24E+06	1.17E+01	5.22E+02

OVER PRESSURE(14)= 2.760

COMPONENT	V	D	то	EO	PC	RS	R
C1	7.36E+05	1.41E+02	4.28E-01	2.51E+06	3.66E+06	1.35E+01	7.79F+02
C2	6.59E+05	1.36E+02	4.12E-01	2.38E+06	4.03E+06	1.23E+01	7.65E+02
C3	6.31E+05	1.34E+02	4.06E-01	2.32E+06	4.17E+06	1.19E+01	7.59E+02
C4	6.17E+05	1.33E+02	4.03E-01	2.29E+06	4.24E+06	1.17E+01	7.56E+02

OVER PRESSURE(15)= 2.000

COMPONENT	V	D	то	EO	PC	RS	R
C1	7.36E+05	1.41E+02	4.28E-01	2.51E+06	3.66E+06	1.35E+01	9 65F+02
C2	6.59E+05	1.36E+02	4.12E-01	2.38E+06	4.03E+06	1.23E+01	9.49F+02
C3	6.31E+05	1.34E+02	4.06E-01	2.32E+06	4.17E+06	1.19E+01	9.41F+02
C4	6.17E+05	1.33E+02	4.03E-01	2.29E+06	4.24E+06	1.17E+01	9.37E+02

OVER PRESSURE(16)= 1.000

COMPONENT	v	D	то	EO	PC	RS	R
C1	7.36E+05	1.41E+02	4.28E-01	2.51E+06	3.66F+06	1 35F+01	1 535.02
C2	6.59E+05	1.36E+02	4.12E-01	2.38E+06	4.03E+06	1.23E+01	1.53E+03
C3	6.31E+05	1.34E+02	4.06E-01	2.32E+06	4.17E+06	1.19E+01	1.49E+03
C4	6.17E+05	1.33E+02	4.03E-01	2.29E+06	4.24E+06	1.17E+01	1.49E+03

OVER PRESSURE(17)= 0.700

COMPONENT	v	D	то	EO	PC	RS	R
C1	7.36E+05	1.41E+02	4.28E-01	2.51E+06	3.66F+06	1 35F+01	1 945.02
C2	6.59E+05	1.36E+02	4.12E-01	2.38E+06	4.03E+06	1.23E+01	1 91F+03
C3	6.31E+05	1.34E+02	4.06E-01	2.32E+06	4.17E+06	1.19E+01	1.89E+03
C4	6.17E+05	1.33E+02	4.03E-01	2.29E+06	4.24E+06	1.17E+01	1.89E+03

OVER PRESSURE(18)= 0.200

COMPONENT	v	D	то	EO	PC	RS	R
C1	7.36E+05	1.41E+02	4.28E-01	2.51E+06	3.66E+06	1.35E+01	4.48E+03
C2	6.59E+05	1.36E+02	4.12E-01	2.38E+06	4.03E+06	1.23E+01	4.40E+03
C3	6.31E+05	1.34E+02	4.06E-01	2.32E+06	4.17E+06	1.19E+01	4.37E+03
C4	6.17E+05	1.33E+02	4.03E-01	2.29E+06	4.24E+06	1.17E+01	4.35E+03

APPENDIX M

Computer program for calculation of spacing distance using Clancey's model

```
C
С
C
C PROGRAM FOR CALCULATION OF SPACING DISTANCE USING CLANCEY MODEL.
C =
            C
C
С
С
      DIMENSION W(100), F1(100), F2(100), C1(100), C2(100), QM(100),
     @RS(100),R(100),N(100),X(100),R1(100),RE(100),WTM(100)
C
C
       READ *,T
C
      PRINT 222
 222
      FORMAT(//15X, 'CLANCEY S MODEL RESULTS: '/, 15X, 23('*')//)
С
С
       READ *, (W(11), 11=1,7)
       READ *, (RS(NN), NN=1, 18)
C
         READ *, (N(KK), F1(KK), F2(KK), WTM(KK), KK=1,4)
C
С
      DO 3 1=1,7
C
      PRINT 100, I, W(1)
 100 FORMAT(10X, 'MASS OF FLAMMABLE GAS IN THE CLOUD(', 12, ')=', F9.1,
     @/10X,40('-')//)
C
C
      DO 6 L=1,18
С
       PRINT 155, L, RS(L)
     FORMAT(10X, 'SCALED RANGE (', 12, ')=', F10.3, /, 10X, 16('-'))
 155
C
      PRINT 150
 150 FORMAT(3X, ' COMPONENT
                                 C1
                                          C2
                                                    QM
                                                             х
                                                                   ۰,
     @'
          R1
                   RE
                              R',/,4X,72('-'))
C
С
С
      DO 8 J=1,4
С
      C1(J)=(12.187*F1(J)*WTM(J))/T
      C2(J)=(12.187*F2(J)*WTM(J))/T
      QM(J)=0.5*((W(I)/C2(J))**0.333)
      X(J) = (10.0*QM(J))**1.14
      R1(J)=0.71*((W(I)/C2(J))**0.333)*((ALOG(C2(J)/C1(J)))**0.5)
      RE(J)=2.0*R1(J)
       R(J)=(RE(J)*RS(L))/0.48
```

CC

	PRINT 200,N(J),C1(J),C2(J),QM(J),X(J),R1(J),RE(J),R(J)
200	FORMAT(7X, 'C', 11, 5X, 7(F8.2, 1X))
С	
8	CONTINUE
С	
	PRINT 220
220	FORMAT(10X,///)
С	
6	CONTINUE
С	
	PRINT 230
230	FORMAT(10X,///)
С	
С	
3	CONTINUE
С	
	STOP
	END

MASS OF FLAMMABLE GAS IN THE CLOUD(1)= 1000.0

SCALED RANGE (1)= 0.480

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	10.75	206.82	15.99	31.99	31 99
C2	0.04	0.16	9.26	174.59	15.35	30.70	30.70
C3	0.04	0.17	8.93	167.54	14.88	29.76	29.76
C4	0.05	0.20	8.49	158.07	14.70	29.39	29.39

SCALE	D RANGE (2)=	0.660				
COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	10.75	206.82	15.99	31.99	43 98
C2	0.04	0.16	9.26	174.59	15.35	30.70	42.22
C3	0.04	0.17	8.93	167.54	14.88	29.76	40.92
C4	0.05	0.20	8.49	158.07	14.70	29.39	40.41

SCALED RANGE (3)= 0.670

C1	C2	QM	x	R1	RE	R
0.03	0.10	10.75	206.82	15.99	31 99	44.65
0.04	0.16	9.26	174.59	15.35	30.70	42 86
0.04	0.17	8.93	167.54	14.88	29.76	41 54
0.05	0.20	8.49	158.07	14.70	29.39	41.02
	C1 0.03 0.04 0.04 0.05	C1 C2 0.03 0.10 0.04 0.16 0.04 0.17 0.05 0.20	C1 C2 GM 0.03 0.10 10.75 0.04 0.16 9.26 0.04 0.17 8.93 0.05 0.20 8.49	C1 C2 CM X 0.03 0.10 10.75 206.82 0.04 0.16 9.26 174.59 0.04 0.17 8.93 167.54 0.05 0.20 8.49 158.07	C1 C2 QM X R1 0.03 0.10 10.75 206.82 15.99 0.04 0.16 9.26 174.59 15.35 0.04 0.17 8.93 167.54 14.88 0.05 0.20 8.49 158.07 14.70	C1 C2 QM X R1 RE 0.03 0.10 10.75 206.82 15.99 31.99 0.04 0.16 9.26 174.59 15.35 30.70 0.04 0.17 8.93 167.54 14.88 29.76 0.05 0.20 8.49 158.07 14.70 29.39

SCALED RANGE (4)= 0.700

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	10.75	206.82	15.99	31 99	46 65
C2	0.04	0.16	9.26	174.59	15.35	30.70	44 78
C3	0.04	0.17	8.93	167.54	14.88	29.76	43.39
C4	0.05	0.20	8.49	158.07	14.70	29.39	42.86

SCALED RANGE (5)= 0.750 COMPONENT QM C1 C2 х R1 RE R C1 0.030.1010.75206.820.040.169.26174.59 15.9931.9915.3530.70 49.98 C2 47.97 0.04 0.17 8.93 167.54 C3 14.88 29.76 46.49 C4 0.05 0.20 8.49 158.07 29.39 14.70 45.92

SCALED RANGE (6)= 0.900

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	10.75	206.82	15.99	31.99	59 97
C2	0.04	0.16	9.26	174.59	15.35	30.70	57.57
C3	0.04	0.17	8.93	167.54	14.88	29.76	55.79
C4	0.05	0.20	8.49	158.07	14.70	29.39	55.11

SCALED RANGE (7)= 0.980

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	10.75	206.82	15.99	31 99	65 21
C2	0.04	0.16	9.26	174.59	15.35	30.70	62 69
C3	0.04	0.17	8.93	167.54	14.88	29.76	60.75
C4	0.05	0.20	8.49	158.07	14.70	29.39	60.01

SCALED RANGE (8)= 1.090

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	10.75	206.82	15.99	31.99	72 64
C2	0.04	0.16	9.26	174.59	15.35	30.70	69.72
C3	0.04	0.17	8.93	167.54	14.88	29.76	67.57
C4	0.05	0.20	8.49	158.07	14.70	29.39	66.74

SCALED	RANGE	(9)=	1.120

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	10.75	206.82	15.99	31.99	74.63
C2	0.04	0.16	9.26	174.59	15.35	30.70	71.64

C3	0.04	0.17	8.93	167.54	14.88	29.76	69.43
C4	0.05	0.20	8.49	158.07	14.70	29.39	68.58

SCALED RANGE (10)= 1.210

COMPONENT	C1	C2	QM	х	R1	RE	R
C1	0.03	0.10	10.75	206.82	15.99	31,99	80.63
C2	0.04	0.16	9.26	174.59	15.35	30.70	77.40
C3	0.04	0.17	8.93	167.54	14.88	29.76	75.01
C4	0.05	0.20	8.49	158.07	14.70	29.39	74.09

SCALED RANGE (11)= 1.530

COMPONENT	C1	C2	QM	х	R1	RE	R
C1	0.03	0.10	10.75	206.82	15.99	31 99	101 96
C2	0.04	0.16	9.26	174.59	15.35	30.70	97 87
C3	0.04	0.17	8.93	167.54	14.88	29.76	94.85
C4	0.05	0.20	8.49	158.07	14.70	29.39	93.68

SCALED RANGE (12)= 2.100

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	10.75	206.82	15.99	31 99	120 04
C2	0.04	0.16	9.26	174.59	15.35	30.70	134 33
C3	0.04	0.17	8.93	167.54	14.88	29.76	130.18
C4	0.05	0.20	8.49	158.07	14.70	29.39	128.58

SCALED RANGE (13)= 2.800

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	10.75	206.82	15.99	31,99	186 59
C2	0.04	0.16	9.26	174.59	15.35	30.70	179.10
C3	0.04	0.17	8.93	167.54	14.88	29.76	173.58
C4	0.05	0.20	8.49	158.07	14.70	29.39	171.45

SCALED RANGE (14)= 4.300

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	10.75	206.82	15.99	31.99	286.54
C2	0.04	0.16	9.26	174.59	15.35	30.70	275.05
C3	0.04	0.17	8.93	167.54	14.88	29.76	266.57
C4	0.05	0.20	8.49	158.07	14.70	29.39	263.29

SCALED RANGE (15)= 5.600

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	10.75	206.82	15.99	31.99	373.17
C2	0.04	0.16	9.26	174.59	15.35	30.70	358.20
C3	0.04	0.17	8.93	167.54	14.88	29.76	347.16
C4	0.05	0.20	8.49	158.07	14.70	29.39	342.89

SCALED RANGE (16)= 9.600

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	10.75	206.82	15.99	31.99	639 73
C2	0.04	0.16	9.26	174.59	15.35	30.70	614.06
C3	0.04	0.17	8.93	167.54	14.88	29.76	595.13
C4	0.05	0.20	8.49	158.07	14.70	29.39	587.81

SCALED RANGE (17)= 13.000

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	10.75	206.82	15.99	31.99	866.29
C2	0.04	0.16	9.26	174.59	15.35	30.70	831.54
C3	0.04	0.17	8.93	167.54	14.88	29.76	805.90
C4	0.05	0.20	8.49	158.07	14.70	29.39	796.00

SCALE	D RANGE	(18)=	34.000				
COMPONENT	C1	C2	QM	x	R1	RE	R
C1 C2	0.03	0.10 0.16	10.75 9.26	206.82 174.59	15.99 15.35	31.99 30.70	2265.69 2174.79

C3	0.04	0.17	8.93	167.54	14.88	29.76	2107.75
C4	0.05	0.20	8.49	158.07	14.70	29.39	2081.84

MASS OF FLAMMABLE GAS IN THE CLOUD(2)= 5000.0

SCALED RANGE (1)= 0.480

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	18.36	381.01	27.33	54.67	54.67
C2	0.04	0.16	15.83	321.63	26.24	52.47	52.47
C3	0.04	0.17	15.27	308.65	25.43	50.86	50.86
C4	0.05	0.20	14.51	291.21	25.12	50.23	50.23

SCALED RANGE (2)= 0.660

COMPONENT	C1	C2	QM	х	R1	RE	R
C1	0.03	0.10	18.36	381.01	27.33	54.67	75 17
C2	0.04	0.16	15.83	321.63	26.24	52.47	72.15
C3	0.04	0.17	15.27	308.65	25.43	50.86	69.93
C4	0.05	0.20	14.51	291.21	25.12	50.23	69.07

SCALED RANGE (3)= 0.670

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	18.36	381.01	27.33	54.67	76.31
C2	0.04	0.16	15.83	321.63	26.24	52.47	73.24
C3	0.04	0.17	15.27	308.65	25.43	50.86	70.99
C4	0.05	0.20	14.51	291.21	25.12	50.23	70.11

SCALE	RANGE	(4)=	0.700				
COMPONENT	C1	C2	QM	x	R1	RE	R

C1	0.03	0.10	18.36	381.01	27.33	54.67	79.72
C2	0.04	0.16	15.83	321.63	26.24	52.47	76.52
C3	0.04	0.17	15.27	308.65	25.43	50.86	74.16
C4	0.05	0.20	14.51	291.21	25.12	50.23	73.25

SCALED RANGE (5)= 0.750

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	18.36	381.01	27.33	54.67	85.42
C2	0.04	0.16	15.83	321.63	26.24	52.47	81.99
C3	0.04	0.17	15.27	308.65	25.43	50.86	79.46
C4	0.05	0.20	14.51	291.21	25.12	50.23	78.49

SCALED RANGE (6)=

0.900

COMPONENT	C1	C2	QM	х	R1	RE	R
C1	0.03	0.10	18.36	381.01	27.33	54.67	102 50
C2	0.04	0.16	15.83	321.63	26.24	52.47	98 39
C3	0.04	0.17	15.27	308.65	25.43	50.86	95.35
C4	0.05	0.20	14.51	291.21	25.12	50.23	94.18

SCALED RANGE (7)= 0.980

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	18.36	381.01	27.33	54 67	111 61
C2	0.04	0.16	15.83	321.63	26.24	52.47	107 13
C3	0.04	0.17	15.27	308.65	25.43	50.86	103.83
C4	0.05	0.20	14.51	291.21	25.12	50.23	102.55

SCALED RANGE (8)= 1.090

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	18.36	381.01	27.33	54 67	124 14
C2	0.04	0.16	15.83	321.63	26.24	52.47	119.16
C3	0.04	0.17	15.27	308.65	25.43	50.86	115.48
C4	0.05	0.20	14.51	291.21	25.12	50.23	114.06

SCALED RANGE (9)= 1.120

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	18.36	381.01	27.33	54.67	127.55
C2	0.04	0.16	15.83	321.63	26.24	52.47	122.44
C3	0.04	0.17	15.27	308.65	25.43	50.86	118.66
C4	0.05	0.20	14.51	291.21	25.12	50.23	117.20

SCALED RANGE (10)= 1.210

COMPONENT	C1	C2	QM	х	R1	RE	R
C1	0.03	0.10	18.36	381.01	27.33	54.67	137.80
C2	0.04	0.16	15.83	321.63	26.24	52.47	132.28
C3	0.04	0.17	15.27	308.65	25.43	50.86	128.20
C4	0.05	0.20	14.51	291.21	25.12	50.23	126.62

SCALED RANGE (11)= 1.530

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	18.36	381.01	27.33	54.67	174 25
C2	0.04	0.16	15.83	321.63	26.24	52.47	167 26
C3	0.04	0.17	15.27	308.65	25.43	50.86	162.10
C4	0.05	0.20	14.51	291.21	25.12	50.23	160.11

SCALED RANGE (12)= 2.100

COMPONENT	C1	C2	QM	х	R1	RE	R
C1	0.03	0.10	18.36	381.01	27.33	54.67	239 17
C2	0.04	0.16	15.83	321.63	26.24	52.47	229.57
C3	0.04	0.17	15.27	308.65	25.43	50.86	222.49
C4	0.05	0.20	14.51	291.21	25.12	50.23	219.76

SCALE	D RANGE (13)=	2.800				
COMPONENT	C1	C2	QM	x	R1	RE	R
C1 C2	0.03	0.10 0.16	18.36 15.83	381.01 321.63	27.33 26.24	54.67 52.47	318.89 306.09

C3	0.04	0.17	15.27	308.65	25.43	50.86	296.66
C4	0.05	0.20	14.51	291.21	25.12	50.23	293.01

SCALED RANGE (14)= 4.300

COMPONENT	C1	C2	QM	х	R1	RE	R
C1	0.03	0.10	18.36	381.01	27.33	54.67	489 72
C2	0.04	0.16	15.83	321.63	26.24	52.47	470.07
C3	0.04	0.17	15.27	308.65	25.43	50.86	455.58
C4	0.05	0.20	14.51	291.21	25.12	50.23	449.98

SCALED RANGE (15)= 5.600

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	18.36	381.01	27.33	54 67	637 77
C2	0.04	0.16	15.83	321.63	26.24	52.47	612 19
C3	0.04	0.17	15.27	308.65	25.43	50.86	593 31
C4	0.05	0.20	14.51	291.21	25.12	50.23	586.02

SCALED RANGE (16)= 9.600

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	18.36	381.01	27 33	54 67	1092 22
C2	0.04	0.16	15.83	321.63	26.24	52.47	1049 46
C3	0.04	0.17	15.27	308.65	25.43	50.86	1017.11
C4	0.05	0.20	14.51	291.21	25.12	50.23	1004.61

SCALED RANGE (17)= 13.000

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	18.36	381.01	27.33	54 67	1480 55
C2	0.04	0.16	15.83	321.63	26.24	52.47	1421 15
C3	0.04	0.17	15.27	308.65	25.43	50.86	1377.34
C4	0.05	0.20	14.51	291.21	25.12	50.23	1360.41

SCALED RANGE (18)= 34.000

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	18.36	381.01	27.33	54.67	3872.20
C2	0.04	0.16	15.83	321.63	26.24	52.47	3716.85
C3	0.04	0.17	15.27	308.65	25.43	50.86	3602.27
C4	0.05	0.20	14.51	291.21	25.12	50.23	3557.99

MASS OF FLAMMABLE GAS IN THE CLOUD(3)= 10000.0

SCALED RANGE (1)= 0.480

COMPONENT	C1	C2	QM	х	R1	RE	R
C1	0.03	0.10	23.13	495.69	34.43	68 86	68 96
C2	0.04	0.16	19.94	418.44	33.05	66.10	66 10
C3	0.04	0.17	19.23	401.55	32.03	64.06	64.06
C4	0.05	0.20	18.27	378.86	31.64	63.27	63.27

SCALED RANGE (2)= 0.660

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	23.13	495.69	34.43	68 86	04 69
C2	0.04	0.16	19.94	418.44	33.05	66.10	90.88
C3	0.04	0.17	19.23	401.55	32.03	64.06	88.08
C4	0.05	0.20	18.27	378.86	31.64	63.27	87.00

SCALED RANGE (3)= 0.670

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	23.13	495.69	34,43	68.86	96 12
C2	0.04	0.16	19.94	418.44	33.05	66.10	92.26
C3	0.04	0.17	19.23	401.55	32.03	64.06	89.42
C4	0.05	0.20	18.27	378.86	31.64	63.27	88.32

SCALED RANGE (4)= 0.700

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	23.13	495.69	34.43	68.86	100.42
C2	0.04	0.16	19.94	418.44	33.05	66.10	96.39
C3	0.04	0.17	19.23	401.55	32.03	64.06	93.42
C4	0.05	0.20	18.27	378.86	31.64	63.27	92.27

SCALED RANGE (5)= 0.750

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	23.13	495.69	34.43	68.86	107 59
C2	0.04	0.16	19.94	418.44	33.05	66.10	103.28
C3	0.04	0.17	19.23	401.55	32.03	64.06	100.09
C4	0.05	0.20	18.27	378.86	31.64	63.27	98.86

SCALED RANGE (6)= 0.900

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	23.13	495.69	34.43	68 86	129 11
C2	0.04	0.16	19.94	418.44	33.05	66.10	123.93
C3	0.04	0.17	19.23	401.55	32.03	64.06	120.11
C4	0.05	0.20	18.27	378.86	31.64	63.27	118.63

SCALED RANGE (7)= 0.980

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	23.13	495.69	34.43	68 86	140 59
C2	0.04	0.16	19.94	418.44	33.05	66.10	134.95
C3	0.04	0.17	19.23	401.55	32.03	64.06	130.79
C4	0.05	0.20	18.27	378.86	31.64	63.27	129.18

SCALE	D RANGE (8)=	1.090				
COMPONENT	C1	C2	QM	x	R1	RE	R
C1 C2	0.03 0.04	0.10 0.16	23.13 19.94	495.69 418.44	34.43 33.05	68.86 66.10	156.37 150.09

C3	0.04	0.17	19.23	401.55	32.03	64.06	145.47
C4	0.05	0.20	18.27	378.86	31.64	63.27	143.68

SCALED RANGE (9)= 1.120

COMPONENT	C1	C2	QM	х	R1	RE	R
C1	0.03	0.10	23.13	495.69	34.43	68.86	160.67
C2	0.04	0.16	19.94	418.44	33.05	66.10	154.23
C3	0.04	0.17	19.23	401.55	32.03	64.06	149.47
C4	0.05	0.20	18.27	378.86	31.64	63.27	147.63

SCALED RANGE (10)= 1.210

COMPONENT	C1	C2	QM	х	R1	RE	R
C1	0.03	0.10	23.13	495.69	34.43	68 86	173 59
C2	0.04	0.16	19.94	418.44	33.05	66.10	166 62
C3	0.04	0.17	19.23	401.55	32.03	64.06	161.48
C4	0.05	0.20	18.27	378.86	31.64	63.27	159.50

SCALED RANGE (11)= 1.530

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	23.13	495.69	34.43	68.86	219 49
C2	0.04	0.16	19.94	418.44	33.05	66.10	210.68
C3	0.04	0.17	19.23	401.55	32.03	64.06	204.19
C4	0.05	0.20	18.27	378.86	31.64	63.27	201.68

SCALED RANGE (12)= 2.100

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	23.13	495.69	34.43	68.86	301 26
C2	0.04	0.16	19.94	418.44	33.05	66,10	289,17
C3	0.04	0.17	19.23	401.55	32.03	64.06	280.26
C4	0.05	0.20	18.27	378.86	31.64	63.27	276.81

SCALED RANGE (13)= 2.800

α	MPONENT	C1	C2	QM	x	R1	RE	R
	C1	0.03	0.10	23.13	495.69	34.43	68,86	401.68
	C2	0.04	0.16	19.94	418.44	33.05	66.10	385.56
	C3	0.04	0.17	19.23	401.55	32.03	64.06	373.68
	C4	0.05	0.20	18.27	378.86	31.64	63.27	369.09

SCALED RANGE (14)= 4.300

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	23.13	495.69	34.43	68.86	616.87
C2	0.04	0.16	19.94	418.44	33.05	66.10	592.12
C3	0.04	0.17	19.23	401.55	32.03	64.06	573.86
C4	0.05	0.20	18.27	378.86	31.64	63.27	566.81

SCALED RANGE (15)= 5.600

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	23.13	495.69	34,43	68.86	803 36
C2	0.04	0.16	19.94	418.44	33.05	66.10	771.13
C3	0.04	0.17	19.23	401.55	32.03	64.06	747.36
C4	0.05	0.20	18.27	378.86	31.64	63.27	738.17

SCALED RANGE (16)= 9.600

COMPONENT	C1	C2	QM	х	R1	RE	R
C1	0.03	0.10	23.13	495.69	34.43	68.86	1377 19
C2	0.04	0.16	19.94	418.44	33.05	66.10	1321.93
C3	0.04	0.17	19.23	401.55	32.03	64.06	1281.18
C4	0.05	0.20	18.27	378.86	31.64	63.27	1265.43

SCALE	D RANGE (17)= 1	13.000				
COMPONENT	C1	C2	QM	x	R1	RE	R
C1 C2	0.03 0.04	0.10 0.16	23.13 19.94	495.69 418.44	34.43 33.05	68.86 66.10	1864.94 1790.12

C3	0.04	0.17	19.23	401.55	32.03	64.06	1734.94
C4	0.05	0.20	18.27	378.86	31.64	63.27	1713.61

SCALED RANGE (18)= 34.000

COMPONENT	C1	C2	QM	х	R1	RE	R
C1	0.03	0.10	23.13	495.69	34.43	68.86	4877.54
C2	0.04	0.16	19.94	418.44	33.05	66.10	4681.85
C3	0.04	0.17	19.23	401.55	32.03	64.06	4537.52
C4	0.05	0.20	18.27	378.86	31.64	63.27	4481.75

MASS OF FLAMMABLE GAS IN THE CLOUD(4)= 20000.0

SCALED RANGE (1)= 0.480

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	29.14	644.90	43.37	86.74	86.74
C2	0.04	0.16	25.11	544.39	41.63	83.26	83.26
C3	0.04	0.17	24.22	522.42	40.35	80.69	80.69
C4	0.05	0.20	23.02	492.90	39.85	79.70	79.70

SCALED RANGE (2)= 0.660

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	29.14	644.90	43.37	86.74	119.26
C2	0.04	0.16	25.11	544.39	41.63	83.26	114.48
C3	0.04	0.17	24.22	522.42	40.35	80.69	110.95
C4	0.05	0.20	23.02	492.90	39.85	79.70	109.59

SCALE	RANGE	(3)=	0.670				
COMPONENT	C1	C2	QM	х	R1	RE	R

C1	0.03	0.10	29.14	644.90	43.37	86.74	121.07
C2	0.04	0.16	25.11	544.39	41.63	83.26	116.21
C3	0.04	0.17	24.22	522.42	40.35	80.69	112.63
C4	0.05	0.20	23.02	492.90	39.85	79.70	111.25

SCALED RANGE (4)= 0.700

COMPONENT	C1	C2	QM	х	R1	RE	R
C1	0.03	0.10	29.14	644.90	43.37	86.74	126 49
C2	0.04	0.16	25.11	544.39	41.63	83.26	121 42
C3	0.04	0.17	24.22	522.42	40.35	80.69	117.67
C4	0.05	0.20	23.02	492.90	39.85	79.70	116.23

SCALED RANGE (5)=

0.750

COMPONENT	C1	C2	QM	х	R1	RE	R
C1	0.03	0.10	29.14	644.90	43.37	86.74	135 53
C2	0.04	0.16	25.11	544.39	41.63	83.26	130.09
C3	0.04	0.17	24.22	522.42	40.35	80.69	126 08
C4	0.05	0.20	23.02	492.90	39.85	79.70	124.53

SCALED RANGE (6)= 0.900

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	29.14	644.90	43.37	86.74	162 63
C2	0.04	0.16	25.11	544.39	41.63	83.26	156 11
C3	0.04	0.17	24.22	522.42	40.35	80.69	151.30
C4	0.05	0.20	23.02	492.90	39.85	79.70	149.44

SCALED RANGE (7)= 0.980

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	29.14	644.90	43.37	86.74	177 09
C2	0.04	0.16	25.11	544.39	41.63	83.26	169.98
C3	0.04	0.17	24.22	522.42	40.35	80,69	164.74
C4	0.05	0.20	23.02	492.90	39.85	79.70	162.72

SCALED RANGE (8)= 1.090

COMPONENT	C1	C2	QM	X R1	R1	R1 RE	R
C1	0.03	0.10	29.14	644.90	43.37	86.74	196.97
C2	0.04	0.16	25.11	544.39	41.63	83.26	189.06
C3	0.04	0.17	24.22	522.42	40.35	80.69	183.24
C4	0.05	0.20	23.02	492.90	39.85	79.70	180.98

SCALED RANGE (9)= 1.120

COMPONENT	C1	C2	QM	х	R1	RE	R
C1	0.03	0.10	29.14	644.90	43.37	86.74	202.39
C2	0.04	0.16	25.11	544.39	41.63	83.26	194.27
C3	0.04	0.17	24.22	522.42	40.35	80.69	188.28
C4	0.05	0.20	23.02	492.90	39.85	79.70	185.96

SCALED RANGE (10)= 1.210

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	29.14	644.90	43.37	86.74	218 65
C2	0.04	0.16	25.11	544.39	41.63	83.26	209.88
C3	0.04	0.17	24.22	522.42	40.35	80.69	203.41
C4	0.05	0.20	23.02	492.90	39.85	79.70	200.91

SCALED RANGE (11)= 1.530

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	29.14	644.90	43.37	86.74	276.48
C2	0.04	0.16	25.11	544.39	41.63	83.26	265.38
C3	0.04	0.17	24.22	522.42	40.35	80.69	257.20
C4	0.05	0.20	23.02	492.90	39.85	79.70	254.04

D RANGE	(12)=	2.100				
C1	C2	QM	x	R1	RE	R
0.03	0.10	29.14	644.90 544.39	43.37	86.74	379.48
	C1 0.03 0.04	C1 C2 0.03 0.10 0.04 0.16	$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	$\begin{array}{c ccccccccccccccccccccccccccccccccccc$

C3	0.04	0.17	24.22	522.42	40.35	80.69	353.02
C4	0.05	0.20	23.02	492.90	39.85	79.70	348.68

SCALED RANGE (13)= 2.800

COMPONENT	C1	C2	QM	х	R1	RE	R
C1	0.03	0.10	29.14	644.90	43.37	86.74	505.97
C2	0.04	0.16	25.11	544.39	41.63	83.26	485.67
C3	0.04	0.17	24.22	522.42	40.35	80.69	470.70
C4	0.05	0.20	23.02	492.90	39.85	79.70	464.91

SCALED RANGE (14)= 4.300

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	29.14	644.90	43.37	86.74	777 02
C2	0.04	0.16	25.11	544.39	41.63	83.26	745.85
C3	0.04	0.17	24.22	522.42	40.35	80.69	722.86
C4	0.05	0.20	23.02	492.90	39.85	79.70	713.97

SCALED RANGE (15)= 5.600

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COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	29.14	644.90	43.37	86.74	1011 94
C2	0.04	0.16	25.11	544.39	41.63	83.26	971.34
C3	0.04	0.17	24.22	522.42	40.35	80.69	941.39
C4	0.05	0.20	23.02	492.90	39.85	79.70	929.82

SCALED RANGE (16)= 9.600

1734 75
1665.15
1613.82
1593.98
SCALED RANGE (17)= 13.000

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	29.14	644.90	43.37	86.74	2349.14
C2	0.04	0.16	25.11	544.39	41.63	83.26	2254.89
C3	0.04	0.17	24.22	522.42	40.35	80.69	2185.38
C4	0.05	0.20	23.02	492.90	39.85	79.70	2158.51

SCALED RANGE (18)= 34.000

COMPONENT	C1	C1 C2	QM	x	R1	RE	R
C1	0.03	0.10	29.14	644.90	43.37	86.74	6143.90
C2	0.04	0.16	25.11	544.39	41.63	83.26	5897.40
C3	0.04	0.17	24.22	522.42	40.35	80.69	5715.60
C4	0.05	0.20	23.02	492.90	39.85	79.70	5645.34

MASS OF FLAMMABLE GAS IN THE CLOUD(5)= 30000.0

SCALED RANGE (1)= 0.480

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	33.35	752.21	49.64	99.28	99 28
C2	0.04	0.16	28.74	634.98	47.65	95.29	95.29
C3	0.04	0.17	27.72	609.35	46.18	92.36	92.36
C4	0.05	0.20	26.35	574.92	45.61	91.22	91.22

SCALED RANGE (2)= 0.660

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	33.35	752.21	49.64	99.28	136.50
C2	0.04	0.16	28.74	634.98	47.65	95.29	131.03
C3	0.04	0.17	27.72	609.35	46.18	92.36	126.99
C4	0.05	0.20	26.35	574.92	45.61	91.22	125.43

SCALED RANGE (3)= 0.670

COMPONENT	C1	C2	QM	х	R1	RE	R
C1	0.03	0.10	33.35	752.21	49.64	99.28	138.57
C2	0.04	0.16	28.74	634.98	47.65	95.29	133.01
C3	0.04	0.17	27.72	609.35	46.18	92.36	128.91
C4	0.05	0.20	26.35	574.92	45.61	91.22	127.33

SCALED RANGE (4)= 0.700

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	33.35	752.21	49.64	99.28	144.78
C2	0.04	0.16	28.74	634.98	47.65	95.29	138.97
C3	0.04	0.17	27.72	609.35	46.18	92.36	134.69
C4	0.05	0.20	26.35	574.92	45.61	91.22	133.03

SCALED RANGE (5)= 0.750

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	33.35	752.21	49.64	99.28	155 12
C2	0.04	0.16	28.74	634.98	47.65	95.29	148.90
C3	0.04	0.17	27.72	609.35	46.18	92.36	144.31
C4	0.05	0.20	26.35	574.92	45.61	91.22	142.53

SCALED RANGE (6)= 0.900

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	33.35	752.21	49.64	99.28	186.14
C2	0.04	0.16	28.74	634.98	47.65	95.29	178.67
C3	0.04	0.17	27.72	609.35	46.18	92.36	173.17
C4	0.05	0.20	26.35	574.92	45.61	91.22	171.04

SCALED	RANGE	(7)=	0.980
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COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	33.35	752.21	49.64	99.28	202.69
C2	0.04	0.16	28.74	634.98	47.65	95.29	194.56

C3	0.04	0.17	27.72	609.35	46.18	92.36	188.56
C4	0.05	0.20	26.35	574.92	45.61	91.22	186.24

SCALED RANGE (8)= 1.090

COMPONENT	C1	C2	QM	х	R1	RE	R
C1	0.03	0.10	33.35	752.21	49.64	99.28	225.44
C2	0.04	0.16	28.74	634.98	47.65	95.29	216.39
C3	0.04	0.17	27.72	609.35	46.18	92.36	209.72
C4	0.05	0.20	26.35	574.92	45.61	91.22	207.15

SCALED RANGE (9)= 1.120

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	33.35	752.21	49.64	99.28	231.64
C2	0.04	0.16	28.74	634.98	47.65	95.29	222.35
C3	0.04	0.17	27.72	609.35	46.18	92.36	215.50
C4	0.05	0.20	26.35	574.92	45.61	91.22	212.85

SCALED RANGE (10)= 1.210

COMPONENT	C1	C2	QM	х	R1	RE	R
C1	0.03	0.10	33.35	752.21	49.64	99.28	250.26
C2	0.04	0.16	28.74	634.98	47.65	95.29	240.22
C3	0.04	0.17	27.72	609.35	46.18	92.36	232.81
C4	0.05	0.20	26.35	574.92	45.61	91.22	229.95

SCALED RANGE (11)= 1.530

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	33.35	752.21	49.64	99.28	316.44
C2	0.04	0.16	28.74	634.98	47.65	95.29	303.75
C3	0.04	0.17	27.72	609.35	46.18	92.36	294.38
C4	0.05	0.20	26.35	574.92	45.61	91.22	290.76

SCALED RANGE (12)= 2.100

COMPONENT	C1	C2	QM	х	R1	RE	R
C1	0.03	0.10	33.35	752.21	49.64	99.28	434.33
C2	0.04	0.16	28.74	634.98	47.65	95.29	416.91
C3	0.04	0.17	27.72	609.35	46.18	92.36	404.06
C4	0.05	0.20	26.35	574.92	45.61	91.22	399.09

SCALED RANGE (13)= 2.800

COMPONENT	C1	C2	QM	х	R1	RE	R
C1	0.03	0.10	33.35	752.21	49.64	99.28	579.11
C2	0.04	0.16	28.74	634.98	47.65	95.29	555.88
C3	0.04	0.17	27.72	609.35	46.18	92.36	538.74
C4	0.05	0.20	26.35	574.92	45.61	91.22	532.12

SCALED RANGE (14)= 4.300

COMPONENT	C1	C2	QM	х	R1	RE	R
C1	0.03	0.10	33.35	752.21	49.64	99.28	889.35
C2	0.04	0.16	28.74	634.98	47.65	95.29	853.67
C3	0.04	0.17	27.72	609.35	46.18	92.36	827.35
C4	0.05	0.20	26.35	574.92	45.61	91.22	817.18

SCALED RANGE (15)= 5.600

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	33.35	752.21	49.64	99.28	1158.22
C2	0.04	0.16	28.74	634.98	47.65	95.29	1111.75
C3	0.04	0.17	27.72	609.35	46.18	92.36	1077.48
C4	0.05	0.20	26.35	574.92	45.61	91.22	1064.24

SCALE	D RANGE (16)=	9.600				
COMPONENT	C1	C2	QM	x	R1	RE	R
C1 C2	0.03 0.04	0.10 0.16	33.35 28.74	752.21 634.98	49.64 47.65	99.28 95.29	1985.52 1905.86

C3	0.04	0.17	27.72	609.35	46.18	92.36	1847.11
C4	0.05	0.20	26.35	574.92	45.61	91.22	1824.40

SCALED RANGE (17)= 13.000

COMPONENT	C1	C2	QM	х	R1	RE	R
C1	0.03	0.10	33.35	752.21	49.64	99.28	2688.73
C2	0.04	0.16	28.74	634.98	47.65	95.29	2580.85
C3	0.04	0.17	27.72	609.35	46.18	92.36	2501.29
C4	0.05	0.20	26.35	574.92	45.61	91.22	2470.55

SCALED RANGE (18)= 34.000

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	33.35	752.21	49.64	99.28	7032 06
C2	0.04	0.16	28.74	634.98	47.65	95.29	6749.93
C3	0.04	0.17	27.72	609.35	46.18	92.36	6541.85
C4	0.05	0.20	26.35	574.92	45.61	91.22	6461.43

MASS OF FLAMMABLE GAS IN THE CLOUD(6)= 40000.0

SCALED RANGE (1)= 0.480

11.11	and the second						
п	C1	C2	QM	x	R1	RE	R
	0.03	0.10	36.70	839.01	54.63	109.26	109.26
	0.04	0.16	31.63	708.26	52.44	104.87	104.87
	0.04	0.17	30.51	679.66	50.82	101.64	101.64
	0.05	0.20	28.99	641.26	50.20	100.39	100.39
ALED	RANGE (2)=	0.660				
п	C1	C2	CM	x	R1	RE	R
	ALED I	TT C1 0.03 0.04 0.04 0.05 CALED RANGE (TT C1	$\begin{array}{c ccccc} & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ \end{array} $	AT C1 C2 GM 0.03 0.10 36.70 0.04 0.16 31.63 0.04 0.17 30.51 0.05 0.20 28.99 CALED RANGE (2)= 0.660 AT C1 C2 GM	$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	$\begin{array}{c ccccccccccccccccccccccccccccccccccc$

C1	0.03	0.10	36.70	839.01	54.63	109.26	150.23
C2	0.04	0.16	31.63	708.26	52.44	104.87	144.20
C3	0.04	0.17	30.51	679.66	50.82	101.64	139.76
C4	0.05	0.20	28.99	641.26	50.20	100.39	138.04

SCALED RANGE (3)= 0.670

	the man line and has been the set						
COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	36.70	839.01	54.63	109.26	152.50
C2	0.04	0.16	31.63	708.26	52.44	104.87	146.39
C3	0.04	0.17	30.51	679.66	50.82	101.64	141.87
C4	0.05	0.20	28.99	641.26	50.20	100.39	140.13

SCALED RANGE (4)= 0.700

COMPONENT	C1	C2	QM	х	R1	RE	R
C1	0.03	0.10	36.70	839.01	54.63	109.26	159 33
C2	0.04	0.16	31.63	708.26	52.44	104.87	152.94
C3	0.04	0.17	30.51	679.66	50.82	101.64	148.23
C4	0.05	0.20	28.99	641.26	50.20	100.39	146.40

SCALED RANGE (5)= 0.750

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	36.70	839.01	54.63	109.26	170 71
C2	0.04	0.16	31.63	708.26	52.44	104.87	163.86
C3	0.04	0.17	30.51	679.66	50.82	101.64	158.81
C4	0.05	0.20	28.99	641.26	50.20	100.39	156.86

SCALED RANGE (6)= 0.900

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	36.70	839.01	54.63	109.26	204.86
C2	0.04	0.16	31.63	708.26	52.44	104.87	196.64
C3	0.04	0.17	30.51	679.66	50.82	101.64	190.58
C4	0.05	0.20	28.99	641.26	50.20	100.39	188.23

SCALED RANGE (7)= 0.980

COMPONENT	C1	C2	QM	X	R1	RE	R
C1	0.03	0.10	36.70	839.01	54.63	109.26	223.07
C2	0.04	0.16	31.63	708.26	52.44	104.87	214.12
C3	0.04	0.17	30.51	679.66	50.82	101.64	207.52
C4	0.05	0.20	28.99	641.26	50.20	100.39	204.97

SCALED RANGE (8)= 1.090

COMPONENT	C1	C2	QM	х	R1	RE	R
C1	0.03	0.10	36.70	839.01	54.63	109.26	248,10
C2	0.04	0.16	31.63	708.26	52.44	104.87	238.15
C3	0.04	0.17	30.51	679.66	50.82	101.64	230.81
C4	0.05	0.20	28.99	641.26	50.20	100.39	227.97

SCALED RANGE (9)= 1.120

COMPONENT	C1	C2	QM	x	R1	RE	R
Cĺ	0.03	0.10	36.70	839.01	54.63	109 26	254 03
C2	0.04	0.16	31.63	708.26	52.44	104.87	244.70
C3	0.04	0.17	30.51	679.66	50.82	101.64	237.16
C4	0.05	0.20	28.99	641.26	50.20	100.39	234.25

SCALED RANGE (10)= 1.210

COMPONENT	C1	C2	QM	x	R1	RE	R
C1	0.03	0.10	36.70	839.01	54,63	109.26	275 42
C2	0.04	0.16	31.63	708.26	52.44	104.87	264.37
C3	0.04	0.17	30.51	679.66	50.82	101.64	256.22
C4	0.05	0.20	28.99	641.26	50.20	100.39	253.07

SCALE	D RANGE	(11)=	1.530				
COMPONENT	C1	 C2	QM	x	R1	RE	R
C1 C2	0.03 0.04	0.10 0.16	36.70 31.63	839.01 708.26	54.63 52.44	109.26 104.87	348.26 334.28

NOMENCLATURE

а	numerical exponent
a	sonic velocity in air (m/sec), equation 7.6
ao	sonic velocity in air at the instant of vessel rupture
	(m/sec)
a ₂	sonic velocity in the sphere at the instant of vessel rupture
	(m/sec)
Α	numerical constant
Α	surface area of pool (m ²), equation 6.73
b	numerical exponent
В	numercial constant
С	specific heat of combusion products (kJ/kg K)
С	calorific value of liquid (kJ/kg), equation 6.69
Ca	specific heat of air (kJ/kg K)
Cpm	mean specific heat of combustion products (kJ/kg K)
C(r)	vapour concentration at distance r from centre (g/m^3)
Co	vapour concentration at centre of cloud (g/m3)
C1	concentration at lower flammable limit (g/m ³)
C ₂	concentration at upper flammable limit (g/m ³)
d	distance from tank to receiver (m)
D	equivalent spherical diameter of fireball (m)
D	diameter of pool (m) equation 6.73
Dc	equivalent spherical diameter at completion of combustion
	(m)
Dc	the initial hemispherical diameter of the cloud at NTP
	(m), equation 7.6
D ₁	equivalent spherical diameter at neutral buoyancy (m)
Do	equivalent spherical diameter of gas cloud (m)
Ds	equivalent spherical diameter of hot shell (m)
(dD/dt) _c	constant rate of increase of diameter during combustion
	(m/sec)
E	flame surface emissive power (kW/m^2), equation 6.73
E ₁	the effective energy stored in a pressurised sphere of
	ideal gas (J)
Eo	total combustion energy released (MJ)

ΔΕ	Isentropic expansion energy (J)
f	fraction of fuel burnt
f	fraction of heat of combustion of fuel which is radiated, equation 6.72
fc	fraction of fuel burnt at the end of combustion
f ₁	fraction of fuel burnt to produce neutral buoyancy
F ₂₁	view factor from target to the fireball surface
h	net calorific value of fuel (kJ/kg)
н	elevation of bottom of fireball above ground (m)
H _c	heat of combustion (kJ/kg), equation 6.72
ΔH	the enthalpy change
ΔH°c	standard heat of combustion (cal/g mole)
ΔH°p	the enthalpy change of the products
k	constant, equation 6.69
k	constant appropriate to various degrees of damage, equation 7.1
L	distance from surface of fireball to target (m)
L	length of flame (m), equation 6.73
m	mass of fuel in cloud initially (kg)
m1	mass of water required to remove the heat (kg)
m	fuel burning rate (kg/m ² .sec)
М	molecular weight of the flammable material
М	mean molecular weight of stoichiometric combustion products
n	increase in total number of moles per mole of flammable
	gas burnt stoichiometrically
n	number of moles, equation 6.63
Pc°	rate of pressure rise at the centre of the initial explosion
Pf	final absolute pressure (N/m ²)
Po	initial absolute pressure (N/m ²)
Po	ambient atmospheric pressure (N/m^2) , equation 5.1
P ₂	initial absolute pressure in the sphere at the instant of
	burst (kN/m^2) , equation 5.1
P ₂	dimensionless overpressure in the sphere
Ps	shock overpressure KN/m ²
Ps	shock overpressure (kN/m ²), equation 7.2
Ps	dimensionless shock overpressure
Pso	initial reduced shock overpressure

q	radiation intensity (kW/m ²)
q ₁	emitted flux density from fireball surface (kW/m ²)
92	incident flux density from a fireball surface (kW/m ²)
Q	heat flux density function (kW/m ²)
Q1	quantity of heat radiated (kW)
Q2	radiant flux intensity (kW/m ²)
r	mass ratio of combustion products per unit mass of
	flammable gas, from stoichiometric combustion
r	liquid burning rate (m/sec), equation 6.69
r	radius (m), equation 7.8
r ₁	radius at which the concentration is at the lower
	flammable limit (m)
ro	initial radius of sphere (m)
R	distance radially along the ground from the centre-line of
	a fireball (m)
R	distance from the explosion (ft), equation 7.1
R	radial distance from centre of explosion (m), equation 7.2
Re	radius when the flame front reaches the periphery (m)
Ro	scaling radius based on energy (m)
Rs	radial distance at shock front condition
Rs	dimensionless shock radius
R _{so}	scaled shock front radius at the instant of failure
Rs	radius at which blast wave becomes shock wave (m),
	equation 7.4
t	time since ignition (sec)
tc	duration of combustion (sec)
te	exposure time (sec)
to	duration of explosive release of energy (sec)
ts	time taken for a pressure wave to transform into a shock
	wave (sec)
t	theoretical flame temperature (c)
Т	absolute temperature (k)
Tc	isothermal temperature of fireball (k)
T _f	flame surface temperature (k)
Ts	temperature of hot shell (k)
ΔΤ	temperature rise of the water (k)
U	average radial velocity of air through the hot shell (m/sec)
v	volumetric rate of flow of air into fireball (m ³ /sec)

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v	volume of sphere (m ³)
v ₁	water capacity of vessel (litres)
Vo	original volume (m ³)
Vo	original volume of sphere (m^3) , equation 5.5
\mathbf{W}_1	maximum permitted LPG fill by weight (kg)
w	weight of TNT (lb), equation 7.1
w	total quantity of vapour in cloud (g), equation 7.16
Z	percent of damage

Greek Letters

τ	atmospheric transmissivity
α	absorptivity
3	emissivity
θ	Stefan-Boltzmann constant (kW/m ² K ⁴)
υ	stoichiometric molar ratio of air to flammable gas
σ	standard deviation of concentration distribution (m)
$\sigma_{\rm m}$	standard deviation when cloud is at maximum potential
	(m)
γ	ratio of specific heats
γο	specific heat ratio of air
γ ₁	specific heat ratio of stored gas
ρ	density (kg/m ³)
Pref	relative density of LPG at the reference temperature for
	filling ratio

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