

Toro Red Dye (for hydraulic oil)

The Toro Company

Chemwatch: 78-1332 Issue Date: 18/05/2017 Version No: 2.1.1.1

8/05/2017 Version No: 2.1.1.1 Print Date: 18/05/2017

Chemwatch Hazard Alert Code: 2

Safety Data Sheet according to WHS and ADG requirements

S.GHS.AUS.EN

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

Product Identifier	
Product name	Toro Red Dye (for hydraulic oil)
Synonyms	44-2500
Other means of identification	Not Available
Relevant identified uses of th	e substance or mixture and uses advised against
Relevant identified uses	Dye.
Details of the supplier of the	safety data sheet
Registered company name	Toro Australia
Address	53 Howards Road, Beverly, South Australia 5009, Australia
Telephone	08 8300 3633
Fax	08 8243 2940
Website	www.toro.com.au
Email	info.au.toro.com
Emergency telephone number	er som
Association / Organisation	State EPA
Emergency telephone numbers	000 (Police, Fire, Ambulance)
Other emergency telephone numbers	Poisons Information 131 126

SECTION 2 HAZARDS IDENTIFICATION

Poisons Schedule	Not Applicable
[1] Classification	Skin Corrosion/Irritation Category 2, Eye Irritation Category 2A, Reproductive Toxicity Category 2, Specific target organ toxicity - single exposure Category 3 (respiratory tract irritation), Specific target organ toxicity - single exposure Category 3 (narcotic effects)
Legend:	1. Classified by Chemwatch; 2. Classification drawn from HSIS ; 3. Classification drawn from EC Directive 1272/2008 - Annex VI
abel elements	
GHS label elements	
SIGNAL WORD	WARNING
azard statement(s)	
	Causes skin irritation.
H315	

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H361	Suspected of damaging fertility or the unborn child.
H335	May cause respiratory irritation.
H336	May cause drowsiness or dizziness.
Precautionary statement(s) F	revention
P201	Obtain special instructions before use.
P271	Use only outdoors or in a well-ventilated area.
P281	Use personal protective equipment as required.
P261	Avoid breathing mist/vapours/spray.
P280	Wear protective gloves/protective clothing/eye protection/face protection.
Precautionary statement(s) F	esponse
P308+P313	IF exposed or concerned: Get medical advice/attention.
P362	Take off contaminated clothing and wash before reuse.
P305+P351+P338	IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.
P312	Call a POISON CENTER or doctor/physician if you feel unwell.
P337+P313	If eye irritation persists: Get medical advice/attention.
P302+P352	IF ON SKIN: Wash with plenty of soap and water.
Precautionary statement(s) S	torage
P405	Store locked up.
P403+P233	Store in a well-ventilated place. Keep container tightly closed.
Precautionary statement(s)	isposal
P501	Dispose of contents/container in accordance with local regulations.

SECTION 3 COMPOSITION / INFORMATION ON INGREDIENTS

Substances

See section below for composition of Mixtures

Mixtures

CAS No	%[weight]	Name
64742-53-6.	50-60	naphthenic distillate, light, hydrotreated (severe)
71819-51-7	40-50	C.I. Solvent Red 164
1330-20-7	<0.9	xylene
100-41-4	<0.3	ethylbenzene

SECTION 4 FIRST AID MEASURES

Eye Contact	 If this product comes in contact with the eyes: 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	If skin contact occurs: 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.
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, without delay.

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Ingestion	 For advice, contact a Poisons Information Centre or a doctor at once. Urgent hospital treatment is likely to be needed. 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.
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Indication of any immediate medical attention and special treatment needed

Treat symptomatically.

- + Heavy and persistent skin contamination over many years may lead to dysplastic changes. Pre-existing skin disorders may be aggravated by exposure to this product.
- In general, emesis induction is unnecessary with high viscosity, low volatility products, i.e. most oils and greases.
- + High pressure accidental injection through the skin should be assessed for possible incision, irrigation and/or debridement.
- NOTE: Injuries may not seem serious at first, but within a few hours tissue may become swollen, discoloured and extremely painful with extensive subcutaneous necrosis. Product may be forced through considerable distances along tissue planes.

Periodic medical surveillance should be carried out on persons in occupations exposed to the manufacture or bulk handling of the product and this should include hepatic function tests and urinalysis examination. [ILO Encyclopaedia]

- For acute or short term repeated exposures to xylene:
 - Gastro-intestinal absorption is significant with ingestions. For ingestions exceeding 1-2 ml (xylene)/kg, intubation and lavage with cuffed endotracheal tube is recommended. The use of charcoal and cathartics is equivocal.
 - Pulmonary absorption is rapid with about 60-65% retained at rest.
 - Primary threat to life from 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 or pCO2 > 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 catecholamine's. Inhaled cardio selective bronchodilators (e.g. Alupent, Salbutamol) are the preferred agents, with aminophylline a second choice.
 - BIOLOGICAL EXPOSURE INDEX BEI

These represent the determinants observed in speci	imens collected from a healthy worker exposed at	the Exposure Standard (ES or TLV):	
Determinant Methylhippu-ric acids in urine	Index 1.5 gm/gm creatinine	Sampling Time End of shift	Comments
	2 mg/min	Last 4 hrs of shift	

SECTION 5 FIREFIGHTING MEASURES

Extinguishing media

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

Special hazards arising from the substrate or mixture

Fire Incompatibility	Avoid contamination with oxidising agents i.e. nitrates, oxidising acids, chlorine bleaches, pool chlorine etc. as ignition may result
Advice for firefighters	
Fire Fighting	 Alert Fire Brigade and tell them location and nature of hazard. Wear full body protective clothing with breathing apparatus. Prevent, by any means available, spillage from entering drains or water course. Use water delivered as a fine spray to control fire and cool adjacent area. Avoid spraying water onto liquid pools. DO NOT approach containers suspected to be hot.
Fire/Explosion Hazard	 Combustible. Slight fire hazard when exposed to heat or flame. Heating may cause expansion or decomposition leading to violent rupture of containers. I on combustion, may emit toxic fumes of carbon monoxide (CO). May emit acrid smoke. Mists containing combustible materials may be explosive. Combustion products include: , carbon dioxide (CO2) , nitrogen oxides (NOx) other pyrolysis products typical of burning organic material. May emit poisonous fumes. May emit corrosive fumes. CARE: Water in contact with hot liquid may cause foaming and a steam explosion with wide scattering of hot oil and possible severe burns.
HAZCHEM	Not Applicable

SECTION 6 ACCIDENTAL RELEASE MEASURES

Personal precautions, protective equipment and emergency procedures

See section 8

Environmental precautions

See section 12

Methods and material for containment and cleaning up

Minor Spills	 Slippery when spilt. Remove all ignition sources. Clean up all spills immediately. Avoid breathing vapours and contact with skin and eyes. Control personal contact with the substance, by using protective equipment. Contain and absorb spill with sand, earth, inert material or
	vermiculite. • Wipe up.
Major Spills	 Slippery when spilt. Moderate hazard. Clear area of personnel and move upwind. Alert Fire Brigade and tell them location and nature of hazard. Wear breathing apparatus plus protective gloves.
	 Prevent, by any means available, spillage from entering drains or water course. No smoking, naked lights or ignition sources.
Personal Protective Equipment	advice is contained in Section 8 of the SDS.

SECTION 7 HANDLING AND STORAGE

	► Avoid all personal contact, including inhalation.
	► Wear protective clothing when risk of exposure occurs.
	► Use in a well-ventilated area.
Safe handling	▶ Prevent concentration in hollows and sumps.
	DO NOT enter confined spaces until atmosphere has been checked.
	Avoid smoking, naked lights or ignition sources.
	▶ Store in original containers.
	▶ Keep containers securely sealed.
Other information	► No smoking, naked lights or ignition sources.
Other Information	▶ Store in a cool, dry, well-ventilated area.
	Store away from incompatible materials and foodstuff containers.
	 Protect containers against physical damage and check regularly for leaks.
itions for safe storage,	ncluding any incompatibilities
	▶ Metal can or drum
Suitable container	Packaging as recommended by manufacturer.
	► Check all containers are clearly labelled and free from leaks.
Storage incompatibility	CARE: Water in contact with heated material may cause foaming or a steam explosion with possible severe burns from wide scattering of hot
moompationity	material. Resultant overflow of containers may result in fire. ► Avoid reaction with oxidising agents

SECTION 8 EXPOSURE CONTROLS / PERSONAL PROTECTION

Control parameters

OCCUPATIONAL EXPOSURE LIMITS (OEL)

INGREDIENT DATA

Source	Ingredient	Material name	TWA	STEL	Peak	Notes
Australia Exposure Standards	naphthenic distillate, light, hydrotreated (severe)	Oil mist, refined mineral	5 mg/m3	Not Available	Not Available	Not Available
Australia Exposure Standards	xylene	Xylene (o-, m-, pisomers)	350 mg/m3 / 80 ppm	655 mg/m3 / 150 ppm	Not Available	Not Available

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Australia Exposure Standards	ethylbenzene	Ethyl benzene	434 mg/m3 / 100 ppm	543 mg/m3 / 125 ppm	Not Available	Not Available
EMERGENCY LIMITS						
Ingredient	Material name	TEEL-1	TEEL-2	·	TEEL-3	
xylene	Xylenes	Not Available	Not Available		Not Available	
ethylbenzene	Ethyl benzene	Not Available	Not Available		Not Available	
Ingredient	Original IDLH		Revised IDLH			
naphthenic distillate, light, hydrotreated (severe)	Not Available		Not Available			
C.I. Solvent Red 164	Not Available		Not Available			
xylene	1,000 ppm		900 ppm			
ethylbenzene	2,000 ppm		800 [LEL] ppm			
oposure controls			1			
Appropriate engineering controls	Engineering controls are used to remove can be highly effective in protecting wor basic types of engineering controls are: Process controls which involve changing Enclosure and/or isolation of emission s "adds" and "removes" air in the work en ventilation system must match the partic	kers and will typically be inde g the way a job activity or pro ource which keeps a selected vironment. Ventilation can rer	pendent of worker intera icess is done to reduce t d hazard "physically" aw move or dilute an air con	actions to provide this I the risk. /ay from the worker and	high level of prot d ventilation that	ection. The strategically
Personal protection						
Eye and face 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 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. 					
Skin protection	See Hand protection below					
Hands/feet protection	 Wear chemical protective gloves, e.g. Wear safety footwear or safety gum The selection of suitable gloves does no manufacturer. Where the chemical is a and has therefore to be checked prior to The exact break through time for substa making a final choice. Personal hygiene is a key element of ef and dried thoroughly. Neoprene rubber gloves 	ooots, e.g. Rubber ot only depend on the materia preparation of several substa o the application. ances has to be obtained from	nces, the resistance of t	the glove material can e protective gloves and	not be calculated	l in advance rved when
Body protection	See Other protection below					
	Overalls. P.V.C. apron. Barrier cream. Skin cleansing cream.					
Other protection	▶ Barrier cream.					

GLOVE SELECTION INDEX

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Glove selection is based on a modified presentation of the:

"Forsberg Clothing Performance Index".

The effect(s) of the following substance(s) are taken into account in the Toro Red Dye (for hydraulic oil)

Material	СРІ
BUTYL	A
BUTYL/NEOPRENE	A
HYPALON	A
NAT+NEOPR+NITRILE	С
NATURAL+NEOPRENE	A
NEOPRENE	A
NEOPRENE/NATURAL	A
NITRILE	A
NITRILE+PVC	A
PE/EVAL/PE	A
PVA	В
PVC	A
PVDC/PE/PVDC	A

* CPI - Chemwatch Performance Index

A: Best Selection

B: Satisfactory; may degrade after 4 hours continuous immersion

C: Poor to Dangerous Choice for other than short term immersion

NOTE: As a series of factors will influence the actual performance of the glove, a final selection must be based on detailed observation. -

* Where the glove is to be used on a short term, casual or infrequent basis, factors such as "feel" or convenience (e.g. disposability), may dictate a choice of gloves which might otherwise be unsuitable following long-term or frequent use. A qualified practitioner should be consulted.

Respiratory protection

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 5 x ES	A-AUS / Class 1 P2	-	A-PAPR-AUS / Class 1 P2
up to 25 x ES	Air-line*	A-2 P2	A-PAPR-2 P2
up to 50 x ES	-	A-3 P2	-
50+ x ES	-	Air-line**	-

* - Continuous-flow; ** - Continuous-flow or positive pressure demand

^ - 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)

Cartridge respirators should never be used for emergency ingress or in areas of unknown vapour concentrations or oxygen content. The wearer must be warned to leave the contaminated area immediately on detecting any odours through the respirator. The odour may indicate that the mask is not functioning properly, that the vapour concentration is too high, or that the mask is not properly fitted. Because of these limitations, only restricted use of cartridge respirators is considered appropriate.

SECTION 9 PHYSICAL AND CHEMICAL PROPERTIES

Information on basic physical and chemical properties

Appearance	Red liquid with petroleum odour; does not mix with water.		
Physical state	Liquid	Relative density (Water = 1)	0.94
Odour	Not Available	Partition coefficient n- octanol / water	Not Available

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Odour threshold	Not Available	Auto-ignition temperature (°C)	307.0
pH (as supplied)	Not Available	Decomposition temperature	Not Available
Melting point / freezing point (°C)	Not Available	Viscosity (cSt)	100-118
Initial boiling point and boiling range (°C)	204-327	Molecular weight (g/mol)	Not Applicable
Flash point (°C)	>93 (Seta flash CC)	Taste	Not Available
Evaporation rate	Not Available	Explosive properties	Not Available
Flammability	Not Applicable	Oxidising properties	Not Available
Upper Explosive Limit (%)	Not Available	Surface Tension (dyn/cm or mN/m)	Not Available
Lower Explosive Limit (%)	Not Available	Volatile Component (%vol)	Not Available
Vapour pressure (kPa)	Not Available	Gas group	Not Available
Solubility in water (g/L)	Immiscible	pH as a solution (1%)	Not Available
Vapour density (Air = 1)	Not Available	VOC g/L	Not Available

SECTION 10 STABILITY AND REACTIVITY

Reactivity	See section 7
Chemical stability	 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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The material can cause respiratory irritation in some persons. The body's response to such irritation can cause further lung damage. Inhalation of vapours may cause drowsiness and dizziness. This may be accompanied by sleepiness, reduced alertness, loss of reflexes, lack of co-ordination, and vertigo. Inhalation of aerosols (mists, fumes), generated by the material during the course of normal handling, may be damaging to the health of the individual. Inhalation hazard is increased at higher temperatures. Inhaling high concentrations of mixed hydrocarbons can cause narcosis, with nausea, vomiting and lightheadedness. Low molecular weight (C2-C12) hydrocarbons can irritate mucous membranes and cause incoordination, giddiness, nausea, vertigo, confusion, headache, appetite loss, drowsiness, tremors and stupor. Central nervous system (CNS) depression may include general discomfort, symptoms of giddiness, headache, dizziness, nausea, anaesthetic effects, slowed reaction time, slurred speech and may progress to unconsciousness. Serious poisonings may result in respiratory depression and may be fatal. Inhalation of oil droplets or aerosols may cause discomfort and may produce chemical inflammation of the lungs. Inhaled The acute toxicity of inhaled alkylbenzenes is best described by central nervous system depression. As a rule, these compounds may also act as general anaesthetics Systemic poisoning produced by general anaesthesia is characterised by lightheadedness, nervousness, apprehension, euphoria, confusion, dizziness, drowsiness, tinnitus, blurred or double vision, vomiting and sensations of heat, cold or numbness, twitching, tremors, convulsions, unconsciousness and respiratory depression and arrest. Cardiac arrest may result from cardiovascular collapse. Bradycardia, and hypotension may also be produced. Inhaled alkylbenzene vapours cause death in animals at air levels that are relatively similar (typically LC50s are in the range 5000 -8000 ppm for 4 to 8 hour exposures). Headache, fatigue, tiredness, irritability and digestive disturbances (nausea, loss of appetite and bloating) are the most common symptoms of xylene overexposure. Injury to the heart, liver, kidneys and nervous system has also been noted amongst workers. Xylene is a central nervous system depressant Accidental ingestion of the material may be damaging to the health of the individual. Ingestion of petroleum hydrocarbons can irritate the pharynx, oesophagus, stomach and small intestine, and cause swellings and ulcers of the mucous. Ingestion Symptoms include a burning mouth and throat; larger amounts can cause nausea and vomiting, narcosis, weakness, dizziness, slow and shallow breathing, abdominal swelling, unconsciousness and convulsions. This material can cause inflammation of the skin on contact in some persons. The material may accentuate any pre-existing dermatitis condition Repeated exposure may cause skin cracking, flaking or drying following normal handling and use. Open cuts, abraded or irritated skin should not be exposed to this material Skin Contact Entry into the blood-stream, through, for example, cuts, abrasions or lesions, may produce systemic injury with harmful effects. Examine the skin prior to the use of the material and ensure that any external damage is suitably protected. Aromatic hydrocarbons may produce sensitivity and redness of the skin. They are not likely to be absorbed into the body through the skin but branched species are more likely to. This material can cause eve irritation and damage in some persons. Direct eye contact with petroleum hydrocarbons can be painful, and the corneal epithelium may be temporarily damaged. Aromatic species can cause irritation Eve and excessive tear secretion. Ample evidence from experiments exists that there is a suspicion this material directly reduces fertility. Substance accumulation, in the human body, may occur and may cause some concern following repeated or long-term occupational exposure. There is some evidence from animal testing that exposure to this material may result in toxic effects to the unborn baby. Constant or exposure over long periods to mixed hydrocarbons may produce stupor with dizziness, weakness and visual disturbance, weight loss and anaemia, and reduced liver and kidney function. Skin exposure may result in drying and cracking and redness of the skin. Chronic Women exposed to xylene in the first 3 months of pregnancy showed a slightly increased risk of miscarriage and birth defects. Evaluation of workers chronically exposed to xylene has demonstrated lack of genetic toxicity. Exposure to the material for prolonged periods may cause physical defects in the developing embryo (teratogenesis). Oil may contact the skin or be inhaled. Extended exposure can lead to eczema, inflammation of hair follicles, pigmentation of the face and warts on the soles of the feet. TOXICITY IRRITATION Toro Red Dye (for hydraulic oil) Not Available Not Available naphthenic distillate, light, IRRITATION TOXICITY hydrotreated (severe) [2] Not Available Dermal (rabbit) LD50: >2000 mg/kg [1] Inhalation (rat) LC50: >3.9 ma/l/4hr [1] Inhalation (rat) LC50: >4.7 mg/l/4hr [1] Inhalation (rat) LC50: >5 mg/l/4hr [1] Inhalation (rat) LC50: >5.2 ma/l/4hr [1] Inhalation (rat) LC50: >5.3 mg/l/4hr [1] Inhalation (rat) LC50: 10.5 mg/l/4hr [1] Inhalation (rat) LC50: 5.7 mg/l/4hr [1] Inhalation (rat) LC50: 9.6 mg/l/4hr

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	[1]	
	Oral (rat) LD50: >2000 mg/kg	
	ΤΟΧΙΟΙΤΥ	IRRITATION
C.I. Solvent Red 164		
	Not Available	Not Available
	TOXICITY	IRRITATION
xylene	Dermal (rabbit) LD50: >1700 mg/kg	Eye (human): 200 ppm irritant
Inhalation (rat) LC50: 5000 ppm/4hr		Eye (rabbit): 5 mg/24h SEVERE
Oral (rat) LD50: 4300 mg/k ²		Eye (rabbit): 87 mg mild
		Skin (rabbit):500 mg/24h moderate
	TOXICITY	IRRITATION
ethylbenzene	Dermal (rabbit) LD50: ca.15432.6 mo	Eye (rabbit): 500 mg - SEVERE
ethylbenzene	Inhalation (mouse) LC50: 35.5 mg/L/2hr	Skin (rabbit): 15 mg/24h mild
	Inhalation (rat) LC50: 55 mg/L/2fr	
	Oral (rat) LD50: 3500 mg/k ²	
Leį		Acute toxicity 2.* Value obtained from manufacturer's SDS. Unless otherwise specified
	data extracted from RTECS - Register of Toxic Effect of ch	emical Substances
	ED refined oils by removing or transforming undesirable components. In comparison to unrefined and mildly refined base oils, the highly and severely	
NAPHTHENIC DISTILI LIGHT, HYDROTRE (SEV C.I. SOLVENT RE	LATE, ATED, VERE) Unrefined & mildly refined distillate base oils contain the molecules and have shown the highest potential carcinoger refined distillate base oils by removing or transforming undesirable compor- refined distillate base oils have a smaller range of hydrocarbon mol carcinogenicity testing of residual oils has been negative, su components are largely non-bioavailable due to their molecu- For highly and severely refined distillate base oils: In animal studies, the acute, oral, semilethal dose is >5g/kg semilethal concentration for inhalation is 2.18 to >4 mg/L. The skin and eye irritation. Testing for sensitisation has been ne testes and lung have been observed, as well as the formation reproductive toxicity or significant increases in birth defects.	ate base oils is inversely related to the degree of processing. highest levels of undesirable components, have the largest variation of hydrocarbon ic and mutagenic activities. Highly and severely refined distillate base oils are produced from unrefined and mildly ments. In comparison to unrefined and mildly refined base oils, the highly and severely ecules and have demonstrated very low mammalian toxicity. Mutagenicity and pporting the belief that these materials lack biologically active components or the lar size. body weight and the semilethal dose by skin contact is >2g/kg body weight. The we materials have varied from "non-irritating" to "moderately irritating" when tested for pative. The effects of repeated exposure vary by species; in animals, effects to the n of granulomas. In animals, these substances have not been found to cause
LIGHT, HYDROTRE (SEV C.I. SOLVENT RE	LATE, ATED, VERE) Unrefined & mildly refined distillate base oils contain the molecules and have shown the highest potential carcinoger refined distillate base oils by removing or transforming undesirable compor- refined distillate base oils have a smaller range of hydrocarbon mol carcinogenicity testing of residual oils has been negative, su components are largely non-bioavailable due to their molecu- For highly and severely refined distillate base oils: In animal studies, the acute, oral, semilethal dose is >5g/kg semilethal concentration for inhalation is 2.18 to >4 mg/L. The skin and eye irritation. Testing for sensitisation has been ne testes and lung have been observed, as well as the formation reproductive toxicity or significant increases in birth defects.	ate base oils is inversely related to the degree of processing. highest levels of undesirable components, have the largest variation of hydrocarbon ic and mutagenic activities. Highly and severely refined distillate base oils are produced from unrefined and mildly ments. In comparison to unrefined and mildly refined base oils, the highly and severely ecules and have demonstrated very low mammalian toxicity. Mutagenicity and pporting the belief that these materials lack biologically active components or the lar size. body weight and the semilethal dose by skin contact is >2g/kg body weight. The we materials have varied from "non-irritating" to "moderately irritating" when tested for pative. The effects of repeated exposure vary by species; in animals, effects to the n of granulomas. In animals, these substances have not been found to cause
LIGHT, HYDROTRE (SEV C.I. SOLVENT RE	LATE, indexUnrefined & mildly refined distillate base oils contain the molecules and have shown the highest potential carcinoger refined distillate base oils by removing or transforming undesirable compo- refined distillate base oils have a smaller range of hydrocarbon mol carcinogenicity testing of residual oils has been negative, su components are largely non-bioavailable due to their molecule For highly and severely refined distillate base oils: In animal studies, the acute, oral, semilethal dose is >5g/kg semilethal concentration for inhalation is 2.18 to >4 mg/L.D 164No significant acute toxicological data identified in literature reproductive effector in ratsCLENEEthylbenzene is readily absorbed when inhaled, swallowed urine. It may irritate the skin, eyes and may cause hearing la liver and lungs, including a tendency to cancer formation, according to animal testing. The disting a tendency to cancer formation, according to animal testing. The stant of the skin and eye is readily absorbed when inhaled to a size the skin and eye is readily absorbed in the skin and eye is readily absorbed when inhaled action in the stant acute toxicological data identified in literatureCLENEEthylbenzene is readily absorbed when inhaled action is a set of the skin and eye is a set of the skin according to animal testing. The stant acute formation, according to animal testing. The stant acute formation is a set of the skin according to animal testing. The stant acute formation, according to animal testing. The stant acute formation is according to animal testing. The stant acute formation	ate base oils is inversely related to the degree of processing. highest levels of undesirable components, have the largest variation of hydrocarbon ic and mutagenic activities. Highly and severely refined distillate base oils are produced from unrefined and mildly ments. In comparison to unrefined and mildly refined base oils, the highly and severely ecules and have demonstrated very low mammalian toxicity. Mutagenicity and pporting the belief that these materials lack biologically active components or the lar size. body weight and the semilethal dose by skin contact is >2g/kg body weight. The we materials have varied from "non-irritating" to "moderately irritating" when tested for pative. The effects of repeated exposure vary by species; in animals, effects to the n of granulomas. In animals, these substances have not been found to cause
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Skin Irritation/Corrosion Reproductivity Serious Eye STOT - Single Exposure Damage/Irritation 0 0 Respiratory or Skin STOT - Repeated sensitisation Exposure 0 Aspiration Hazard Mutagenicity Legend: - Data available but does not fill the criteria for classification

- Data available to make classification

- Data Not Available to make classification

SECTION 12 ECOLOGICAL INFORMATION

Ingredient	Endpoint	Test Duration (hr)	Species	Value	Source
naphthenic distillate, light, hydrotreated (severe)	EC50	48	Crustacea	>1000mg/L	1
naphthenic distillate, light, hydrotreated (severe)	EC50	96	Algae or other aquatic plants	>1000mg/L	1
naphthenic distillate, light, hydrotreated (severe)	EC50	96	Algae or other aquatic plants	>1000mg/L	1
naphthenic distillate, light, hydrotreated (severe)	NOEC	504	Crustacea	>1mg/L	1
xylene	LC50	96	Fish	2.6mg/L	2
xylene	EC50	48	Crustacea	>3.4mg/L	2
xylene	EC50	72	Algae or other aquatic plants	4.6mg/L	2
xylene	EC50	24	Crustacea	0.711mg/L	4
xylene	NOEC	73	Algae or other aquatic plants	0.44mg/L	2
ethylbenzene	LC50	96	Fish	0.0043mg/L	4
ethylbenzene	EC50	48	Crustacea	1.184mg/L	4
ethylbenzene	EC50	96	Algae or other aquatic plants	3.6mg/L	2
ethylbenzene	EC50	96	Crustacea	=0.49mg/L	1
ethylbenzene	NOEC	168	Crustacea	0.96mg/L	5

(QSAR) - Aquatic Toxicity Data (Estimated) 4. US EPA, Ecotox database - Aquatic Toxicity Data 5. ECETOC Aquatic Hazard Assessment Data 6. NITE (Japan) - Bioconcentration Data 7. METI (Japan) - Bioconcentration Data 8. Vendor Data

DO NOT discharge into sewer or waterways.

Persistence and degradability

Ingredient	Persistence: Water/Soil	Persistence: Air
xylene	HIGH (Half-life = 360 days)	LOW (Half-life = 1.83 days)
ethylbenzene	HIGH (Half-life = 228 days)	LOW (Half-life = 3.57 days)
Bioaccumulative potential		
Ingredient	Bioaccumulation	
xylene	MEDIUM (BCF = 740)	
ethylbenzene	LOW (BCF = 79.43)	
Mobility in soil		
Ingredient	Mobility	
ethylbenzene	LOW (KOC = 517.8)	

Toro Red Dye (for hydraulic oil)

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Recycle wherever possible or consult manufacture	er for recycling options.
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Consult State Land Waste Authority for disposal.

- Bury or incinerate residue at an approved site.
- Recycle containers if possible, or dispose of in an authorised landfill.

SECTION 13 DISPOSAL CONSIDERATIONS

Waste treatment methods	
Product / Packaging disposal	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. DO NOT allow wash water from cleaning or process equipment to enter drains. It may be necessary to collect all wash water for treatment before disposal. 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.
	Where in doubt contact the responsible authority.

SECTION 14 TRANSPORT INFORMATION

Labels Required Marine Pollutant NO HAZCHEM Not Applicable Land transport (ADG): NOT REGULATED FOR TRANSPORT OF DANGEROUS GOODS Air transport (ICAO-IATA / DGR): NOT REGULATED FOR TRANSPORT OF DANGEROUS GOODS

Sea transport (IMDG-Code / GGV See): NOT REGULATED FOR TRANSPORT OF DANGEROUS GOODS

Transport in bulk according to 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

NAPHTHENIC DISTILLATE, LIGHT, HYDROTREATED (SEVERE)(64742-53-6.) IS FOUND ON THE FOLLOWING REGULATORY LISTS

Australia Exposure Standards		Australia Inventory of Chemical Substances (AICS)			
Australia Hazardous Substances Information System - Consolidated Lists					
C.I. SOLVENT RED 164(71819-5	C.I. SOLVENT RED 164(71819-51-7) IS FOUND ON THE FOLLOWING REGULATORY LISTS				
Australia Inventory of Chemical S	Substances (AICS)				
XYLENE(1330-20-7) IS FOUND	ON THE FOLLOWING REGULATORY LISTS				
Australia Exposure Standards		Australia Inventory of Chemical Substances (AICS)			
Australia Hazardous Substance	Australia Hazardous Substances Information System - Consolidated Lists				
ETHYLBENZENE(100-41-4) IS F	OUND ON THE FOLLOWING REGULATORY LISTS				
Australia Exposure Standards		Australia Inventory of Chemical Substances (AICS)			
Australia Hazardous Substances Information System - Consolidated Lists					
National Inventory	Status				
Australia - AICS	Y				
Canada - DSL	Υ				

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Canada - NDSL	N (naphthenic distillate, light, hydrotreated (severe); xylene; ethylbenzene; C.I. Solvent Red 164)
China - IECSC	Y
Europe - EINEC / ELINCS / NLP	Y
Japan - ENCS	N (C.I. Solvent Red 164)
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

Other information

Ingredients with multiple cas numbers

Name	CAS No
C.I. Solvent Red 164	71819-51-7, 92257-31-3

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.

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.

Definitions and abbreviations

PC-TWA: Permissible Concentration-Time Weighted Average

PC-STEL: Permissible Concentration-Short Term Exposure Limit

IARC: International Agency for Research on Cancer

- ACGIH: American Conference of Governmental Industrial Hygienists
- STEL: Short Term Exposure Limit
- TEEL: Temporary Emergency Exposure Limit。
- IDLH: Immediately Dangerous to Life or Health Concentrations
- OSF: Odour Safety Factor
- NOAEL :No Observed Adverse Effect Level

LOAEL: Lowest Observed Adverse Effect Level

- TLV: Threshold Limit Value
- LOD: Limit Of Detection
- OTV: Odour Threshold Value BCF: BioConcentration Factors
- BEI: Biological Exposure Index

BEI. Diological Exposure maex

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