According to OSHA Hazard Communication Standard, 29 CFR 1910.1200

# Fuel gases refinery

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#### **SECTION 1. IDENTIFICATION**

Product name : Fuel gases refinery

Product code : 002D4317

CAS-No. : 68308-27-0

Manufacturer or supplier's details

Manufacturer/Supplier : Vertex Refining Alabama LLC

400 Industrial Pkwy

Ext. East

Saraland, AL 36571

SDS Request : 251-679-7180 Customer Service : 251-679-7180

**Emergency telephone number** 

Spill Information : 1-800-424-9300 Health Information : 1-800-424-9300

Recommended use of the chemical and restrictions on use

Recommended use

:

Intermediate Refinery Stream.

Restrictions on use : This product must not be used in applications other than those

listed in Section 1 without first seeking the advice of the sup-

plier.

#### **SECTION 2. HAZARDS IDENTIFICATION**

GHS classification in accordance with 29 CFR 1910.1200

Flammable gases : Category 1

Gases under pressure : Compressed gas

Carcinogenicity : Category 1A

Germ cell mutagenicity : Category 1B

Reproductive toxicity : Category 1A

Acute toxicity : Category 4

Specific target organ toxicity

Category 2

- repeated exposure

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## **GHS** label elements

Hazard pictograms









Signal word : Danger

Hazard statements : PHYSICAL HAZARDS:

H220 Extremely flammable gas.

H280 Contains gas under pressure; may explode if heated.

HEALTH HAZARDS: H350 May cause cancer.

H340 May cause genetic defects.

H360 May damage fertility or the unborn child.

H332 Harmful if inhaled.

H373 May cause damage to organs through prolonged or re-

peated exposure.

**ENVIRONMENTAL HAZARDS:** 

Not classified as an environmental hazard under GHS criteria.

Precautionary statements : P102 Keep out of reach of children.

## Prevention:

P201 + P202 Obtain special instructions before use. Do not handle until all safety precautions have been read and understood.

P210 Keep away from heat/sparks/open flames/hot surfaces. No smoking.

P260 Do not breathe dust/ fume/ gas/ mist/ vapours/ spray. P261 Avoid breathing dust/ fume/ gas/ mist/ vapours/ spray.

P271 Use only outdoors or in a well-ventilated area.

P280 Wear protective gloves/ protective clothing/ eye protection/ face protection.

## Response:

P304 + P340 IF INHALED: Remove victim to fresh air and keep at rest in a position comfortable for breathing.

P308 + P313 IF exposed or concerned: Get medical advice/ attention.

P312 Call a POISON CENTER or doctor/ physician if you feel unwell.

P377 Leaking gas fire: Do not extinguish, unless leak can be stopped safely.

P381 Eliminate all ignition sources if safe to do so.

## Storage:

P410 + P403 Protect from sunlight. Store in a well-ventilated place.

P405 Store locked up.

#### Disposal:

P501 Dispose of contents and container to appropriate waste

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site or reclaimer in accordance with local and national regula-

tions.

#### Other hazards which do not result in classification

Vapours are heavier than air. Vapours may travel across the ground and reach remote ignition sources causing a flashback fire danger.

This material has the potential to be a static accumulator.

Electrostatic charges may be generated during pumping. Electrostatic discharge may cause fire. High gas concentrations will displace available air; unconsciousness and death may occur suddenly from lack of oxygen.

This product contains benzene which may cause leukaemia (AML - acute myelogenous leukaemia).

May cause MDS (Myelodysplastic Syndrome).

Hydrogen sulphide (H2S), an extremely flammable and toxic gas, and other hazardous vapours may evolve and collect in the headspace of storage tanks, transport vessels and other enclosed containers.

The classification of this material is based on OSHA HCS 2012 criteria.

#### **SECTION 3. COMPOSITION/INFORMATION ON INGREDIENTS**

Substance / Mixture : Substance

#### **Hazardous components**

Chemical name	Synonyms	CAS-No.	Concentration (% w/w)
Fuel Gases, Refinery	Fuel gases,	68308-27-0	100
	refinery		

#### **Further information**

### Contains:

Chemical name	Identification number	Concentration [%]
1,3-butadiene	106-99-0, 203-450-8	0.1 - 5
benzene	71-43-2, 200-753-7	0.1 - 0.3
Hydrogen sulfide	7783-06-4, 231-977-3	- < 1
Carbon dioxide	124-38-9, 204-696-9	- < 1

#### **SECTION 4. FIRST-AID MEASURES**

General advice : Vapourisation of H2S that has been trapped in clothing can be

dangerous to rescuers. Maintain respiratory protection to avoid contamination from the victim to rescuer. Mechanical ventilation should be used to resuscitate if at all possible.

If inhaled : Remove to fresh air. Do not attempt to rescue the victim un-

less proper respiratory protection is worn. If the victim has difficulty breathing or tightness of the chest, is dizzy, vomiting, or unresponsive, give 100% oxygen with rescue breathing or Cardio-Pulmonary Resuscitation as required and transport to

the nearest medical facility.

Respiratory irritation signs and symptoms may include a temporary burning sensation of the nose and throat, coughing,

and/or difficulty breathing.

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In case of skin contact : Remove contaminated clothing. Immediately flush skin with

large amounts of water for at least 15 minutes, and follow by washing with soap and water if available. If redness, swelling, pain and/or blisters occur, transport to the nearest medical

facility for additional treatment.

Do not remove clothing that adheres to skin due to freezing. In the event of frostbite, slowly warm the exposed area by

rinsing with warm water. Otherwise:

Contaminated clothing may be a fire hazard and therefore should be soaked with water before being removed.

Loosen tight clothing.

Keep warm and at rest.

In case of eye contact : Flush eyes with water while holding eyelids open. Rest eyes

for 30 minutes. If redness, burning, blurred vision, or swelling persist transport to the nearest medical facility for additional

treatment.

Eye irritation signs and symptoms may include a burning sen-

sation, redness, swelling, and/or blurred vision.

In the event of frostbite, slowly warm the exposed area by

rinsing with warm water. Otherwise:

If swallowed : In the unlikely event of ingestion, obtain medical attention

immediately.

Most important symptoms

and effects, both acute and

delayed

High concentrations may cause central nervous system depression resulting in headaches, dizziness and nausea; continued inhalation may result in unconsciousness and/or death.

Protection of first-aiders : When administering first aid, ensure that you are wearing the

appropriate personal protective equipment according to the

incident, injury and surroundings.

Indication of any immediate medical attention and special

treatment needed

Treat symptomatically.

Administer oxygen if necessary.

Hydrogen sulphide (H2S) - CNS asphyxiant. May cause rhinitis, bronchitis and occasionally pulmonary oedema after severe exposure. CONSIDER: Oxygen therapy. Consult a Poi-

son Control Center for guidance.

Potential for cardiac sensitisation, particularly in abuse situations. Hypoxia or negative inotropes may enhance these ef-

fects. Consider: oxygen therapy.

#### **SECTION 5. FIRE-FIGHTING MEASURES**

Suitable extinguishing media : Shut off supply. If not possible and no risk to surroundings, let

the fire burn itself out.

Use foam, water fog for major fires.

Use dry chemical powder, carbon dioxide, sand or earth for

minor fires.

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Unsuitable extinguishing

media

 Do not use direct water jets on the burning product as they could cause a steam explosion and spread of the fire.
 Simultaneous use of foam and water on the same surface is

to be avoided as water destroys the foam.

Specific hazards during fire-

fighting

Hazardous combustion products may include:

Carbon monoxide may be evolved if incomplete combustion

occurs.

Unidentified organic and inorganic compounds.

Sustained fire attack on vessels may result in a Boiling Liquid

Expanding Vapor Explosion (BLEVE).

Contents are under pressure and can explode when exposed

to heat or flames.

The vapour is heavier than air, spreads along the ground and

distant ignition is possible.

Further information : Keep adjacent containers cool by spraying with water.

Special protective equipment

for firefighters

Proper protective equipment including chemical resistant gloves are to be worn; chemical resistant suit is indicated if large contact with spilled product is expected. Self-Contained Breathing Apparatus must be worn when approaching a fire in a confined space. Select fire fighter's clothing approved to

relevant Standards (e.g. Europe: EN469).

## **SECTION 6. ACCIDENTAL RELEASE MEASURES**

Personal precautions, protective equipment and emergency procedures

Shut off leaks, if possible without personal risks. Remove all possible sources of ignition in the surrounding area and evacuate all personnel. Attempt to disperse the gas or to direct its flow to a safe location for example by using fog sprays. Take precautionary measures against static discharge. Ensure electrical continuity by bonding and grounding (earthing) all equipment. Monitor area with combustible gas meter. Test atmosphere for flammable gas concentrations to ensure safe working conditions before personnel are allowed to enter

the area.

Environmental precautions : Use appropriate containment to avoid environmental contami-

nation.

Methods and materials for containment and cleaning up

Allow to evaporate.

Attempt to disperse the gas or to direct its flow to a safe loca-

tion, for example by using fog sprays.

Take precautionary measures against static discharges.

Avoid contact with skin, eyes and clothing. Evacuate the area of all non-essential personnel.

Ventilate contaminated area thoroughly.

Take precautionary measures against static discharges.

Additional advice : For guidance on selection of personal protective equipment

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see Chapter 8 of this Safety Data Sheet.

Notify authorities if any exposure to the general public or the

environment occurs or is likely to occur.

For guidance on disposal of spilled material see Chapter 13 of

this Safety Data Sheet.

Vapour may form an explosive mixture with air.

Risk of explosion. Inform the emergency services if product

enters surface water drains.

#### **SECTION 7. HANDLING AND STORAGE**

Technical measures : Avoid breathing of or direct contact with material. Only use in

well ventilated areas. Wash thoroughly after handling. For guidance on selection of personal protective equipment see

Chapter 8 of this Safety Data Sheet.

Use the information in this data sheet as input to a risk assessment of local circumstances to help determine appropriate controls for safe handling, storage and disposal of this

material.

Air-dry contaminated clothing in a well-ventilated area before

laundering.

Use local exhaust ventilation if there is risk of inhalation of

vapours, mists or aerosols.

Take precautionary measures against static discharges.

Advice on safe handling : The inherent toxic and olfactory (sense of smell) fatiguing

properties of hydrogen sulphide require that air monitoring alarms be used if concentrations are expected to reach harmful levels such as in enclosed spaces, heated transport vessels and spill or leak situations. If the air concentration exceeds 10 ppm, the area should be evacuated unless respira-

tory protection is in use.

Ensure that all local regulations regarding handling and stor-

age facilities are followed.

This product is intended for use in closed systems only.

This product can create a low temperature exposure hazard

when released as a liquid.

Extinguish any naked flames. Do not smoke. Remove ignition

sources. Avoid sparks.

Avoid prolonged or repeated contact with skin.

Electrostatic charges may be generated during pumping. Elec-

trostatic discharge may cause fire.

Earth all equipment.

Use local exhaust ventilation if there is risk of inhalation of

vapours, mists or aerosols.

Avoidance of contact : Oxidizing agents

Product Transfer : Do not use compressed air for filling discharge or handling.

Electrostatic charges may be generated during pumping. Electrostatic discharge may cause fire. Delivery lines may become cold enough to present a cold burns hazard. Ensure electrical continuity by bonding and grounding (earthing) all equipment.

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Restrict line velocity during pumping in order to avoid genera-

tion of electrostatic discharge.

Further information on stor-

age stability

Store only in purpose-designed, appropriately labelled pressure vessels or cylinders.

Must be stored in a well-ventilated area, away from sunlight,

ignition sources and other sources of heat.

Do not store near cylinders containing compressed oxygen or

other strong oxidizers.

The vapours in the head space of the storage vessel may lie in the flammable/explosive range and hence may be flamma-

ble.

Refer to section 15 for any additional specific legislation cov-

ering the packaging and storage of this product.

Packaging material : Suitable material: For containers and container linings, use

materials specifically approved for use with this product., Examples of suitable materials are: PA-11, PEEK, PVDF, PTFE, GRE (Epoxy), GRVE (vinyl ester), Viton (FKM), type F and

GB, Neoprene (CR).

Unsuitable material: Some forms of cast iron., Examples of materials to avoid are: ABS, polymethyl methacrylate (PMMA), polyethylene (PE / HDPE), polypropylene (PP), PVC, natural rubber (NR), Nitrile (NBR) ethylene propylene rubber (EPDM), Butyl (IIR), Hypalon (CSM), polystyrene, polyvinyl chloride (PVC), polyisobutylene., For containers and container linings, aluminium should not be used if there is a

risk of caustic contamination of the product.

Container Advice : Containers, even those that have been emptied, can contain

explosive vapours. Do not cut, drill, grind, weld or perform

similar operations on or near containers.

Specific use(s) : Not applicable

See additional references that provide safe handling practices for liquids that are determined to be static accumulators: American Petroleum Institute 2003 (Protection Against Ignitions Arising out of Static, Lightning and Stray Currents) or National Fire Protection Agency 77 (Recommended Practices

on Static Electricity).

IEC/TS 60079-32-1: Electrostatic hazards, guidance

#### **SECTION 8. EXPOSURE CONTROLS AND PERSONAL PROTECTION**

## Components with workplace control parameters

Components	CAS-No.	Value type (Form of	Control parameters / Permissible concentration	Basis
		exposure)	Concentiation	
1,3-butadiene	106-99-0	TWA	2 ppm	ACGIH

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1,3-butadiene		PEL	1 ppm	OSHA CARC		
1,3-butadiene		STEL	5 ppm	OSHA CARC		
1,3-butadiene		TWA	1 ppm	OSHA Z-1		
1,3-butadiene		STEL	5 ppm	OSHA Z-1		
benzene		TWA	0.5 ppm	ACGIH		
benzene		STEL	2.5 ppm	ACGIH		
benzene		PEL	1 ppm	OSHA CARC		
benzene		STEL	5 ppm	OSHA CARC		
benzene		TWA	10 ppm	OSHA Z-2		
benzene		CEIL	25 ppm	OSHA Z-2		
benzene		Peak	50 ppm (10 minutes)	OSHA Z-2		
Hydrogen sulfide	7783-06-4	TWA	5 ppm 7 mg/m3	2009/161/EU		
		nation: This value availabl	alue is for information when	nere there is no		
Hydrogen sulfide		STEL	10 ppm 14 mg/m3	2009/161/EU		
		Further information: This value is for information where there is no national limit value available.				
Hydrogen sulfide		STEL	5 ppm	ACGIH		
		Further information: Central Nervous System impairment, Upper Respiratory Tract irritation				
Hydrogen sulfide		CEIL	20 ppm	OSHA Z-2		
Hydrogen sulfide		Peak	50 ppm (10 minutes once only if no other measured expo- sure occurs)	OSHA Z-2		
Hydrogen sulfide		TWA	1 ppm	ACGIH		
Hydrogen sulfide		STEL	5 ppm	ACGIH		
Carbon dioxide	124-38-9	TWA	5,000 ppm	ACGIH		
Carbon dioxide		STEL	30,000 ppm	ACGIH		
Carbon dioxide		TWA	5,000 ppm 9,000 mg/m3	OSHA Z-1		

## **Biological occupational exposure limits**

Components	CAS-No.	Control parameters	Biological specimen	Sam- pling time	Permissible concentration	Basis
1,3-butadiene	106-99-0	1,2 Dihy- droxy-4-(N- acetylcyste- inyl)-butane	Urine	End of shift (As soon as possible	2.5 mg/l	ACGIH BEI

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		Mixture of N-1 and N- 2(hydroxybu tenyl)valine	Hemoglo- bin (Hb) adducts in blood	after exposure ceases) Not criti- cal	2.5 picomoles per gram Hemoglobin	ACGIH BEI
benzene	71-43-2	S- Phenylmer- capturic acid	Urine	End of shift (As soon as possible after exposure ceases)	25 µg/g creatinine	ACGIH BEI
		t,t-Muconic acid	Urine	End of shift (As soon as possible after exposure ceases)	500 μg/g creatinine	ACGIH BEI
1,3-butadiene	106-99-0	1,2 Dihy- droxy-4-(N- acetylcyste- inyl)-butane	Urine	End of shift (As soon as possible after exposure ceases)	2.5 mg/l	ACGIH BEI
		Mixture of N-1 and N- 2(hydroxybu tenyl)valine	Hemoglo- bin (Hb) adducts in blood	Not criti- cal	2.5 picomoles per gram Hemoglobin	ACGIH BEI
benzene	71-43-2	S- Phenylmer- capturic acid	Urine	End of shift (As soon as possible after exposure ceases)	25 μg/g creatinine	ACGIH BEI
		t,t-Muconic acid	Urine	End of shift (As soon as possible after exposure ceases)	500 μg/g creatinine	ACGIH BEI

# **Monitoring Methods**

Monitoring of the concentration of substances in the breathing zone of workers or in the general workplace may be required to confirm compliance with an OEL and adequacy of exposure controls. For some substances biological monitoring may also be appropriate.

Validated exposure measurement methods should be applied by a competent person and sam-

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ples analysed by an accredited laboratory.

Examples of sources of recommended exposure measurement methods are given below or contact the supplier. Further national methods may be available.

National Institute of Occupational Safety and Health (NIOSH), USA: Manual of Analytical Methods http://www.cdc.gov/niosh/

Occupational Safety and Health Administration (OSHA), USA: Sampling and Analytical Methods http://www.osha.gov/

Health and Safety Executive (HSE), UK: Methods for the Determination of Hazardous Substances http://www.hse.gov.uk/

Institut für Arbeitsschutz Deutschen Gesetzlichen Unfallversicherung (IFA), Germany http://www.dguv.de/inhalt/index.isp

L'Institut National de Recherche et de Securité, (INRS), France http://www.inrs.fr/accueil

## **Engineering measures**

The level of protection and types of controls necessary will vary depending upon potential exposure conditions. Select controls based on a risk assessment of local circumstances. Appropriate measures include:

Use sealed systems as far as possible.

Firewater monitors and deluge systems are recommended. Adequate explosion-proof ventilation to control airborne concentrations below the exposure guidelines/limits.

Local exhaust ventilation is recommended. Eye washes and showers for emergency use.

## General Information:

Consider technical advances and process upgrades (including automation) for the elimination of releases. Minimise exposure using measures such as closed systems, dedicated facilities and suitable general/local exhaust ventilation. Drain down systems and clear transfer lines prior to breaking containment. Clean/flush equipment, where possible, prior to maintenance. Where there is potential for exposure: restrict access to authorised persons; provide specific activity training to operators to minimise exposures; wear suitable gloves and coveralls to prevent skin contamination; wear respiratory protection when there is potential for inhalation; clear up spills immediately and dispose of wastes safely. Ensure safe systems of work or equivalent arrangements are in place to manage risks. Regularly inspect, test and maintain all control measures. Consider the need for risk based health surveillance.

Do not ingest. If swallowed then seek immediate medical assistance

## Personal protective equipment

Respiratory protection

If engineering controls do not maintain airborne concentrations to a level which is adequate to protect worker health, select respiratory protection equipment suitable for the specific conditions of use and meeting relevant legislation. Check with respiratory protective equipment suppliers.

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Where air-filtering respirators are unsuitable (e.g. airborne concentrations are high, risk of oxygen deficiency, confined space) use appropriate positive pressure breathing apparatus.

Where air-filtering respirators are suitable, select an appropriate combination of mask and filter.

All respiratory protection equipment and use must be in accordance with local regulations.

Respirator selection, use and maintenance should be in accordance with the requirements of the OSHA Respiratory Protection Standard, 29 CFR 1910.134.

Hand protection Remarks

Personal hygiene is a key element of effective hand care. Gloves must only be worn on clean hands. After using gloves, hands should be washed and dried thoroughly. Application of a non-perfumed moisturizer is recommended. Suitability and durability of a glove is dependent on usage, e.g. frequency and duration of contact, chemical resistance of glove material, dexterity. Always seek advice from glove suppliers. Contaminated gloves should be replaced. Where hand contact with the product may occur the use of gloves approved to relevant standards (e.g. Europe: EN374, US: F739) made from the following materials may provide suitable chemical protection. Neoprene rubber. Nitrile rubber. If contact with liquefied product is possible or anticipated, gloves should be thermally insulated to prevent cold burns. For continuous contact we recommend gloves with breakthrough time of more than 240 minutes with preference for > 480 minutes where suitable gloves can be identified. For shortterm/splash protection we recommend the same, but recognize that suitable gloves offering this level of protection may not be available and in this case a lower breakthrough time maybe acceptable so long as appropriate maintenance and replacement regimes are followed. Glove thickness is not a good predictor of glove resistance to a chemical as it is dependent on the exact composition of the glove material.

Eye protection : Wear safety glasses and face shield (preferably with a chin

guard) if splashes are likely to occur.

Skin and body protection : Chemical and cold resistant gloves/gauntlets, boots, and

apron.

Protective measures : Personal protective equipment (PPE) should meet recom-

mended national standards. Check with PPE suppliers.

Hygiene measures : Always observe good personal hygiene measures, such as

washing hands after handling the material and before eating, drinking, and/or smoking. Routinely wash work clothing and protective equipment to remove contaminants. Discard contaminated clothing and footwear that cannot be cleaned.

Practice good housekeeping.

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## **Environmental exposure controls**

General advice : Local guidelines on emission limits for volatile substances

must be observed for the discharge of exhaust air containing

vapour.

#### **SECTION 9. PHYSICAL AND CHEMICAL PROPERTIES**

Appearance : Gas.

Colour : Not applicable

Odour : Not applicable

Odour Threshold : Data not available

pH : Not applicable

Melting point/freezing point : Data not available

Initial boiling point and boiling

range

Data not available

Flash point : Data not available

Evaporation rate : Data not available

Flammability (solid, gas) : Extremely flammable.

Upper explosion limit / upper

flammability limit

ca. 15 %(V)

Lower explosion limit / Lower

flammability limit

ca. 1.8 %(V)

Vapour pressure : Not applicable

Relative vapour density : Data not available

Relative density : Data not available

Density : Data not available

Solubility(ies)

Water solubility : negligible

Solubility in other solvents : Data not available

Partition coefficient: n-

octanol/water

Data not available

Auto-ignition temperature :  $> 200 \, ^{\circ}\text{C} \, / > 392 \, ^{\circ}\text{F}$ 

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Decomposition temperature : Data not available

Viscosity

Viscosity, kinematic : Data not available

Explosive properties : Classification Code: NOT CLASS: Not classified

## **SECTION 10. STABILITY AND REACTIVITY**

Reactivity : No, product will not become self-reactive.

Chemical stability : Stable under normal conditions of use.

Possibility of hazardous reac-

tions

No. Hazardous, exothermical polymerization cannot occur.

Conditions to avoid : Heat, open flames, sparks and flammable atmospheres.

In certain circumstances product can ignite due to static elec-

tricity.

Incompatible materials : Oxidizing agents

Hazardous decomposition

products

Hazardous decomposition products are not expected to form

during normal storage.

### **SECTION 11. TOXICOLOGICAL INFORMATION**

Basis for assessment : Information given is based on product data, a knowledge of

the components and the toxicology of similar products.
Unless indicated otherwise, the data presented is representative of the product as a whole, rather than for individual com-

ponent(s).

## Information on likely routes of exposure

Inhalation is the primary route of exposure although exposure may occur through skin or eye contact.

## **Acute toxicity**

**Product:** 

Acute oral toxicity : Remarks: Not applicable

Acute inhalation toxicity : LC 50 (Rat): Exposure time: 4 h

Remarks: Harmful if inhaled. LC50 > 2500 - <= 20000 ppmV

Acute dermal toxicity : Remarks: Not applicable

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#### Skin corrosion/irritation

## **Product:**

Remarks: Not irritating to skin.

## Serious eye damage/eye irritation

## **Product:**

Remarks: Irritating to eyes. (Hydrogen Sulfide)

## Respiratory or skin sensitisation

## **Product:**

Remarks: Not a sensitiser.

Based on available data, the classification criteria are not met.

## Germ cell mutagenicity

## **Product:**

: Remarks: May cause heritable genetic damage, Mutagen classification based on Butadiene content at >= 0.1%., Contains benzene.

## Carcinogenicity

#### **Product:**

Remarks: Causes cancer in laboratory animals., Carcinogen classification based on Butadiene content at >= 0.1%.

Remarks: Contains Benzene, CAS # 71-43-2., Known human carcinogen.

Remarks: Contains Benzene, CAS # 71-43-2., May cause leukaemia (AML - acute myelogenous

leukaemia).

IARC Group 1: Carcinogenic to humans

1,3-butadiene 106-99-0

benzene 71-43-2

OSHA specifically regulated carcinogen

1,3-butadiene 106-99-0

benzene 71-43-2

NTP Known to be human carcinogen

1,3-butadiene 106-99-0

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benzene 71-43-2

## Reproductive toxicity

### **Product:**

Remarks: Causes foetotoxicity in animals at doses which are maternally toxic.

#### STOT - single exposure

#### **Product:**

Remarks: Contains hydrogen sulphide., Inhalation of vapours or mists may cause irritation to the respiratory system.

## STOT - repeated exposure

#### **Product:**

Remarks: May cause damage to organs or organ systems through prolonged or repeated exposure.. Blood

## **Aspiration toxicity**

#### **Product:**

Not an aspiration hazard.

#### **Further information**

## **Product:**

Remarks: H2S has a broad range of effects dependent on the airborne concentration and length of exposure: 0.02 ppm odour threshold, smell of rotten eggs; 10 ppm eye and respiratory tract irritation; 100 ppm coughing, headache, dizziness, nausea, eye irritation, loss of sense of smell in minutes; 200 ppm potential for pulmonary oedema after >20-30 minutes; 500 ppm loss of consciousness after short exposures, potential for respiratory arrest; >1000ppm immediate loss of consciousness, may lead rapidly to death, prompt cardiopulmonary resuscitation may be required. Do not depend on sense of smell for warning. H2S causes rapid olfactory fatigue (deadens sense of smell). There is no evidence that H2S will accumulate in the body tissue after repeated exposure., Rapid release of gases which are liquids under pressure may cause frost burns of exposed tissues (skin, eye) due to evaporative cooling., High gas concentrations will displace available air; unconsciousness and death may occur suddenly from lack of oxygen., Exposure to very high concentrations of similar materials has been associated with irregular heart rhythms and cardiac arrest.

Remarks: Classifications by other authorities under varying regulatory frameworks may exist.

## **SECTION 12. ECOLOGICAL INFORMATION**

Basis for assessment : Information given is based on product testing, and/or similar

products, and/or components.

According to OSHA Hazard Communication Standard, 29 CFR 1910.1200

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Unless indicated otherwise, the data presented is representative of the product as a whole, rather than for individual com-

ponent(s).

Physical properties indicate that petroleum gases will rapidly volatilise from the aquatic environment and that acute and

chronic effects would not be observed in practice.

**Ecotoxicity** 

**Product:** 

Toxicity to fish (Acute toxici-

ty)

Remarks: Practically non toxic:

LL/EL/IL50 > 100 mg/l

Toxicity to daphnia and other :

aquatic invertebrates (Acute

toxicity)

Remarks: Practically non toxic:

LL/EL/IL50 > 100 mg/l

Toxicity to algae (Acute tox-

icity)

Remarks: Practically non toxic:

LL/EL/IL50 > 100 mg/l

Toxicity to fish (Chronic tox-

icity)

Remarks: Data not available

Toxicity to daphnia and other : aquatic invertebrates (Chron-

ic toxicity)

Remarks: Data not available

Toxicity to microorganisms

(Acute toxicity)

Remarks: LL/EL/IL50 > 100 mg/l

Practically non toxic:

Based on available data, the classification criteria are not met.

Persistence and degradability

**Product:** 

Biodegradability : Remarks: Oxidises rapidly by photo-chemical reactions in air.

Readily biodegradable.

**Bioaccumulative potential** 

**Product:** 

Bioaccumulation : Remarks: Does not bioaccumulate significantly.

Mobility in soil

**Product:** 

Mobility : Remarks: Because of their extreme volatility, air is the only

environmental compartment that hydrocarbon gases will be

found.

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#### Other adverse effects

## **Product:**

Additional ecological infor-

mation

In view of the high rate of loss from solution, the product is

unlikely to pose a significant hazard to aquatic life.

#### **SECTION 13. DISPOSAL CONSIDERATIONS**

#### **Disposal methods**

Waste from residues : It is the responsibility of the waste generator to determine the

toxicity and physical properties of the material generated to determine the proper waste classification and disposal meth-

ods in compliance with applicable regulations.

Waste arising from a spillage or tank cleaning should be disposed of in accordance with prevailing regulations, preferably to a recognised collector or contractor. The competence of the collector or contractor should be established beforehand. Do not dispose into the environment, in drains or in water

courses

Given the nature and uses of this product, the need for disposal seldom arises. If necessary, dispose by controlled combustion in purpose-designed equipment. If this is not possible,

contact the supplier.

Contaminated packaging : Drain container thoroughly.

After draining, vent in a safe place away from sparks and fire.

Residues may cause an explosion hazard.

Do not pollute the soil, water or environment with the waste

container.

Return part-used or empty cylinders to the supplier. For tanks seek specialist advice from suppliers.

Dispose in accordance with prevailing regulations, preferably to a recognized collector or contractor. The competence of the collector or contractor should be established beforehand.

## **Local legislation**

Remarks : Disposal should be in accordance with applicable regional,

national, and local laws and regulations.

Local regulations may be more stringent than regional or na-

tional requirements and must be complied with.

#### **SECTION 14. TRANSPORT INFORMATION**

## **National Regulations**

**US Department of Transportation Classification (49 CFR Parts 171-180)** 

UN/ID/NA number : UN 3160

Proper shipping name : LIQUEFIED GAS, TOXIC, FLAMMABLE, N.O.S.

(Hydrogen sulphide)

Class : 2.3

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Subsidiary risk : 2.1

Packing group : Not Assigned Labels : 2.3 (2.1) ERG Code : 115 Marine pollutant : no

#### **International Regulations**

IATA-DGR

UN/ID No. : UN 3160 (Not permitted for transport)

Proper shipping name : LIQUEFIED GAS, TOXIC, FLAMMABLE, N.O.S.

(Hydrogen sulphide)

Class : 2.3

Packing group : Not Assigned

**IMDG-Code** 

UN number : UN 3160

Proper shipping name : LIQUEFIED GAS, TOXIC, FLAMMABLE, N.O.S.

(Hydrogen sulphide)

Class : 2.3 Subsidiary risk : 2.1

Packing group : Not Assigned Labels : 2.3 (2.1)

Marine pollutant : no

## Transport in bulk according to Annex II of MARPOL 73/78 and the IBC Code

Not applicable for product as supplied. MARPOL Annex 1 rules apply for bulk shipments by sea.

# Special precautions for user

Remarks : Special Precautions: Refer to Chapter 7, Handling & Storage,

for special precautions which a user needs to be aware of or

needs to comply with in connection with transport.

## **SECTION 15. REGULATORY INFORMATION**

# **EPCRA - Emergency Planning and Community Right-to-Know Act**

# **CERCLA Reportable Quantity**

Components	CAS-No.	Component RQ	Calculated product RQ	
		(lbs)	(lbs)	
1,3-butadiene	106-99-0	10	200	
benzene	71-43-2	10	10 (D018)	

<sup>\*:</sup> Vertex HSSE classifies this material as an "oil" under the CERCLA Petroleum Exclusion, therefore re-leases to the environment are not reportable under CERCLA., The components with RQs are given for information.

## SARA 304 Extremely Hazardous Substances Reportable Quantity

This material does not contain any components with a section 304 EHS RQ.

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## SARA 302 Extremely Hazardous Substances Threshold Planning Quantity

This material does not contain any components with a section 302 EHS TPQ.

SARA 311/312 Hazards : Flammable (gases, aerosols, liquids, or solids)

Gases under pressure

Carcinogenicity

Germ cell mutagenicity Reproductive toxicity

Acute toxicity (any route of exposure)

Specific target organ toxicity (single or repeated exposure)

SARA 313 : The following components are subject to reporting levels es-

tablished by SARA Title III, Section 313:

1,3-butadiene 106-99-0 >= 5 - < 10 %

benzene 71-43-2 >= 0.1 - < 1 %

#### Clean Water Act

The following Hazardous Chemicals are listed under the U.S. CleanWater Act, Section 311, Table 117.3:

 Hydrogen sulfide
 7783-06-4
 0.9999 %

 benzene
 71-43-2
 0.3 %

#### **US State Regulations**

## Pennsylvania Right To Know

 1,3-butadiene
 106-99-0

 Hydrogen sulfide
 7783-06-4

 benzene
 71-43-2

#### California Prop. 65

WARNING: This product can expose you to chemicals including 1,3-butadiene, benzene, which is/are known to the State of California to cause cancer and birth defects or other reproductive harm. For more information go to www.P65Warnings.ca.gov.

## California List of Hazardous Substances

1,3-butadiene 106-99-0

## California Regulated Carcinogens

 1,3-butadiene
 106-99-0

 benzene
 71-43-2

## Other regulations:

The regulatory information is not intended to be comprehensive. Other regulations may apply to this material.

#### **SECTION 16. OTHER INFORMATION**

#### **Further information**

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NFPA Rating (Health, Fire, Reac- 2, 4, 0

tivity)

## Full text of other abbreviations

2009/161/EU : 2009/161/EU

ACGIH : USA. ACGIH Threshold Limit Values (TLV)
ACGIH BEI : ACGIH - Biological Exposure Indices (BEI)

OSHA CARC : OSHA Specifically Regulated Chemicals/Carcinogens
OSHA Z-1 : USA. Occupational Exposure Limits (OSHA) - Table Z-1 Lim-

its for Air Contaminants

OSHA Z-2 : USA. Occupational Exposure Limits (OSHA) - Table Z-2

2009/161/EU / STEL : Short term exposure limit
2009/161/EU / TWA : Limit Value - eight hours
ACGIH / TWA : 8-hour, time-weighted average
ACGIH / STEL : Short-term exposure limit

ACGIH / STEL : Short-Term Exposure Limit (STEL)
OSHA CARC / PEL : Permissible exposure limit (PEL)

OSHA CARC / STEL : Excursion limit

OSHA Z-1 / TWA : 8-hour time weighted average
OSHA Z-1 / STEL : Short Term Exposure Limit
OSHA Z-2 / TWA : 8-hour time weighted average
OSHA Z-2 / CEIL : Acceptable ceiling concentration

OSHA Z-2 / Peak : Acceptable maximum peak above the acceptable ceiling con-

centration for an 8-hr shift

Abbreviations and Acronyms : The standard abbreviations and acronyms used in this docu-

ment can be looked up in reference literature (e.g. scientific

dictionaries) and/or websites.

ACGIH = American Conference of Governmental Industrial

Hygienists

ADR = European Agreement concerning the International

Carriage of Dangerous Goods by Road

AICS = Australian Inventory of Chemical Substances ASTM = American Society for Testing and Materials

BEL = Biological exposure limits

BTEX = Benzene, Toluene, Ethylbenzene, Xylenes

CAS = Chemical Abstracts Service

CEFIC = European Chemical Industry Council CLP = Classification Packaging and Labelling

COC = Cleveland Open-Cup

DIN = Deutsches Institut fur Normung
DMEL = Derived Minimal Effect Level
DNEL = Derived No Effect Level
DSL = Canada Domestic Substance List

EC = European Commission EC50 = Effective Concentration fifty

ECETOC = European Center on Ecotoxicology and Toxicolo-

gy Of Chemicals

ECHA = European Chemicals Agency

EINECS = The European Inventory of Existing Commercial

Chemical Substances

EL50 = Effective Loading fifty

ENCS = Japanese Existing and New Chemical Substances

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Inventory

EWC = European Waste Code

GHS = Globally Harmonised System of Classification and

Labelling of Chemicals

IARC = International Agency for Research on Cancer

IATA = International Air Transport Association

IC50 = Inhibitory Concentration fifty

IL50 = Inhibitory Level fifty

IMDG = International Maritime Dangerous Goods

INV = Chinese Chemicals Inventory

IP346 = Institute of Petroleum test method N° 346 for the determination of polycyclic aromatics DMSO-extractables

KECI = Korea Existing Chemicals Inventory

LC50 = Lethal Concentration fifty

LD50 = Lethal Dose fifty per cent.

LL/EL/IL = Lethal Loading/Effective Loading/Inhibitory loading

LL50 = Lethal Loading fifty

MARPOL = International Convention for the Prevention of

Pollution From Ships

NOEC/NOEL = No Observed Effect Concentration / No Observed Effect Level

served Effect Level

OE\_HPV = Occupational Exposure - High Production Volume

PBT = Persistent, Bioaccumulative and Toxic

PICCS = Philippine Inventory of Chemicals and Chemical

Substances

PNEC = Predicted No Effect Concentration

REACH = Registration Evaluation And Authorisation Of

Chemicals

RID = Regulations Relating to International Carriage of Dan-

gerous Goods by Rail

SKIN\_DES = Skin Designation

STEL = Short term exposure limit

TRA = Targeted Risk Assessment

TSCA = US Toxic Substances Control Act

TWA = Time-Weighted Average

vPvB = very Persistent and very Bioaccumulative

Due to the conversion of this product to GHS classification and labelling, there has been a significant change to the nature of the information presented in chapter 2.

A vertical bar (I) in the left margin indicates an amendment from the previous version.

Due to a change in detail in Section 15, this document has been released as a significant change.

Sources of key data used to compile the Safety Data Sheet

The quoted data are from, but not limited to, one or more sources of information (e.g. toxicological data from Vertex HSSE, material suppliers' data, CONCAWE, EU IUCLID date base, EC 1272 regulation, etc).

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The information provided in this Safety Data Sheet is correct to the best of our knowledge, information and belief at the date of its publication. The information given is designed only as a guidance for safe handling, use, processing, storage, transportation, disposal and release and is not to be considered a warranty or quality specification. The information relates only to the specific material designated and may not be valid for such material used in combination with any other materials or in any process, unless specified in the text.

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