According to OSHA Hazard Communication Standard, 29 CFR 1910.1200

PROPYLENE OXIDE Sustainable

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SECTION 1. IDENTIFICATION

Product name : PROPYLENE OXIDE Sustainable

Product code : U1151

CAS-No. : 75-56-9

Other means of identification : Epoxy propane, 1,2-, Methyl ethylene oxide, Methyl oxirane,

PO, Propylene epoxide, Propylene oxide, 1,2-

Manufacturer or supplier's details

Company : Shell Chemical LP

PO Box 576

HOUSTON TX 77001

USA

SDS Request : 1-800-240-6737

Customer Service : 1-855-697-4355

Emergency telephone number

Chemtrec Domestic (24 hr) : 1-800-424-9300

Chemtrec International (24

hr)

: 1-703-527-3887

Recommended use of the chemical and restrictions on use

Recommended use : Chemical intermediate.

Restrictions on use : Restricted to professional users., This product must not be

used in applications other than the above without first seeking

the advice of the supplier.

SECTION 2. HAZARDS IDENTIFICATION

GHS classification in accordance with the OSHA Hazard Communication Standard (29 CFR 1910.1200)

Flammable liquids : Category 1

Acute toxicity (Oral) : Category 4

Acute toxicity (Dermal) : Category 3

Acute toxicity (Inhalation) : Category 3

Eye irritation : Category 2A

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Specific target organ toxicity

- single exposure

Category 3 (Respiratory Tract)

Germ cell mutagenicity : Category 1B

Carcinogenicity : Category 1B

GHS label elements

Hazard pictograms







Signal word : Danger

Hazard statements : PHYSICAL HAZARDS:

H224 Extremely flammable liquid and vapour.

HEALTH HAZARDS:

H302 Harmful if swallowed. H311 Toxic in contact with skin.

H331 Toxic if inhaled.

H319 Causes serious eye irritation. H335 May cause respiratory irritation. H340 May cause genetic defects.

H350 May cause cancer. ENVIRONMENTAL HAZARDS:

Not classified as an environmental hazard under GHS criteria.

Precautionary statements : Prevention:

P201 Obtain special instructions before use.

P202 Do not handle until all safety precautions have been read

and understood.

P210 Keep away from heat/ sparks/ open flames/ hot surfaces.

No smoking.

P233 Keep container tightly closed.

P240 Ground/bond container and receiving equipment.

P241 Use explosion-proof electrical/ ventilating/ lighting equip-

ment.

P242 Use only non-sparking tools.

P243 Take precautionary measures against static discharge. P261 Avoid breathing dust/ fume/ gas/ mist/ vapours/ spray.

P264 Wash hands thoroughly after handling.

P270 Do not eat, drink or smoke when using this product.

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

P280 Wear protective gloves/ protective clothing/ eye protection/

face protection.

Response:

P301 + P310 IF SWALLOWED: Immediately call a POISON

CENTER or doctor/ physician.

P302 + P352 IF ON SKIN: Wash with plenty of soap and water. P303 + P361 + P353 IF ON SKIN (or hair): Remove/ Take off

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immediately all contaminated clothing. Rinse skin with water/ shower.

P304 + P340 IF INHALED: Remove victim to fresh air and keep

at rest in a position comfortable for breathing.

P305 + P351 + P338 IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.

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

P330 Rinse mouth.

P337 + P313 If eye irritation persists: Get medical advice/ attention

P361 + P364 Take off immediately all contaminated clothing and wash it before reuse.

P370 + P378 In case of fire: Use appropriate media to extin-

guish.

Storage:

P403 + P233 Store in a well-ventilated place. Keep container tightly closed.

P235 Keep cool. P405 Store locked up.

Disposal:

P501 Dispose of contents/ container to an approved waste disposal plant.

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.

Vapours may ignite and explode.

This material is a static accumulator.

Even with proper grounding and bonding, this material can still accumulate an electrostatic charge.

If sufficient charge is allowed to accumulate, electrostatic discharge and ignition of flammable airvapour mixtures can occur.

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)
propylene oxide	methyloxirane	75-56-9	<= 100

SECTION 4. FIRST-AID MEASURES

General advice : DO NOT DELAY.

Keep victim calm. Obtain medical treatment immediately.

If inhaled : Call emergency number for your location / facility.

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Remove to fresh air. Do not attempt to rescue the victim unless 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 percent modical facility.

the nearest medical facility.

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.

In case of eye contact : Immediately flush eye(s) with plenty of water.

Remove contact lenses, if present and easy to do. Continue

rinsing.

Transport to the nearest medical facility for additional treat-

ment.

If swallowed : If swallowed, do not induce vomiting: transport to nearest

medical facility for additional treatment. If vomiting occurs spontaneously, keep head below hips to prevent aspiration.

Rinse mouth.

Most important symptoms and effects, both acute and

delayed

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

and/or difficulty breathing.

Breathing of high vapour concentrations may cause central nervous system (CNS) depression resulting in dizziness, lightheadedness, headache, nausea and loss of coordination. Continued inhalation may result in unconsciousness and death.

Skin irritation signs and symptoms may include a burning sensation, redness, swelling, and/or blisters.

Eye irritation signs and symptoms may include a burning sen-

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

Ingestion may result in nausea, vomiting and/or diarrhoea. Peripheral nerve damage may be evidenced by impairment of motor function (incoordination, unsteady walk, or muscle weakness in the extremities, and/or loss of sensation in the

arms and legs).

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

: IMMEDIATE TREATMENT IS EXTREMELY IMPORTANT!

Artificial respiration may be required.

Call a doctor or poison control center for guidance.

Treat symptomatically.

SECTION 5. FIRE-FIGHTING MEASURES

According to OSHA Hazard Communication Standard, 29 CFR 1910.1200

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Suitable extinguishing media : Large fires should only be fought by properly trained fire fight-

ers.

Alcohol-resistant foam, water spray or fog. Dry chemical powder, carbon dioxide, sand or earth may be used for small fires

only.

Do not discharge extinguishing waters into the aquatic envi-

ronment.

Unsuitable extinguishing

media

Do not use water in a jet.

Specific hazards during fire-

fighting

Flammable vapours may be present even at temperatures

below the flash point.

Will float and can be reignited on surface water.

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

distant ignition is possible.

Carbon monoxide may be evolved if incomplete combustion

occurs.

Contents are under pressure and can explode when exposed

to heat or flames.

Specific extinguishing meth-

ods

Standard procedure for chemical fires.

Further information : Clear fire area of all non-emergency personnel.

All storage areas should be provided with adequate fire

fighting facilities.

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

Observe all relevant local and international regulations.

Risk of explosion. Inform the emergency services if liquid enters surface water drains.

Notify authorities if any exposure to the general public or the

environment occurs or is likely to occur.

Local authorities should be advised if significant spillages

cannot be contained.

Avoid contact with spilled or released material. For guidance on selection of personal protective equipment see Section 8 of

this Safety Data Sheet.

Isolate hazard area and deny entry to unnecessary or unpro-

tected personnel.

Stay upwind and keep out of low areas.

Extinguish any naked flames. Do not smoke. Remove ignition

sources. Avoid sparks.

Note - Since the danger of fire is so great, bunker gear worn

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over protective clothing is highly recommended.

Environmental precautions : Shut off leaks, if possible without personal risks.

Remove all possible sources of ignition in the surrounding

area.

Use appropriate containment to avoid environmental contamination. Prevent from spreading or entering drains, ditches or rivers by using sand, earth, or other appropriate barriers. Attempt to disperse the vapour or to direct its flow to a safe

location, for example by using fog sprays.

Methods and materials for containment and cleaning up

Large spillage:

Prevent from spreading by making a barrier with sand, earth

or other containment material.

Remove with explosion-proof vacuum trucks or pump to stor-

age/salvage vessels.

Test atmosphere for vapours to ensure safe working condi-

tions before other personnel are allowed into area.

Treat residues as for small spillage.

Small spillage:

Soak up residue with an absorbent such as clay, sand or other

suitable material and dispose of properly.

Allow to evaporate.

Retain washings as contaminated waste.

Note that aqueous solutions have a low flash point unless very

dilute.

Observe all relevant local and international regulations.

Additional advice

: For guidance on selection of personal protective equipment

see Section 8 of this Safety Data Sheet.

For guidance on disposal of spilled material see Section 13 of

this Safety Data Sheet.

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

Section 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.

Ensure that all local regulations regarding handling and stor-

age facilities are followed.

Advice on safe handling : Avoid exposure. Obtain special instructions before use.

Avoid inhaling vapour and/or mists.

Avoid contact with skin, eyes and clothing.

Monitor concentrations in air at regular intervals.

Extinguish any naked flames. Do not smoke. Remove ignition

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sources. Avoid sparks.

This product is intended for use in closed systems only.

Handling Temperature:

Ambient.

Use local exhaust ventilation if there is risk of inhalation of vapours, mists or aerosols.

Bulk storage tanks should be diked (bunded).

Properly dispose of any contaminated rags or cleaning materials in order to prevent fires.

Even with proper grounding and bonding, this material can still accumulate an electrostatic charge.

If sufficient charge is allowed to accumulate, electrostatic discharge and ignition of flammable air-vapour mixtures can occur.

Be aware of handling operations that may give rise to additional hazards that result from the accumulation of static charges.

These include but are not limited to pumping (especially turbulent flow), mixing, filtering, splash filling, cleaning and filling of tanks and containers, sampling, switch loading, gauging, vacuum truck operations, and mechanical movements.

These activities may lead to static discharge e.g. spark formation.

Restrict line velocity during pumping in order to avoid generation of electrostatic discharge (≤ 1 m/s until fill pipe submerged to twice its diameter, then ≤ 7 m/s). Avoid splash filling. Do NOT use compressed air for filling, discharging, or handling operations.

Avoidance of contact

Clay-based absorbents.

Bases, ammonia, primary and secondary amines, water and acids

Heavy metals, alkali metals, alkali metal hydroxides, anhydrous chlorides of aluminium. iron, tin, copper and its alloys.

Strong oxidising agents.

Product Transfer

If positive displacement pumps are used, these must be fitted with a non-integral pressure relief valve. Lines should be purged with nitrogen before and after product transfer. Refer to supplier for further product transfer instructions if required. Refer to guidance under Handling section.

Conditions for safe storage

Refer to section 15 for any additional specific legislation covering the packaging and storage of this product.

Further information on storage stability

Keep away from aerosols, flammables, oxidizing agents, corrosives and from other flammable products which are not harmful or toxic to man or to the environment.

A reliable fixed sprinkler/deluge system should be installed.

Tanks must be clean, dry and rust-free.

Prevent ingress of water.

Must be stored in a diked (bunded) well- ventilated area, away from sunlight, ignition sources and other sources of heat. Tanks should be fitted with a vapour recovery system. Vapours from tanks should not be released to atmosphere.

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Breathing losses during storage should be controlled by a suitable vapour treatment system.

Tanks must be specifically designed for use with this product. Cleaning, inspection and maintenance of storage tanks is a specialist operation, which requires the implementation of strict procedures and precautions.

These include issuing of work permits, gas-freeing of tanks, using a manned harness and lifelines and wearing air-supplied breathing apparatus.

Storage Temperature:

30 °C / 86 °F maximum.

Use lowest practicable storage temperatures and avoid through-draughts of air to minimise risk of generating a flammable condition in the tank space.

Electrostatic charges will be generated during pumping. Electrostatic discharge may cause fire. Ensure electrical continuity by bonding and grounding (earthing) all equipment to reduce the risk.

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

Packaging material : Suitable material: Stainless steel., Mild steel.

Unsuitable material: Plastics, Aluminum

Specific use(s) : Not applicable

Ensure that all local regulations regarding handling and stor-

age facilities are followed.

See additional references that provide safe handling practices: 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 exposure)	Control parameters / Permissible concentration	Basis
propylene oxide	75-56-9	TWA	1 ppm 2.4 mg/m3	Shell Internal Standard (SIS) for 8-12 hour TWA.
propylene oxide		TWA	2 ppm	ACGIH
propylene oxide		TWA	100 ppm 240 mg/m3	OSHA Z-1

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Biological occupational exposure limits

No biological limit allocated.

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 samples 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.jsp

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:

Adequate explosion-proof ventilation to control airborne concentrations.

Local exhaust ventilation is recommended.

Firewater monitors and deluge systems are recommended. Items that cannot be decontaminated should be destroyed (see Chapter 13).

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.

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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. 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.

If air-filtering respirators are suitable for conditions of use:

Select a filter suitable for organic gases and vapours [boiling point <65 °C (149 °F)]

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

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. Longer term protection: Butyl rubber. Incidental contact/Splash protection: Nitrile rubber gloves. Silver Shield. 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 short-term/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. Glove thickness should be typically greater than 0.35 mm depending on the glove make and model. 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. 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 nonperfumed moisturizer is recommended.

Eye protection : Wear goggles for use against liquids and gas, combined with

face shield.

Skin and body protection : Wear antistatic and flame-retardant clothing.

Wear chemical resistant gloves/gauntlets and boots. Where

risk of splashing, also wear an apron.

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

mended national standards. Check with PPE suppliers.

Thermal hazards : When handling cold material that can cause frost burns, wear

cryogenic gloves, safety hat and visor, cold resistant overalls (with cuffs over gloves and legs over boots) and heavy duty

boots e.g. leather for cold resistance.

Hygiene measures : Wash hands before eating, drinking, smoking and using the

toilet.

Launder contaminated clothing before re-use.

Environmental exposure controls

General advice : Local guidelines on emission limits for volatile substances

must be observed for the discharge of exhaust air containing

vapour.

Minimise release to the environment. An environmental assessment must be made to ensure compliance with local envi-

ronmental legislation.

Information on accidental release measures are to be found in

section 6.

SECTION 9. PHYSICAL AND CHEMICAL PROPERTIES

Appearance : Oily liquid.

Colour : Colourless to yellowish

Odour : Ethereal

Odour Threshold : 35 ppm

pH : Data not available

Melting / freezing point : -112 °C / -170 °F

Boiling point/boiling range : 35 °C / 95 °F

Flash point : $-37 \,^{\circ}\text{C} / -35 \,^{\circ}\text{F}$

Method: Tag Closed Cup (ASTM D56)

Evaporation rate : ca. 12

Method: ASTM D 3539, nBuAc=1

Flammability

Flammability (solid, gas) : Not applicable

Lower explosion limit and upper explosion limit / flammability limit

Upper explosion limit / up- : 37.0 %(V)

per flammability limit

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Lower explosion limit /

Lower flammability limit

: 1.7 %(V)

Vapour pressure : 25.1 kPa (0 °C / 32 °F)

Relative vapour density : 3.6

Relative density : 0.824 (3.89 °C / 39.00 °F)

Method: ASTM D4052

Density : 830 kg/m3 (20 °C / 68 °F)

Method: ASTM D4052

Solubility(ies)

Water solubility : 405 kg/m3 (20 °C / 68 °F

)

Solubility in other solvents : Data not available

Partition coefficient: n-

octanol/water

log Pow: 0.03

Auto-ignition temperature : 490 °C / 914 °F

Decomposition temperature : Data not available

Viscosity

Viscosity, dynamic : 0.58 mPa.s (20 °C / 68 °F)

Method: ASTM D445

Viscosity, kinematic : 0.374 mm2/s (20 °C / 68 °F)

Method: ASTM D445

0.447 mm2/s (0 °C / 32 °F)

Method: ASTM D445

Explosive properties : Not classified

Oxidizing properties : Data not available

Surface tension : 71.5 mN/m, 15 °C / 59 °F

Conductivity: < 100 pS/m

The conductivity of this material makes it a static accumulator., A liquid is typically considered nonconductive if its con-

ductivity is below 100 pS/m and is considered semi-

conductive if its conductivity is below 10,000 pS/m., Whether a liquid is nonconductive or semi-conductive, the precautions are the same., A number of factors, for example liquid tem-

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perature, presence of contaminants, and anti-static additives

can greatly influence the conductivity of a liquid

Molecular weight 58.01 g/mol

Particle size Data not available

SECTION 10. STABILITY AND REACTIVITY

Reactivity Material will polymerise at elevated temperatures 122 °F (50

°C) or if contaminated with water.

Chemical stability No hazardous reaction is expected when handled and stored

according to provisions

Possibility of hazardous reac-

tions

Reacts violently with strong oxidising agents.

Reacts with strong acids.

Conditions to avoid Heat, flames, and sparks.

> Prevent vapour accumulation. Temperatures above 30 °C / 86 °F.

Incompatible materials Clay-based absorbents.

Bases, ammonia, primary and secondary amines, water and

acids.

Heavy metals, alkali metals, alkali metal hydroxides, anhydrous chlorides of aluminium. iron, tin, copper and its alloys.

Strong oxidising agents.

Hazardous decomposition

products

Thermal decomposition is highly dependent on conditions. A complex mixture of airborne solids, liquids and gases includ-

ing carbon monoxide, carbon dioxide, sulphur oxides and unidentified organic compounds will be evolved when this material undergoes combustion or thermal or oxidative degra-

dation.

Unknown toxic products may be formed.

SECTION 11. TOXICOLOGICAL INFORMATION

Basis for assessment : Information given is based on product testing.

> 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.

Acute toxicity

Components:

propylene oxide:

: LD 50 (Rat, male and female): > 300 - <= 2000 mg/kg Acute oral toxicity

Method: Test(s) equivalent or similar to OECD Test Guideline

401

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Remarks: Harmful if swallowed.

Acute inhalation toxicity : LC 50 (Rat, male and female): > 2 -<= 10 mg/l

Exposure time: 4 h
Test atmosphere: vapour

Method: Test(s) equivalent or similar to OECD Test Guideline

403

Remarks: Toxic if inhaled.

High concentrations may cause central nervous system depression resulting in headaches, dizziness and nausea.

Acute dermal toxicity : LD 50 (Rabbit): > 200 - <= 1000 mg/kg

Method: Literature data

Remarks: Toxic in contact with skin.

The current CLP-based labelling of acute dermal toxicity (Category 3; H311) is inaccurate due to a mathematical error having occurred when conducting a unit conversion for the referenced dermal LD50 value of 1.5 mL/kg bw to 950 mg/kg bw. The referenced dermal LD50 is correctly converted to 1,245 mg/kg bw (Category 4; H312) based on propylene oxide rela-

tive density (0.830 at 20°C).

Skin corrosion/irritation

Components:

propylene oxide: Species: Rabbit

Method: OECD Test Guideline 404

Remarks: Not irritating to skin., Based on available data, the classification criteria are not met.

Serious eye damage/eye irritation

Components:

propylene oxide: Species: Rabbit

Method: Literature data

Remarks: Causes serious eye irritation.

Respiratory or skin sensitisation

Components:

propylene oxide:

Species: Guinea pig

Method: Acceptable non-standard method.

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

Germ cell mutagenicity

Components:

propylene oxide:

Genotoxicity in vitro : Method: OECD Test Guideline 471

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Remarks: May cause genetic defects.

: Method: OECD Test Guideline 473 Remarks: May cause genetic defects.

: Method: OECD Test Guideline 476 Remarks: May cause genetic defects.

Genotoxicity in vivo : Remarks: May cause genetic defects.

Carcinogenicity

Components:

propylene oxide:

Species: Mouse, (male and female) Application Route: Inhalation

Method: Test(s) equivalent or similar to OECD Test Guideline 451

Remarks: May cause cancer.

IARC Group 2B: Possibly carcinogenic to humans

propylene oxide 75-56-9

OSHA No component of this product present at levels greater than or

equal to 0.1% is on OSHA's list of regulated carcinogens.

NTP Reasonably anticipated to be a human carcinogen

propylene oxide 75-56-9

Reproductive toxicity

Components:

propylene oxide:

Effects on fertility :

Species: Rat

Sex: male and female Application Route: Inhalation

Method: Equivalent or similar to OECD Test Guideline 416 Remarks: Based on available data, the classification criteria

are not met.

Effects on foetal develop-

ment

Species: Rat, female

Application Route: Inhalation Method: Other guideline method.

Remarks: Based on available data, the classification criteria

are not met.

According to OSHA Hazard Communication Standard, 29 CFR 1910.1200

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STOT - single exposure

Components:

propylene oxide:

Exposure routes: Inhalation
Target Organs: Respiratory Tract

Remarks: May cause respiratory irritation.

STOT - repeated exposure

Components:

propylene oxide:

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

Repeated dose toxicity

Components:

propylene oxide:

Species: Rat, male and female Application Route: Inhalation Test atmosphere: vapour

Method: OECD Test Guideline 453

Target Organs: No specific target organs noted

Aspiration toxicity

Components:

propylene oxide:

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

Further information

Components:

propylene oxide:

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.

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

ponent(s).

Ecotoxicity

Components:

propylene oxide:

Toxicity to fish (Acute toxici-

: LC50 (Oncorhynchus mykiss (rainbow trout)): 52 mg/l

ty) Exposure time: 96 h

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Method: Test(s) equivalent or similar to OECD Guideline 203

Remarks: Harmful

LL/EL/IL50 >10 <= 100 mg/l

Toxicity to daphnia and other : aquatic invertebrates (Acute

toxicity)

EC50 (Daphnia magna (Water flea)): 350 mg/l

Exposure time: 48 h

Method: Test(s) equivalent or similar to OECD Guideline 202

Remarks: Practically non toxic:

LL/EL/IL50 > 100 mg/l

Toxicity to algae (Acute tox-

icity)

EC50 (Pseudokirchneriella subcapitata (algae)): 240 mg/l

Exposure time: 96 h

Method: Test(s) equivalent or similar to OECD Test Guideline

201

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: Practically non toxic:

LL/EL/IL50 > 100 mg/l

Persistence and degradability

Components:

propylene oxide:

Biodegradability : Biodegradation: 89 %

Exposure time: 28 d

Method: OECD Test Guideline 301C Remarks: Readily biodegradable.

Bioaccumulative potential

Components:

propylene oxide:

Bioaccumulation : Remarks: Does not bioaccumulate significantly.

Mobility in soil

Components:

propylene oxide:

Mobility : Remarks: Dissolves in water.

If the product enters soil, one or more constituents will or may

be mobile and may contaminate groundwater.

According to OSHA Hazard Communication Standard, 29 CFR 1910.1200

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Other adverse effects

no data available

SECTION 13. DISPOSAL CONSIDERATIONS

Disposal methods

Waste from residues : Recover or recycle if possible.

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 methods are the controlled to the controlled t

ods in compliance with applicable regulations.

Do not dispose into the environment, in drains or in water

courses.

Waste product should not be allowed to contaminate soil or

water.

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.

Waste, spills or used product is dangerous waste.

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 puncture.

cut or weld uncleaned drums.

Send to drum recoverer or metal reclaimer.

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

national, and local laws and regulations.

SECTION 14. TRANSPORT INFORMATION

National Regulations

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

UN/ID/NA number : UN 1280

Proper shipping name : PROPYLENE OXIDE

Class : 3
Packing group : 1
Labels : 3

Reportable quantity Propylene oxide

(100 lb)

ERG Code : 127P Marine pollutant : no

International Regulations

According to OSHA Hazard Communication Standard, 29 CFR 1910.1200

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IATA-DGR

UN/ID No. : UN 1280

Proper shipping name : PROPYLENE OXIDE

Class : 3
Packing group : 1
Labels : 3

IMDG-Code

UN number : UN 1280

Proper shipping name : PROPYLENE OXIDE

Class : 3
Packing group : I
Labels : 3
Marine pollutant : no

Maritime transport in bulk according to IMO instruments

Pollution category : Y Ship type : 2

Product name : Propylene oxide

Special precautions for user

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

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

needs to comply with in connection with transport.

Additional Information: This product may be transported under nitrogen blanketing.

Nitrogen is an odourless and invisible gas. Exposure to nitrogen enriched atmospheres displaces available oxygen which may cause asphyxiation or death. Personnel must observe strict safety precautions when involved with a confined space

entry.

Transport in bulk according to Annex II of Marpol and the IBC

Code

SECTION 15. REGULATORY INFORMATION

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

CERCLA Reportable Quantity

Components	CAS-No.	lo. Component RQ Calculated	
		(lbs)	(lbs)
propylene oxide	75-56-9	100	100

*: SARA 304 Extremely Hazardous Substances Reportable Quantity

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

SARA 302 Extremely Hazardous Substances Threshold Planning Quantity

Components	CAS-No.	Component TPQ (lbs)
propylene oxide	75-56-9	10000

According to OSHA Hazard Communication Standard, 29 CFR 1910.1200

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SARA 302 Extremely Hazardous Substances Threshold Planning Quantity

propylene oxide 75-56-9 >= 90 - <= 100 %

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

Serious eye damage or eye irritation

Specific target organ toxicity (single or repeated exposure)

Germ cell mutagenicity

Carcinogenicity

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

tablished by SARA Title III, Section 313:

propylene oxide 75-56-9 >= 90 - <= 100 %

Clean Water Act

This product does not contain any Hazardous Chemicals listed under the U.S. CleanWater Act, Section 311, Table 117.3.

US State Regulations

Pennsylvania Right To Know

propylene oxide 75-56-9

California Prop. 65

WARNING: This product can expose you to chemicals including propylene oxide, which is/are known to the State of California to cause cancer. For more information go to www.P65Warnings.ca.gov.

California List of Hazardous Substances

propylene oxide 75-56-9

Other regulations:

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

The components of this product are reported in the following inventories:

AIIC : Listed

DSL : Listed

IECSC : Listed

ENCS : Listed

KECI : Listed

NZIoC : Listed

PICCS : Listed

TSCA : Listed

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TCSI : Listed

SECTION 16. OTHER INFORMATION

Further information

NFPA Rating (Health, Fire, Reac- 3, 4, 2

tivity)

Full text of other abbreviations

ACGIH : USA. ACGIH Threshold Limit Values (TLV)

OSHA Z-1 : USA. Occupational Exposure Limits (OSHA) - Table Z-1 Lim-

its for Air Contaminants

ACGIH / TWA : 8-hour, time-weighted average OSHA Z-1 / TWA : 8-hour time weighted average

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

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

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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 Ob-

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

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

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 Shell Health Services, material suppliers' data, CONCAWE, EU

IUCLID date base, EC 1272 regulation, etc).

Revision Date : 12/28/2023

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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