TRP Tyre Sheen

Paccar Australia Pty Ltd

Chemwatch Hazard Alert Code: 3

Chemwatch: 24-3441TRP Safety Data Sheet according to WHS and ADG requirements

Issue Date: 12/03/2015 Version No: 3.1.1.1

Print Date: 23/03/2016

Initial Date: Not Available S.GHS.AUS.EN

SECTION 1 IDENTIFICATION OF THE SUBSTANCE / MIXTURE AND OF THE COMPANY / UNDERTAKING

Product name TRP Tyre Sheen Synonyms Not Available Proper shipping name Other means of identification Identification Relevant identified uses of the substance or mixture and uses advised against The use of a quantity of material in an unventilated or confined space may result in increased exposure and an irritating atmosphere developing. Before starting consider control of exposure by mechanical ventilation. Use according to manufacturer's directions. Automotive tyre rejuvenator. Details of the supplier of the safety data sheet Registered company name Address 20 Canterbury Rd, Bayswater, VIC 3152 Australia Telephone +61 3 9721 1500 Fax N/A Website www.paccar.com.au Email Not Available Emergency telephone number Association / Organisation Final Not Available Not Available	Product Identifier	
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SECTION 2 HAZARDS IDENTIFICATION	SECTION 2 HAZARDS ID	ENTIFICATION

Classification of the substance or mixture

HAZARDOUS CHEMICAL. DANGEROUS GOODS. According to the WHS Regulations and the ADG Code.

CHEMWATCH HAZARD RATINGS

	MinMax_	
Flammability	3	
Toxicity	2	0 = Minimum
Body Contact	1	1 = Low 2 = Moderate
Reactivity	1	3 = High
Chronic	2	4 = Extreme

Poisons Schedule	S5
[1] Classification	Flammable Liquid Category 2, Specific target organ toxicity - single exposure Category 3 (narcotic effects), Specific target organ toxicity - repeated exposure Category 2, Aspiration Hazard Category 1
Legend:	1. Classified by Chemwatch; 2. Classification drawn from HSIS; 3. Classification drawn from EC Directive 1272/2008 - Annex VI

Label elements

GHS label elements







SIGNAL WORD

DANGER

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Hazard statement(s)	
H225	Highly flammable liquid and vapour.
H336	May cause drowsiness or dizziness.
H373	May cause damage to organs.
H304	May be fatal if swallowed and enters airways.
AUH066	Repeated exposure may cause skin dryness and cracking
Precautionary statement(s) P	Prevention
P210	Keep away from heat/sparks/open flames/hot surfaces No smoking.
P260	Do not breathe dust/fume/gas/mist/vapours/spray.
P271	Use only outdoors or in a well-ventilated area.
P240	Ground/bond container and receiving equipment.
P241	Use explosion-proof electrical/ventilating/lighting/intrinsically safe equipment.
P242	Use only non-sparking tools.
P243	Take precautionary measures against static discharge.
Precautionary statement(s) R	Response
P301+P310	IF SWALLOWED: Immediately call a POISON CENTER or doctor/physician.
P331	Do NOT induce vomiting.
P370+P378	In case of fire: Use alcohol resistant foam or normal protein foam for extinction.
P312	Call a POISON CENTER or doctor/physician if you feel unwell.
P303+P361+P353	IF ON SKIN (or hair): Remove/Take off 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.
Precautionary statement(s) S	Storage
P403+P235	Store in a well-ventilated place. Keep cool.
P405	Store locked up.
P403+P233	Store in a well-ventilated place. Keep container tightly closed.
Precautionary statement(s) D	Disposal
P501	Dispose of contents/container in accordance with local regulations.
SECTION 3 COMPOSITIO	N / INFORMATION ON INGREDIENTS

SECTION 3 COMPOSITION / INFORMATION ON INGREDIENTS

Substances

See section below for composition of Mixtures

Mixtures

CAS No	%[weight]	Name
64742-49-0.	>60	naphtha petroleum, light, hydrotreated
63148-62-9	10-30	polydimethylsiloxane

SECTION 4 FIRST AID MEASURES

Description	OI III	st alu	measu	res

If this product comes in contact with the eyes:

Eye Contact

▶ Wash out immediately with fresh running water. ▶ Ensure complete irrigation of the eye by keeping eyelids apart and away from eye and moving the eyelids by occasionally lifting the upper and lower lids. ▶ Seek medical attention without delay; if pain persists or recurs seek medical attention.

▶ Removal of contact lenses after an eye injury should only be undertaken by skilled personnel.

Skin Contact

- ▶ Immediately remove all contaminated clothing, including footwear.
- ▶ Flush skin and hair with running water (and soap if available).
- ▶ Seek medical attention in event of irritation.

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Inhalation	 ▶ If fumes or combustion products are inhaled remove from contaminated area. ▶ Lay patient down. Keep warm and rested. ▶ Prostheses such as false teeth, which may block airway, should be removed, where possible, prior to initiating first aid procedures. ▶ Apply artificial respiration if not breathing, preferably with a demand valve resuscitator, bag-valve mask device, or pocket mask as trained. ▶ Perform CPR if necessary. ▶ Transport to hospital, or doctor.
Ingestion	 ▶ If swallowed do NOT induce vomiting. ▶ If vomiting occurs, lean patient forward or place on left side (head-down position, if possible) to maintain open airway and prevent aspiration ▶ Observe the patient carefully. ▶ Never give liquid to a person showing signs of being sleepy or with reduced awareness; i.e. becoming unconscious. ▶ Give water to rinse out mouth, then provide liquid slowly and as much as casualty can comfortably drink. ▶ Seek medical advice. ▶ Avoid giving milk or oils. ▶ Avoid giving alcohol. ▶ If spontaneous vomiting appears imminent or occurs, hold patient's head down, lower than their hips to help avoid possible aspiration of vomitus.

Indication of any immediate medical attention and special treatment needed

For acute or short term repeated exposures to petroleum distillates or related hydrocarbons:

- Primary threat to life, from pure petroleum distillate ingestion and/or inhalation, is respiratory failure.
- Patients should be quickly evaluated for signs of respiratory distress (e.g. cyanosis, tachypnoea, intercostal retraction, obtundation) and given oxygen. Patients with inadequate tidal volumes or poor arterial blood gases (pO2 50 mm Hg) should be intubated.
- Arrhythmias complicate some hydrocarbon ingestion and/or inhalation and electrocardiographic evidence of myocardial injury has been reported; intravenous lines and cardiac monitors should be established in obviously symptomatic patients. The lungs excrete inhaled solvents, so that hyperventilation improves clearance.
- A chest x-ray should be taken immediately after stabilisation of breathing and circulation to document aspiration and detect the presence of pneumothorax.
- ▶ Epinephrine (adrenalin) is not recommended for treatment of bronchospasm because of potential myocardial sensitisation to catecholamines. Inhaled cardioselective bronchodilators (e.g. Alupent, Salbutamol) are the preferred agents, with aminophylline a second choice.
- Lavage is indicated in patients who require decontamination; ensure use of cuffed endotracheal tube in adult patients. [Ellenhorn and Barceloux: Medical Toxicology]

Any material aspirated during vomiting may produce lung injury. Therefore emesis should not be induced mechanically or pharmacologically. Mechanical means should be used if it is considered necessary to evacuate the stomach contents; these include gastric lavage after endotracheal intubation. If spontaneous vomiting has occurred after ingestion, the patient should be monitored for difficult breathing, as adverse effects of aspiration into the lungs may be delayed up to 48 hours. Treat symptomatically.

SECTION 5 FIREFIGHTING MEASURES

Extinguishing media

- Foam.
- Dry chemical powder.
- BCF (where regulations permit).
- Carbon dioxide.
- Water spray or fog Large fires only.

Fire Incompatibility	Avoid contamination with oxidising agents i.e. nitrates, oxidising acids, chlorine bleaches, pool chlorine etc. as ignition may result
vice for firefighters	
	▶ Alert Fire Brigade and tell them location and nature of hazard.
	▶ May be violently or explosively reactive.
	▶ Wear breathing apparatus plus protective gloves in the event of a fire.
Fire Fighting	▶ Prevent, by any means available, spillage from entering drains or water course. ▶
	Consider evacuation (or protect in place).
	▶ Fight fire from a safe distance, with adequate cover.
	▶ If safe, switch off electrical equipment until vapour fire hazard removed.
	▶ High temperature decomposition products include silicon dioxide, small amounts of formaldehyde, formic acid, acetic acid and traces of silico
	polymers. ▶ These gases may ignite and, depending on circumstances, may cause the resin/polymer to ignite.
	▶ An outer skin of silica may also form. Extinguishing of fire, beneath the skin, may be difficult.
	▶ Liquid and vapour are highly flammable.
Fire/Explosion	▶ Severe fire hazard when exposed to heat, flame and/or oxidisers.
Hazard	▶ Vapour may travel a considerable distance to source of ignition.
	▶ Heating may cause expansion or decomposition leading to violent rupture of
	containers. ▶ On combustion, may emit toxic fumes of carbon monoxide (CO).
	Combustion products include:, carbon dioxide (CO2), silicon dioxide (SiO2), other pyrolysis products typical of burning organic material Contain
	low boiling substance: Closed containers may rupture due to pressure buildup under fire conditions.

SECTION 6 ACCIDENTAL RELEASE MEASURES

Personal precautions, protective equipment and emergency procedures ▶ Remove all ignition sources. ▶ Clean up all spills immediately. Avoid breathing vapours and contact with skin and eves. **Minor Spills** ▶ Control personal contact with the substance, by using protective equipment. ▶ Contain and absorb small quantities with vermiculite or other absorbent material. Wipe up. ▶ Collect residues in a flammable waste container.

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- ▶ Silicone fluids, even in small quantities, may present a slip hazard.
- \blacktriangleright It may be necessary to rope off area and place warning signs around perimeter.
- ▶ Clean up area from spill, with suitable absorbant, as soon as practically possible.
- ▶ Final cleaning may require use of steam, solvents or detergents.
- ▶ Clear area of personnel and move upwind. **Major Spills**
 - ▶ Alert Fire Brigade and tell them location and nature of hazard.
 - ▶ May be violently or explosively reactive.
 - ▶ Wear breathing apparatus plus protective gloves.
 - ▶ Prevent, by any means available, spillage from entering drains or water course.
 - ▶ Consider evacuation (or protect in place).
 - ▶ No smoking, naked lights or ignition sources.

Personal Protective Equipment advice is contained in Section 8 of the SDS.

SECTION 7 HANDLING AND STORAGE

	▶ Containers, even those that have been emptied, may contain explosive vapours.
Cofe hor -III	▶ Do NOT cut, drill, grind, weld or perform similar operations on or near containers.
Safe handling	Contains low boiling substance:
	Storage in sealed containers may result in pressure buildup causing violent rupture of containers not rated appropriately.
	▶ Check for bulging containers.
	▶ Vent periodically
	Always release caps or seals slowly to ensure slow dissipation of
	vapours ▶ DO NOT allow clothing wet with material to stay in contact
	with skin ▶ Avoid all personal contact, including inhalation.
	▶ Wear protective clothing when risk of exposure occurs. ▶
	Use in a well-ventilated area.
	Prevent concentration in hollows and sumps.
	DO NOT enter confined spaces until atmosphere has been checked.
	Avoid smoking, naked lights, heat or ignition sources.
	▶ When handling, DO NOT eat, drink or smoke.
	▶ Store in original containers in approved flame-proof area.
	▶ No smoking, naked lights, heat or ignition sources.
	▶ DO NOT store in pits, depressions, basements or areas where vapours may be trapped.
Other information	▶ Keep containers securely sealed.
	▶ Store away from incompatible materials in a cool, dry well ventilated area.
	▶ Protect containers against physical damage and check regularly for leaks.
	Observe manufacturer's storage and handling recommendations contained within this SDS.
onditions for safe storage,	including any incompatibilities
	▶ Packing as supplied by manufacturer.
	▶ Plastic containers may only be used if approved for flammable liquid. ▶
	Check that containers are clearly labelled and free from leaks.
	▶ For low viscosity materials (i): Drums and jerry cans must be of the non-removable head type. (ii): Where a can is to be used as an inner
	package, the can must have a screwed enclosure.
	▶ For materials with a viscosity of at least 2680 cSt. (23 deg. C)
Suitable container	▶ For manufactured product having a viscosity of at least 250 cSt. (23 deg. C)
	▶ Manufactured product that requires stirring before use and having a viscosity of at least 20 cSt (25 deg. C): (i) Removable head packaging; (ii)
	Cans with friction closures and (iii) low pressure tubes and cartridges may be used.
	▶Where combination packages are used, and the inner packages are of glass, there must be sufficient inert cushioning material in contact with
	inner and outer packages
	▶ In addition, where inner packagings are glass and contain liquids of packing group I there must be sufficient inert absorbent to absorb any
	spillage, unless the outer packaging is a close fitting moulded plastic box and the substances are not incompatible with the plastic.
	Traces of benzene, a carcinogen, may form when silicones are heated in air above 230 degrees C. Concentrated acids and bases cause degradati
Storage incompatibility	
	▶ Avoid reaction with oxidising agents

SECTION 8 EXPOSURE CONTROLS / PERSONAL PROTECTION

Control parameters

OCCUPATIONAL EXPOSURE LIMITS (OEL)

INGREDIENT DATA

Not Available

EMERGENCY LIMITS					
Ingredient	Material name		TEEL-1	TEEL-2	TEEL-3
polydimethylsiloxane	Dimethyl siloxane; (Dimethylpolysiloxane; Syltherm XLT; Syltherm 800; Silicone 360) 1.5 mg/m			16 mg/m3	990 mg/m3
Ingredient	Original IDLH	Revised IDLH			
naphtha petroleum, light, hydrotreated	Not Available	Not Available			

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polydimethylsiloxane Not Available Not Available **Exposure controls** CARE: Use of a quantity of this material in confined space or poorly ventilated area, where rapid build up of concentrated atmosphere may occur, could require increased ventilation and/or protective gear Engineering controls are used to remove a hazard or place a barrier between the worker and the hazard. Well-designed engineering controls can be highly effective in protecting workers and will typically be independent of worker interactions to provide this high level of protection. The basic Appropriate engineering types of engineering controls are: controls Process controls which involve changing the way a job activity or process is done to reduce the risk. Enclosure and/or isolation of emission source which keeps a selected hazard "physically" away from the worker and ventilation that strategically "adds" and "removes" air in the work environment. Ventilation can remove or dilute an air contaminant if designed properly. The design of a ventilation system must match the particular process and chemical or contaminant in use. Employers may need to use multiple types of controls to prevent employee overexposure. Personal protection ▶ Safety glasses with side shields. ▶ Chemical goggles • Contact lenses may pose a special hazard; soft contact lenses may absorb and concentrate irritants. A written policy document, describing the Eye and face protection wearing of lenses or restrictions on use, should be created for each workplace or task. This should include a review of lens absorption and adsorption for the class of chemicals in use and an account of injury experience. Medical and first-aid personnel should be trained in their removal and suitable equipment should be readily available. In the event of chemical exposure, begin eye irrigation immediately and remove contact lens Skin protection See Hand protection below Wear chemical protective gloves, e.g. PVC. ▶ Wear safety footwear or safety gumboots, e.g. Rubber The selection of suitable gloves does not only depend on the material, but also on further marks of quality which vary from manufacturer to manufacturer. Where the chemical is a preparation of several substances, the resistance of the glove material can not be calculated in advance and has therefore to be checked prior to the application. The exact break through time for substances has to be obtained from the manufacturer of the protective gloves and has to be observed when Suitability and durability of glove type is dependent on usage. Important factors in the selection of gloves include: • frequency and duration of contact, • Hands/feet protection chemical resistance of glove material, ▶ glove thickness and dexterity Select gloves tested to a relevant standard (e.g. Europe EN 374, US F739, AS/NZS 2161.1 or national equivalent). When prolonged or frequently repeated contact may occur, a glove with a protection class of 5 or higher (breakthrough time greater than 240 minutes according to EN 374, AS/NZS 2161.10.1 or national equivalent) is recommended. When only brief contact is expected, a glove with a protection class of 3 or higher (breakthrough time greater than 60 minutes according to EN 374, AS/NZS 2161.10.1 or national equivalent) is recommended. Neoprene rubber gloves See Other protection below **Body protection** Overalls. ▶ PVC Apron ▶ PVC protective suit may be required if exposure severe. ▶ Evewash unit. ▶ Ensure there is ready access to a safety shower. Other protection Some plastic personal protective equipment (PPE) (e.g. gloves, aprons, overshoes) are not recommended as they may produce static electricity. For large scale or continuous use wear tight-weave non-static clothing (no metallic fasteners, cuffs or pockets). Non sparking safety or conductive footwear should be considered. Conductive footwear describes a boot or shoe with a sole made from a conductive compound chemically bound to the bottom components, for permanent control to electrically ground the foot an shall dissipate static electricity from the body to reduce the possibility of ignition of volatile compounds. Electrical resistance must range between 0 to 500,000 ohms. Conductive shoes should be stored in lockers close to the room in which they are worn. Personnel who have been issued conductive footwear should not wear them from their place of work to their homes and return

Respiratory protection

Thermal hazards

Type A-P Filter of sufficient capacity. (AS/NZS 1716 & 1715, EN 143:2000 & 149:2001, ANSI Z88 or national equivalent)

Where the concentration of gas/particulates in the breathing zone, approaches or exceeds the "Exposure Standard" (or ES), respiratory protection is required. Degree of protection varies with both face-piece and Class of filter; the nature of protection varies with Type of filter.

Required Minimum Protection Factor	Half-Face Respirator	Full-Face Respirator	Powered Air Respirator
up to 10 x ES	A-AUS P2	-	A-PAPR-AUS / Class 1 P2
up to 50 x ES	-	A-AUS / Class 1 P2	-
up to 100 x ES	-	A-2 P2	A-PAPR-2 P2 ^

^{^ -} Full-face

A(All classes) = Organic vapours, B AUS or B1 = Acid gasses, B2 = Acid gas or hydrogen cyanide(HCN), B3 = Acid gas or hydrogen cyanide(HCN), E = Sulfur dioxide(SO2), G = Agricultural chemicals, K = Ammonia(NH3), Hg = Mercury, NO = Oxides of nitrogen, MB = Methyl bromide, AX = Low boiling point organic compounds(below 65 degC)

SECTION 9 PHYSICAL AND CHEMICAL PROPERTIES

Not Available

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Information	on h	hasic	nhysical	and	chemical	properties

Appearance	Clear, highly flammable liquid with a solvent odour; does not mixes in water.		
Physical state	Liquid	Relative density (Water = 1)	0.7
Odour	Not Available	Partition coefficient n- octanol / water	Not Available
Odour threshold	Not Available	Auto-ignition temperature (°C)	Not Available
pH (as supplied)	Not Applicable	Decomposition temperature	Not Available
Melting point / freezing point (°C)	Not Available	Viscosity (cSt)	Not Available
Initial boiling point and boiling range (°C)	>80	Molecular weight (g/mol)	Not Applicable
Flash point (°C)	35	Taste	Not Available
Evaporation rate	Not Available	Explosive properties	Not Available
Flammability	Flammable.	Oxidising properties	Not Available
Upper Explosive Limit (%)	6	Surface Tension (dyn/cm or mN/m)	Not Available
Lower Explosive Limit (%)	1	Volatile Component (%vol)	80
Vapour pressure (kPa)	Not Applicable	Gas group	Not Available
Solubility in water (g/L)	Immiscible	pH as a solution (1%)	Not Applicable
Vapour density (Air = 1)	Not Available	VOC g/L	Not Available

SECTION 10 STABILITY AND REACTIVITY

Reactivity	See section 7
Chemical stability	 ▶ Silicone fluids are stable under normal storage conditions. ▶ Hazardous polymerisation will not occur. ▶ At temperatures > 150 C, silicones can slowly react with the oxygen in air. ▶ When heated > 300 C, silicones can slowly depolymerise to volatile siloxanes whether or not air is present. ▶ Unstable in the presence of incompatible materials. ▶ Product is considered stable. ▶ Hazardous polymerisation will not occur.
Possibility of hazardous reactions	See section 7
Conditions to avoid	See section 7
Incompatible materials	See section 7
Hazardous decomposition products	See section 5

SECTION 11 TOXICOLOGICAL INFORMATION

Information on toxicological effects

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Low molecular weight silicone fluids may exhibit solvent action and may Skin exposure to isoparaffins may produce slight to moderate irritation in occurred. Open cuts, abraded or irritated skin should not be exposed to tentry into the blood-stream, through, for example, cuts, abrasions or lesion prior to the use of the material and ensure that any external damage is sue the exposure to silicone fluids causes temporary irritation of the conjuctorneal scarring, permanent eye damage, allergic reactions and catara produces only slight irritation. Limited evidence or practical experience suggests, that the material may contact may cause inflammation characterised by a temporary redness of Harmful: danger of serious damage to health by prolonged exposure through the support of the conjuctors.	roduce skin irritation. animals and humans. Rare sensitisation reactions in humans have iis material name produce systemic injury with harmful effects. Examine the skin itably protected. activa. Injection into the specific structures of the eye, however, causes it, and may lead to blindness. Instillation of isoparaffins into rabbit eyes cause eye irritation in a substantial number of individuals. Prolonged eye the conjunctiva (similar to windburn). agh inhalation. eriods. It can be assumed that it contains a substance which can produce ion and possible dermatitis following. ations but there is not enough data to make an assessment. some concern following repeated or long-term occupational affinic), to mouse skin, induced skin tumours; no tumours were ment and liver and blood changes. [PATTYS] uce stupor with dizziness, weakness and visual disturbance, weight loss	
Repeated exposure may cause skin cracking, flaking or drying following Skin contact with the material may damage the health of the individual; st. Low molecular weight silicone fluids may exhibit solvent action and may Skin exposure to isoparaffins may produce slight to moderate irritation in occurred. Open cuts, abraded or irritated skin should not be exposed to t Entry into the blood-stream, through, for example, cuts, abrasions or lesi prior to the use of the material and ensure that any external damage is st. Eye exposure to silicone fluids causes temporary irritation of the conju corneal scarring, permanent eye damage, allergic reactions and catara produces only slight irritation. Limited evidence or practical experience suggests, that the material may contact may cause inflammation characterised by a temporary redness of Harmful: danger of serious damage to health by prolonged exposure through the serious damage of one is exposed to it for long produced and the serious damage if one is exposed to it for long produced and the serious damage if one is exposed to it for long produced and the serious damage if one is exposed to it for long produced and the serious damage if one is exposed to it for long produced and the serious damage if one is exposed to it for long produced and the serious damage if one is exposed to it for long produced and the serious damage if one is exposed to it for long produced and the serious damage if one is exposed to it for long produced and the serious damage if one is exposed to it for long produced exposure. Repeated application of mildly hydrotreated oils (principally painduced with severely hydrotreated oils. Chronic solvent inhalation exposures may result in nervous system impact Constant or exposure over long periods to mixed hydrocarbons may produced with severely hydrotreated oils. Chronic solvent inhalation exposures may result in nervous system impact constant or exposure and reduced liver and kidney function. Skin exposure may contact the produced in the serious fu	roduce skin irritation. animals and humans. Rare sensitisation reactions in humans have is material ns, may produce systemic injury with harmful effects. Examine the skin itably protected. activa. Injection into the specific structures of the eye, however, causes it, and may lead to blindness. Instillation of isoparaffins into rabbit eyes cause eye irritation in a substantial number of individuals. Prolonged eye the conjunctiva (similar to windburn). Tagh inhalation. Tagh inhalation. Tagh inhalation but there is not enough data to make an assessment. The concern following repeated or long-term occupational affinic), to mouse skin, induced skin tumours; no tumours were sent and liver and blood changes. [PATTYS] The provided systems and visual disturbance, weight loss estupor with dizziness, weakness and visual disturbance, weight loss estupor with dizziness, weakness of the skin. TARITATION Not Available IRRITATION Not Available IRRITATION	
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The liquid may be miscible with fats or oils and may degrease the skin, p	ectives .	
(ICSC13733) Accidental ingestion of the material may be damaging to the health of the individual. Silicone fluids do not have a high acute toxicity. They may have a laxative effect and produce central nervous system depression. Isoparaffinic hydrocarbons cause temporary lethargy, weakness, inco-ordination and diarrhoea.		
Material is highly volatile and may quickly form a concentrated atmosphere in confined or unventilated areas. The vapour may displace and replace air in breathing zone, acting as a simple asphyxiant. This may happen with little warning of overexposure. Swallowing of the liquid may cause aspiration into the lungs with the risk of chemical pneumonitis; serious consequences may result.		
may be fatal. Nerve damage can be caused by some non-ring hydrocarbons. Symptor	as are temporary, and include weakness, tremors, increased saliva, some	
	Vapours of silicones are generally fairly well tolerated, however very high concentrations can cause death within minutes due to respiratory failure. At high temperatures, the fumes and oxidation products can be irritating and toxic and can cause depression leading to death in very high doses. Inhalation of high concentrations of gas/vapour causes lung irritation with coughing and nausea, central nervous depression with headache and dizziness, slowing of reflexes, fatigue and inco-ordination.	
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Version No: **3.1.1.1** Print Date: **23/03/2016**

Studies indicate that normal, branched and cyclic paraffins are absorbed from the mammalian gastrointestinal tract and that the absorption of n-paraffins is inversely proportional to the carbon chain length, with little absorption above C30. With respect to the carbon chain lengths likely to be present in mineral oil.

n-paraffins may be absorbed to a greater extent that iso- or cyclo-paraffins.

The major classes of hydrocarbons have been shown to be well absorbed by the gastrointestinal tract in various species. In many cases, the hydrophobic hydrocarbons are ingested in association with dietary lipids. The dependence of hydrocarbon absorption on concomitant triglyceride digestion and absorption, is known as the "hydrocarbon continuum hypothesis", and asserts that a series of solubilising phases in the intestinal lumen, created by dietary triglycerides and their digestion products, afford hydrocarbons a route to the lipid phase of the intestinal absorptive cell (enterocyte) membrane. While some hydrocarbons may traverse the mucosal epithelium unmetabolised and appear as solutes in lipoprotein particles in intestinal lymph, there is evidence that most hydrocarbons partially separate from nutrient lipids and undergo metabolic transformation in the enterocyte. The enterocyte may play a major role in determining the proportion of an absorbed hydrocarbon that, by escaping initial biotransformation, becomes available for deposition in its unchanged form in peripheral tissues such as adipose tissue, or in the liver.

The High Represe Naphthas (HBNs) contain mainly because hut its adverse health effect is more with other components, which may cause

adverse health effects involving a variety of organs. They may produce genetic damage as well as effects on reproduction and the unborn baby (generally at levels toxic to the mother). They may also cause cancers, for petroleum:

This product contains benzene which is known to cause acute myeloid leukaemia and n-hexane which has been shown to metabolize to compounds which are neuropathic.

This product contains toluene. There are indications from animal studies that prolonged exposure to high concentrations of toluene may lead to hearing loss.

This product contains ethyl benzene and naphthalene from which there is evidence of tumours in rodents

Carcinogenicity: Inhalation exposure to mice causes liver tumours, which are not considered relevant to humans. Inhalation exposure to rats causes kidney tumours which are not considered relevant to humans.

Mutagenicity: There is a large database of mutagenicity studies on gasoline and gasoline blending streams, which use a wide variety of endpoints and give predominantly negative results. All in vivo studies in animals and recent studies in exposed humans (e.g. petrol service station attendants) have shown negative

results in mutagenicity assays.

DHC Solvent Chemie (for EC No.: 926-605-8)

POLYDIMETHYLSILOXANE

NAPHTHA PETROLEUM.

LIGHT, HYDROTREATED

Siloxanes may impair liver and hormonal function, as well as the lung and kidney. They have not been found to be irritating to the skin and eyes. They may potentially cause cancer (tumours of the womb in females) and may cause impaired fertility or infertility.

The material may be irritating to the eye, with prolonged contact causing inflammation. Repeated or prolonged exposure to irritants may produce conjunctivitis

No toxic response noted during 90 day subchronic inhalation toxicity studies The no observable effect level is 450 mg/m3. Non-irritating and non-sensitising in human patch test. [Xerox]*

Acute Toxicity		Carcinogenicity	
Skin Irritation/Corrosion		Reproductivity	
Serious Eye Damage/Irritation	0	STOT - Single Exposure	•
Respiratory or Skin sensitisation	0	STOT - Repeated Exposure	•
Mutagenicity	0	Aspiration Hazard	~

Legend: X − Data available but does not fill the criteria for classification

Issue Date: 12/03/2015

Data required to make classification available
 Data Not Available to make classification

SECTION 12 ECOLOGICAL INFORMATION

Toxicity

Ingredient	Endpoint	Test Duration (hr)	Species	Value	Source
naphtha petroleum, light, hydrotreated	LC50	96	Fish	2.1- 61.1mg/L	2
naphtha petroleum, light, hydrotreated	EC50	48	Crustacea	4.7mg/L	2
naphtha petroleum, light, hydrotreated	EC50	72	Algae or other aquatic plants	12.4mg/L	2
naphtha petroleum, light, hydrotreated	EC50	96	Algae or other aquatic plants	1.6- 16.3mg/L	2
naphtha petroleum, light, hydrotreated	NOEC	72	Algae or other aquatic plants	6.47mg/L	2
polydimethylsiloxane	LC50	96	Fish	3.16mg/L	4

Extracted from 1. IUCLID Toxicity Data 2. Europe ECHA Registered Substances - Ecotoxicological Information - Aquatic Toxicity 3. EPIWIN Suite V3.12 -

Legend: Aquatic Toxicity Data (Estimated) 4. US EPA, Ecotox database - Aquatic Toxicity Data 5. ECETOC Aquatic Hazard Assessment Data 6. NITE (Japan) -

Harmful to aquatic organisms.

When spilled this product may act as a typical oil, causing a film, sheen, emulsion or sludge at or beneath the surface of the body of water. The oil film on water surface may physically affect the aquatic organisms, due to the interruption of the oxygen transfer between the air and the water Oils of any kind can cause:

representation of drowning of water-fowl due to lack of buoyancy, loss of insulating capacity of feathers, starvation and vulnerability to predators due to lack of mobility

Bioconcentration Data 7, METI (Japan) - Bioconcentration Data 8, Vendor Data

- ▶ lethal effects on fish by coating gill surfaces, preventing respiration
- ▶ asphyxiation of benthic life forms when floating masses become engaged with surface debris and settle on the bottom and ▶ adverse aesthetic effects of fouled shoreline and beaches

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In case of accidental releases on the soil, a fine film is formed on the soil, which prevents the plant respiration process and the soil particle saturation. It may cause deep water infestation. For Siloxanes:

Environmental Fate: Siloxanes are used in cosmetics, wax, polishes, and to a minor extent in several other applications.

Atmospheric Fate: In the presence of nitrate ions, short chain siloxanes are broken down by sunlight to the level of silicate within days. The main source atmospheric siloxane release to the air is via evaporation.

Aquatic Fate: It is well accepted that polydimethylsiloxane fluids become permanent residents of sediment but should not have adverse environmental effects. Silicone fluids are very surface active on surface waters. These substances tend to move into the aquatic compartment attached to textiles, sewage sludge, hair, algae, sediment, etc. Non-evaporating silicone fluids used in cosmetics, wax, polishes, cleaning products and those used in textile applications, (softeners), will, to a large extent, end up in wastewater and be directed to wastewater treatment plants.

Ecotoxicity: Siloxanes are chemically stable which makes them very persistent in the environment, where they are expected to remain for many years. For High Benzene Naphthas, (HBNs):

Environmental Fate: Some of these substances occur in crude oil and are formed as by-products from the combustion of natural materials, (e.g. during forest fires).

Atmospheric Fate: The chemical components in HBNs and are expected to evaporate significantly to the air where they are subject to rapid physical degradation through hydroxyl radical attack. Terrestrial Fate: These substances would only be found in soils where localized spills of petroleum products have occurred and during production of products containing these substances. HBNs have the potential to exhibit a high extent of biodegradability. Upon release to soil, some of these substances are expected to sink through the soil to groundwater. Many of these substances have a low to moderate tendency to sorb to soil particles. Sorption potential also varies with changes in other soil properties, (e.g. number of available adsorption sites, porosity and water content), and

environmental conditions.

DO NOT discharge into sewer or waterways.

Persistence and degradability

Ingredient	Persistence: Water/Soil	Persistence: Air
	No Data available for all ingredients	No Data available for all ingredients

Bioaccumulative potential

Ingredient	Bioaccumulation
	No Data available for all ingredients

Mobility in soil

Ingredient	Mobility
	No Data available for all ingredients

SECTION 13 DISPOSAL CONSIDERATIONS

Waste treatment methods

Legislation addressing waste disposal requirements may differ by country, state and/ or territory. Each user must refer to laws operating in their area. In some areas, certain wastes must be tracked.

A Hierarchy of Controls seems to be common - the user should investigate:

• Reduction •

Reuse

▶ Recycling

Disposal (if all else fails)

This material may be recycled if unused, or if it has not been contaminated so as to make it unsuitable for its intended use. If it has been contaminated, it may be possible to reclaim the product by filtration, distillation or some other means. Shelf life considerations should also be applied in making decisions of this type.

Product / Packaging disposal

Note that properties of a material may change in use, and recycling or reuse may not always be appropriate.

• DO NOT allow wash water from cleaning or process equipment to enter drains.

- t may be presented to collect all week water for treatment before dispensely
- \blacktriangleright It may be necessary to collect all wash water for treatment before disposal.
- ullet In all cases disposal to sewer may be subject to local laws and regulations and these should be considered

first. ► Where in doubt contact the responsible authority.

- ▶ Recycle wherever possible.
- ► Consult manufacturer for recycling options or consult local or regional waste management authority for disposal if no suitable treatment or disposal facility can be identified.
- ▶ Dispose of by: burial in a land-fill specifically licenced to accept chemical and / or pharmaceutical wastes or Incineration in a licenced apparatus (after admixture with suitable combustible material).
- ▶ Decontaminate empty containers. Observe all label safeguards until containers are cleaned and destroyed.

SECTION 14 TRANSPORT INFORMATION

Labels Required



Marine Pollutant

NO

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HAZCHEM Land transport (ADG)	•3YE			
Land transport (ADG)		•3YE		
	1003			
UN number	1993			
Packing group				
UN proper shipping name	FLAMMABLE LIQUID, N.O.S. (contains naphtha petroleum, light, hydrotreated)			
Environmental hazard	Not Applicable			
Transport hazard class(es)	Class 3 Subrisk Not Applicable			
Special precautions for user	Special provisions 274 Limited quantity 1 L			
Air transport (ICAO-IATA / DO	GR)			
UN number	1993			
Packing group	П			
UN proper shipping name	Flammable liquid, n.o.s. * (contains naphtha petroleum,	light, hydrotreated)		
Environmental hazard	Not Applicable			
Transport hazard class(es)	ICAO/IATA Class 3 ICAO / IATA Subrisk Not Applicable ERG Code 3H			
Special precautions for user	Special provisions Cargo Only Packing Instructions Cargo Only Maximum Qty / Pack Passenger and Cargo Packing Instructions Passenger and Cargo Maximum Qty / Pack Passenger and Cargo Limited Quantity Packing Instru Passenger and Cargo Limited Maximum Qty / Pack	A3 364 60 L 353 5 L Ctions Y341		
Sea transport (IMDG-Code / G	GGVSee)			
UN number	1993			
Packing group	II			
UN proper shipping name	FLAMMABLE LIQUID, N.O.S. (contains naphtha petroleum, light, hydrotreated)			
Environmental hazard	Not Applicable			
Transport hazard class(es)	IMDG Class 3 IMDG Subrisk Not Applicable			
Special precautions for user	EMS Number F-E, S-E Special provisions 274 Limited Quantities 1 L o Annex II of MARPOL and the IBC code			

Not Applicable

SECTION 15 REGULATORY INFORMATION

Safety, health and environmental regulations / legislation specific for the substance or mixture

NAPHTHA PETROLEUM, LIGHT, HYDROTREATED(64742-49-0.) IS FOUND ON THE FOLLOWING REGULATORY LISTS

Australia Hazardous Substances Information System - Consolidated Lists Australia Inventory of Chemical Substances (AICS)

POLYDIMETHYLSILOXANE(63148-62-9) IS FOUND ON THE FOLLOWING REGULATORY LISTS

Australia Inventory of Chemical Substances (AICS)

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National Inventory	Status
Australia - AICS	Y
Canada - DSL	Y
Canada - NDSL	N (polydimethylsiloxane; naphtha petroleum, light, hydrotreated)
China - IECSC	Y
Europe - EINEC / ELINCS / NLP	N (polydimethylsiloxane)
Japan - ENCS	N (naphtha petroleum, light, hydrotreated)
Korea - KECI	Y
New Zealand - NZIoC	Y
Philippines - PICCS	Y
USA - TSCA	Y
Legend:	Y = All ingredients are on the inventory N = Not determined or one or more ingredients are not on the inventory and are not exempt from listing(see specific ingredients in brackets)

SECTION 16 OTHER INFORMATION

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Other information

Classification of the preparation and its individual components has drawn on official and authoritative sources as well as independent review by the Chemwatch Classification committee using available literature references.

A list of reference resources used to assist the committee may be found at: www.chemwatch.net

The SDS is a Hazard Communication tool and should be used to assist in the Risk Assessment. Many factors determine whether the reported Hazards are Risks in the workplace or other settings. Risks may be determined by reference to Exposures Scenarios. Scale of use, frequency of use and current or available engineering controls must be considered.

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