

21696 Octane Booster LIQUI MOLY Australia Pty Limited

Chemwatch: 5529-01

Version No: 2.1

Safety Data Sheet according to WHS Regulations (Hazardous Chemicals) Amendment 2020 and ADG requirements

SECTION 1 Identification of the substance / mixture and of the company / undertaking

Product Identifier		
Product name	21696 Octane Booster	
Chemical Name	Not Applicable	
Synonyms	Not Available	
Chemical formula	Not Applicable	
Other means of identification	Not Available	

Relevant identified uses of the substance or mixture and uses advised against

Relevant identified uses	Additive.
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Details of the supplier of the safety data sheet

Registered company name	LIQUI MOLY Australia Pty Limited		
Address	Suite 106, 26-32 Pirrama Road Pyrmont NSW 2009 Australia		
Telephone	1300 318 961		
Fax	Not Available		
Website	www.liqui-moly.com.au		
Email	Not Available		

Emergency telephone number

Association / Organisation	LIQUI MOLY Australia Pty Limited	CHEMWATCH EMERGENCY RESPONSE
Emergency telephone numbers	13 11 26 (Poisons Information Centre)	+61 1800 951 288
Other emergency telephone numbers	Not Available	+61 2 9186 1132

Once connected and if the message is not in your prefered language then please dial 01

SECTION 2 Hazards identification

Classification of the substance or mixture

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

COMBUSTIBLE LIQUID, regulated for storage purposes only

ChemWatch Hazard Ratings

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

Poisons Schedule	Not Applicable
Classification [1]	Flammable Liquids Category 4, Aspiration Hazard Category 1, Sensitisation (Skin) Category 1, Serious Eye Damage/Eye Irritation Category 1, Acute Toxicity (Inhalation) Category 4, Specific Target Organ Toxicity - Single Exposure (Narcotic Effects) Category 3, Hazardous to the Aquatic Environment Long-Term Hazard Category 3
Legend:	1. Classified by Chemwatch; 2. Classification drawn from HCIS; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI

Chemwatch Hazard Alert Code: 2

Issue Date: 22/02/2022 Print Date: 22/02/2022 S.GHS.AUS.EN



Signal word Danger

Hazard statement(s)

AUH066	Repeated exposure may cause skin dryness and cracking.	
H227	Combustible liquid.	
H304	May be fatal if swallowed and enters airways.	
H317	May cause an allergic skin reaction.	
H318	Causes serious eye damage.	
H332	Harmful if inhaled.	
H336	May cause drowsiness or dizziness.	
H412	Harmful to aquatic life with long lasting effects.	

Precautionary statement(s) Prevention

P210	Keep away from heat, hot surfaces, sparks, open flames and other ignition sources. No smoking.
P271	Use only outdoors or in a well-ventilated area.
P280	Wear protective gloves, protective clothing, eye protection and face protection.
P261	Avoid breathing mist/vapours/spray.
P273	Avoid release to the environment.
P272	Contaminated work clothing should not be allowed out of the workplace.

Precautionary statement(s) Response

P301+P310	IF SWALLOWED: Immediately call a POISON CENTER/doctor/physician/first aider.
P331	Do NOT induce vomiting.
P305+P351+P338	IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.
P370+P378	In case of fire: Use alcohol resistant foam or normal protein foam to extinguish.
P302+P352	IF ON SKIN: Wash with plenty of water and soap.
P333+P313	If skin irritation or rash occurs: Get medical advice/attention.
P362+P364	Take off contaminated clothing and wash it before reuse.
P304+P340	IF INHALED: Remove person to fresh air and keep comfortable for breathing.

Precautionary statement(s) Storage

P405	Store locked up.
P403+P233	Store in a well-ventilated place. Keep container tightly closed.

Precautionary statement(s) Disposal

P501 Dispose of contents/container to authorised hazardous or special waste collection point in accordance with any local regulation.

SECTION 3 Composition / information on ingredients

Substances

See section below for composition of Mixtures

Mixtures

CAS No	%[weight]	Name
64742-48-9.	80-90	naphtha petroleum, isoparaffin, hydrotreated
76-22-2	1-5	camphor
64742-47-8	1-5	Stoddard Solvent
64742-94-5	1-<2.5	solvent naphtha petroleum, heavy aromatic
12108-13-3	0.1-<1	manganese 2-methylcyclopentadienyl tricarbonyl
91-20-3	0.1-<0.25	naphthalene
Legend:	1. Classified by Chemwatch; 2. Classification drawn from C&L	. Classification drawn from HCIS; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 4. * EU IOELVs available

SECTION 4 First aid measures

Description of first aid measures

Eye Contact

If this product comes in contact with the eyes: Immediately hold eyelids apart and flush the eye continuously with running water.

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	 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. Continue flushing until advised to stop by the Poisons Information Centre or a doctor, or for at least 15 minutes. Transport to hospital or doctor without delay. 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.
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

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. For petroleum distillates

- In case of ingestion, gastric lavage with activated charcoal can be used promptly to prevent absorption decontamination (induced emesis or lavage) is controversial and should be considered on the merits of each individual case; of course the usual precautions of an endotracheal tube should be considered prior to lavage, to prevent aspiration.
- Individuals intoxicated by petroleum distillates should be hospitalized immediately, with acute and continuing attention to neurologic and cardiopulmonary function.
 Positive pressure ventilation may be pressure.
- Positive pressure ventilation may be necessary.
- Acute central nervous system signs and symptoms may result from large ingestions of aspiration-induced hypoxia.
- After the initial episode, individuals should be followed for changes in blood variables and the delayed appearance of pulmonary oedema and chemical pneumonitis. Such
 patients should be followed for several days or weeks for delayed effects, including bone marrow toxicity, hepatic and renal impairment Individuals with chronic pulmonary
 disease will be more seriously impaired, and recovery from inhalation exposure may be complicated.
- Gastrointestinal symptoms are usually minor and pathological changes of the liver and kidneys are reported to be uncommon in acute intoxications.
- Chlorinated and non-chlorinated hydrocarbons may sensitize the heart to epinephrine and other circulating catecholamines so that arrhythmias may occur.Careful consideration of this potential adverse effect should precede administration of epinephrine or other cardiac stimulants and the selection of bronchodilators.

BP America Product Safety & Toxicology Department

SECTION 5 Firefighting measures

Extinguishing media

- ▶ Foam.
- Dry chemical powder.
- BCF (where regulations permit).
- Carbon dioxide.
- Water spray or fog Large fires only.
- Do not use water jets.

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. Cool fire exposed containers with water spray from a protected location. If safe to do so, remove containers from path of fire. 		
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. 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) metal oxides other pyrolysis products typical of burning organic material. Contains low boiling substance: Closed containers may rupture due to pressure buildup under fire conditions. May emit poisonous fumes. May emit corrosive fumes. 		
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

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Minor Spills	 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. Place in a suitable, labelled container for waste disposal. Slippery when spilt.
Major Spills	 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. Increase ventilation. Stop leak if safe to do so. Contain spill with sand, earth or vermiculite. Collect recoverable product into labelled containers for recycling. Absorb remaining product with sand, earth or vermiculite. Collect solid residues and seal in labelled drums for disposal. Wash area and prevent runoff into drains. If contamination of drains or waterways occurs, advise emergency services. Slippery when spilt.

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

SECTION 7 Handling and storage

Precautions for safe handling

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Safe handling	The conductivity of this material may make it a static accumulator., A liquid is typically considered nonconductive if its conductivity 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 temperature, presence of contaminants, and anti-static additives can greatly influence the conductivity of a liquid. • Containers, even those that have been emptied, may contain explosive vapours. • Do NOT cut, drill, grind, weld or perform similar operations on or near containers. Containes 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 well with material to stay in contact with skin • Electrostatic discharge may be generated during pumping - this may result in fire. • Ensure electrical continuity by bonding and grounding (earthing) all equipment. • Restrict line velocity during pumping in order to avoid generation of electrostatic discharge (<=1 m/sec until fill pipe submerged to twice its diameter, then <= 7 m/sec). • Avoid splash filling. • Do NOT est compressed air for filling discharging or handling operations. • 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 est, drink or smoke. • Avoid somking, naked lights or ignition sources. • Avoid somking, naked handling recommendations containers. • Wore contact with incomp
Other information	 Store in original containers. Keep containers securely sealed. No smoking, naked lights or ignition sources. 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. Observe manufacturer's storage and handling recommendations contained within this SDS.

Conditions for safe storage, including any incompatibilities

Suitable container	▶ Metal can or drum
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	Packaging as recommended by manufacturer.
	Check all containers are clearly labelled and free from leaks.
Storage incompatibility	 Avoid reaction with oxidising agents Avoid strong acids, bases.

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	naphtha petroleum, isoparaffin, hydrotreated	Oil mist, refined mineral	5 mg/m3	Not Available	Not Available	Not Available
Australia Exposure Standards	camphor	Camphor, synthetic	2 ppm / 12 mg/m3	19 mg/m3 / 3 ppm	Not Available	Not Available
Australia Exposure Standards	Stoddard Solvent	White spirits	790 mg/m3	Not Available	Not Available	Not Available
Australia Exposure Standards	manganese 2-methylcyclopentadienyl tricarbonyl	Methylcyclopentadienyl manganese tricarbonyl (as Mn)	0.2 mg/m3	Not Available	Not Available	Not Available
Australia Exposure Standards	naphthalene	Naphthalene	10 ppm / 52 mg/m3	79 mg/m3 / 15 ppm	Not Available	Not Available

Ingredient	TEEL-1 TEEL-2			TEEL-3
naphtha petroleum, isoparaffin, hydrotreated	350 mg/m3	1,800 mg/m3		40,000 mg/m3
naphtha petroleum, isoparaffin, hydrotreated	1,100 mg/m3	1,800 mg/m3		40,000 mg/m3
Stoddard Solvent	300 mg/m3	1,800 mg/m3		29500** mg/m3
manganese 2-methylcyclopentadienyl tricarbonyl	0.3 mg/m3	0.6 mg/m3		6.9 mg/m3
naphthalene	15 ppm	83 ppm		500 ppm
Ingredient	Original IDLH		Revised IDI H	
naphtha petroleum, isoparaffin, hydrotreated	2,500 mg/m3		Not Available	
camphor	200 mg/m3		Not Available	
Stoddard Solvent	20,000 mg/m3		Not Available	
solvent naphtha petroleum, heavy aromatic	Not Available		Not Available	
manganese 2-methylcyclopentadienyl tricarbonyl	500 mg/m3		Not Available	
naphthalene	250 ppm		Not Available	

Exposure controls

Appropriate engineering controls	 CARE: Use of a quantity of this material in confined space or poorly ventilated area, where rapid build up of concentrate 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 be highly effective in protecting workers and will typically be independent of worker interactions to provide this high level The basic types of engineering controls are: 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 ven "adds" and "removes" air in the work environment. Ventilation can remove or dilute an air contaminant if designed proper 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. Local exhaust ventilation usually required. If risk of overexposure exists, wear approved respirator. Correct fit is essential protection. Supplied-air type respirator may be required in special circumstances. Correct fit is essential to ensure adequ An approved self contained breathing apparatus (SCBA) may be required in some situations. Provide adequate ventilation in warehouse or closed storage area. Air contaminants generated in the workplace possess velocities which, in turn, determine the "capture velocities" of fresh circulating air required to effectively remove the contain 	d atmosphere may occur, engineering controls can of protection. tilation that strategically rly. The design of a I to obtain adequate uate protection. s varying "escape" aminant.
	Type of Contaminant:	Air Speed:
	solvent, vapours, degreasing etc., evaporating from tank (in still air).	0.25-0.5 m/s (50-100 f/min.)
	aerosols, fumes from pouring operations, intermittent container filling, low speed conveyer transfers, welding, spray drift, plating acid fumes, pickling (released at low velocity into zone of active generation)	0.5-1 m/s (100-200 f/min.)
	direct spray, spray painting in shallow booths, drum filling, conveyer loading, crusher dusts, gas discharge (active	1-2.5 m/s (200-500

grinding, abrasive blasting, tumbling, high speed wheel generated dusts (released at high initial velocity into zone of very high rapid air motion).

2.5-10 m/s (500-2000 f/min.)

within each range the appropriate value depends on:		
Lower end of the range	Upper end of the range	
1: Room air currents minimal or favourable to capture	1: Disturbing room air currents	
2: Contaminants of low toxicity or of nuisance value only.	2: Contaminants of high toxicity	
3: Intermittent, low production.	3: High production, heavy use	
4: Large hood or large air mass in motion	4: Small hood-local control only	

Simple theory shows that air velocity falls rapidly with distance away from the opening of a simple extraction pipe. Velocity generally decreases with the square of distance from the extraction point (in simple cases). Therefore the air speed at the extraction point should be adjusted, accordingly, after reference to distance from the contaminating source. The air velocity at the extraction fan, for example, should be a minimum of 1-2 m/s (200-400 f/min) for extraction of solvents generated in a tank 2 meters distant from the extraction point. Other mechanical considerations, producing performance deficits within the extraction apparatus, make it essential that theoretical air velocities are multiplied by factors of 10 or more when extraction systems are installed or used.

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. In the event of chemical exposure, begin eye irrigation immediately and remove contact lens as soon as practicable. Lens should be removed at the first signs of eye redness or irritation - lens should be removed in a clean environment only after workers have washed hands thoroughly. [CDC NIOSH Current Intelligence Bulletin 59], [AS/NZS 1336 or national equivalent]
Skin protection	See Hand protection below
Hands/feet protection	 Wear schemical protective gloves, e.g. PVC. Wear stelly footwear or safety gumboots, e.g. Rubber NOTE: The material may produce skin sensitisation in predisposed individuals. Care must be taken, when removing gloves and other protective equipment, to avoid all possible skin contact. Contaminated leather items, such as shoes, bells and watch-bands should be removed and destroyed. 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 making a final choice. Personal hygiene is a key element of effective hand care. Gloves must only be worn on clean hands. After using gloves, hands should be washed and dried thoroughly. Application of a non-perfumed moisturiser is recommended. Suitability and durability of glove type is dependent on usage. Important factors in the selection of gloves include: - chemical resistance of glove material, - glove thickness and - devicerity Select gloves tested to a relevant standard (e.g. Europe EN 374, US F739, ASNZS 2161.1 or national equivalent), - When proteoged 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, ASNZ 2161.1.0.1 or national equivalent) is recommended. - Some glove polymer types are less affected by movement and this should be taken into account when considering gloves for long-term use. - Contaminated gloves should be replaced. - Some glove polymer types are less affected by mo
Body protection	See Other protection below
Other protection	 Overalls. P.V.C apron. Barrier cream. Skin cleansing cream. Eve wash unit

Recommended material(s)

GLOVE SELECTION INDEX

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 *computer-generated* selection:

21696 Octane Booster

21000 Obtaile Bood

Material	СРІ
TEFLON	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 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)

- 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.
- Cartridge performance is affected by humidity. Cartridges should be changed after 2 hr of continuous use unless it is determined that the humidity is less than 75%, in which case, cartridges can be used for 4 hr. Used cartridges should be discarded daily, regardless of the length of time used

SECTION 9 Physical and chemical properties

Information on basic physical and chemical properties

Appearance Clear yellow liquid with characteristic odour; does not mix with water.

Physical state	Liquid	Relative density (Water = 1)	0.81 @15C
Odour	Characteristic	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)	<7 @40C
Initial boiling point and boiling range (°C)	Not Available	Molecular weight (g/mol)	Not Applicable
Flash point (°C)	>63	Taste	Not Available
Evaporation rate	Not Available	Explosive properties	Not Available
Flammability	Combustible.	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	Immiscible	pH as a solution (Not Available%)	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	See section 5

Continued...

SECTION 11 Toxicological information

Information on toxicological ef	fects	
Inhaled	Inhalation of vapours or aerosols (mists, tumes), generated by the material during the course of normal handling, may be harmful. 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. There is some evidence to suggest that the material can cause respiratory irritation in some persons. The body's response to such irritation can cause further lung damage. 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. Nerve damage can be caused by some non-ring hydrocarbons. Symptoms are temporary, and include weakness, tremors, increased saliva, some convulsions, excessive tears with discolouration and inco-ordination lasting up to 24 hours. 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.	
Ingestion	Accidental ingestion of the material may be damaging to the health of the individual. Swallowing of the liquid may cause aspiration into the lungs with the risk of chemical pneumonitis; serious consequences may result. (ICSC13733) Isoparaffinic hydrocarbons cause temporary lethargy, weakness, inco-ordination and diarrhoea. Ingestion of petroleum hydrocarbons can irritate the pharynx, oesophagus, stomach and small intestine, and cause swellings and ulcers of the mucous. Symptoms include a burning mouth and throat; larger amounts can cause nausea and vomiting, narcosis, weakness, dizziness, slow and shallow breathing, abdominal swelling, upconsciousness and convulsions	
Skin Contact	Repeated exposure may cause skin cracking, flaking or drying following normal handling and use. Skin contact with the material may damage the health of the individual; systemic effects may result following absorption. There is some evidence to suggest that this material can cause inflammation of the skin on contact in some persons. Skin exposure to isoparaffins may produce slight to moderate irritation in animals and humans. Rare sensitisation reactions in humans have occurred. Workers sensitised to naphthalene and related compounds show an inflammation of the skin with scaling and reddening. Some individuals show an allergic reaction. Open cuts, abraded or irritated skin should not be exposed to this material The material may accentuate any pre-existing dermatitis condition 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.	
Eye	If applied to the eyes, this material causes severe eye damage. Instillation of isoparaffins into rabbit eyes produces only slight irritation. Direct eye contact with petroleum hydrocarbons can be painful, and the corneal epithelium may be temporarily damaged. Aromatic species can cause irritation and excessive tear secretion.	
Chronic	Skin contact with the material is more likely to cause a sensitisation reac Substance accumulation, in the human body, may occur and may cause There has been some concern that this material can cause cancer or mu Prolonged or repeated skin contact may cause drying with cracking, irrita Constant or exposure over long periods to mixed hydrocarbons may proo and anaemia, and reduced liver and kidney function. Skin exposure may	tion in some persons compared to the general population. some concern following repeated or long-term occupational exposure. tations but there is not enough data to make an assessment. tion and possible dermatitis following. duce stupor with dizziness, weakness and visual disturbance, weight loss result in drying and cracking and redness of the skin.
21696 Octane Booster	TOXICITY Dermal (None) LD50: >2000 mg/kg*[2] Inhalation (None) LC50: 14.87 mg/l/4h(vapours)*[2] Oral (None) LD50: >2000 mg/kg*[2]	IRRITATION Not Available
naphtha petroleum, isoparaffin, hydrotreated	TOXICITY Dermal (rabbit) LD50: >1900 mg/kg ^[1] Inhalation(Rat) LC50; >4.42 mg/L4h ^[1] Oral (Rat) LD50; >4500 mg/kg ^[1]	IRRITATION Eye: no adverse effect observed (not irritating) ^[1] Skin: adverse effect observed (irritating) ^[1]

camphor	TOXICITY dermal (rat) LD50: >2000 mg/kg ^[1] Oral (Rat) LD50; >5000 mg/kg ^[1]	IRRITATION Not Available
Stoddard Solvent	TOXICITY Dermal (rabbit) LD50: >3000 mg/kg ^[1] Inhalation(Rat) LC50; >5.5 mg/l4h ^[1] Oral (Rat) LD50; >5000 mg/kg ^[1]	IRRITATION Eye (hmn) 470 ppm/15m irrit. Eye (rabbit) 500 mg/24h moderate Eye: no adverse effect observed (not irritating) ^[1] Skin: adverse effect observed (irritating) ^[1]
	τοχιριτγ	Skin: no adverse effect observed (not irritating) ^[1]
solvent naphtha petroleum, heavy aromatic	Dermal (rabbit) LD50: >2000 mg/kg ^[2] Inhalation(Rat) LC50; >0.003 mg/L4h ^[1]	Eye (rabbit): Irritating Eye: no adverse effect observed (not irritating) ^[1]

	Oral (Rat) LD50; 512 mg/kg ^[1]	Skin: adverse effect observed (irritating) ^[1]
	τοχιςιτγ	IRRITATION
manganese	Dermal (rabbit) LD50: 140 mg/kg ^[2]	Skin (rabbit): 100 mg/24h - mild
2-methylcyclopentadienyl	Inhalation(Rat) LC50: 0.055 mg/L4h ^[2]	
tricarbonyi	Oral (Rat) D50: 9 mg/kg ^[2]	
	ΤΟΧΙΟΙΤΥ	IRRITATION
	dermal (rat) LD50: >2500 mg/kg ^[2]	Eye (rabbit): 100 mg - mild
naphthalene	Inhalation(Rat) LC50; >0.4 mg/l4h ^[1]	Skin (rabbit):495 mg (open) - mild
	Oral (Rat) LD50; 490 mg/kg ^[2]	
Legend:	1. Value obtained from Europe ECHA Registered Substances - Acu specified data extracted from RTECS - Register of Toxic Effect of c	ite toxicity 2.* Value obtained from manufacturer's SDS. Unless otherwise hemical Substances
NAPHTHA PETROLEUM, ISOPARAFFIN, HYDROTREATED	, No significant acute toxicological data identified in literature searc	ch.
Самрног	 for (+/-) -camphor: [CAS No. 21368-68-3] Nil reported The following information refers to contact allergens as a group a contact allergies quickly manifest themselves as contact eczema eczema involves a cell-mediated (T lymphocytes) immune reaction involve antibody-mediated immune reactions. The significance of distribution of the substance and the opportunities for contact with distributed can be a more important allergen than one with strong clinical point of view, substances are noteworthy if they produce a Asthma-like symptoms may continue for months or even years af known as reactive airways dysfunction syndrome (RADS) which or criteria for diagnosing RADS include the absence of previous airw asthma-like symptoms within minutes to hours of a documented or reversible airflow pattern on lung function tests, moderate to sever minimal lymphocytic inflammation, without eosinophila. RADS (or related to the concentration of and duration of exposure to the irri occurs as a result of exposure due to high concentrations of irrita ceases. The disorder is characterized by difficulty breathing, coug Adverse reactions to fragrances in perfumes and fragranced cost sensitivity to light, immediate contact reactions, and pigmented callergy is a lifelong condition, so symptoms may occur on re-expo significant impairment of quality of life and potential consequence of the perfume contains a sensitizing component, intolerance to proughing, phlegm, wheezing, chest tightness, headache, shorthe other respiratory diseases. Perfumes can induce excess reactivity through a carbon filter mask had no protective effect. Occupational asthma caused by perfume substances, such as is persistent symptoms, even though the exposure is below occupa important objective of public health risk management. Hands: Contact sensitization may be the primary cause of hand eczema is a disease involving many factors, and the clinical signip be clear. Underarm: Skin inflammation of the armpits	nd may not be specific to this product. , more rarely as uticaria or Quincke's oedema. The pathogenesis of contact on of the delayed type. Other allergic skin reactions, e.g. contact uticaria, the contact allergen is not simply determined by its sensitisation potential: the in it are equally important. A weakly sensitising substance which is widely ger sensitising potential with which few individuals come into contact. From a nallergic test reaction in more than 1% of the persons tested. ter exposure to the material ends. This may be due to a non-allergic condition can occur after exposure to high levels of highly irritating compound. Main vays disease in a non-atopic individual, with sudden onset of persistent exposure to the irritant. Other criteria for diagnosis of RADS include a re bronchial hyperreactivity on methacholine challenge testing, and the lack of rasthma) following an irritating inhalation is an infrequent disorder with rates tating substance. On the other hand, industrial bronchittis is a disorder that ting substance. On the other hand, industrial bronchittis is a disorder that ting substance infer and connubial contact dermatitis, niticat dermatitis. Airborne and connubial contact dermatitis, products include allergic contact dermatitis, inritant contact dermatitis, so therass for work. effumes by inhalation may occur. Symptoms may include general unwellness, ss ob breath with exertion, acute respiratory illness, hayfever, asthma and y of the airway without producing allergy or airway obstruction. Breathing onamyl acetate, limonene, cinnamaldehyde and benzaldehyde, tend to give tional exposure limits. Prevention of contact sensitization to fragrances is an eczema or a complication of irritant or atopic hand eczema. However hand ficance of fragrance contact allergy in severe, chronic hand eczema may not affume hedodrants and, if the reaction is severe, it may spread down the a skin specialist, a history of such first-time symptoms was significantly to akom specialist,

	without the requirement of an enzyme. For prehaptens, it is possible to prevent activation or exposure during handling and storage of the ingredit are used, care should be taken that they will not be a Prehaptens: Most terpenes with oxidisable allylic po- oxidation products that are formed, the oxidized prod lavender oil increased the potential for sensitization. Prohaptens: Compounds that are bioactivated in the being activated cannot be avoided by outside measu substances. Various enzymes play roles in both acti grouped into chemical classes based on knowledge QSAR prediction: Prediction of sensitization activity and prohaptens.	utside the body to a certain extent by ents and the final product, and by the activated themselves, and thereby for sitions can be expected to self-oxidis ducts will have differing levels of sens e skin and thereby form haptens are re- ures. Activation processes increase the vating and deactivating prohaptens. So of xenobiotic bioactivation reactions, of these substances is complex, espe	different measures, for example, prevention of air addition of suitable antioxidants. When antioxidants m new sensitisers. e on air exposure. Depending on the stability of the itization potential. Tests shows that air exposure of eferred to prohaptens. The possibility of a prohapten he risk for cross-reactivity between fragrance skin-sensitizing prohaptens can be recognized and clinical observations and/or studies of sensitization. ecially for those substances that can act both as pre-
MANGANESE 2-METHYLCYCLOPENTADIENYL TRICARBONYL	The toxicity of metal carbonyls is due to toxicity of ca Exposure occurs by inhalation, or for liquid metal ca Metal carbonyls are toxic by skin contact, inhalation carboxyhemoglobin, which prevents the binding of C For manganese 2-methylcyclopentadienyl tricarbony aberrations occurred, but only in the presence of me toxicity. NOAEL (inhalation) 6.2 mg/m3 (rats and mice)* * We	arbon monoxide, the metal, and becar rbonyls by ingestion or due to the goc or ingestion, in part because of their i 22 I (MMT): Animal testing showed that tabolic activation. MMT does not see orksafe Australia	use of the volatility and instability of the complexes. I fat solubility by skin resorption. ability to carbonylate haemoglobin to give the substance did not cause mutations. Chromosome m to cause genetic, developmental or reproductive
NAPHTHALENE	The material may be irritating to the eye, with prolonged contact causing inflammation. Repeated or prolonged exposure to irritants may produce conjunctivitis.		
NAPHTHA PETROLEUM, ISOPARAFFIN, HYDROTREATED & SOLVENT NAPHTHA PETROLEUM, HEAVY AROMATIC	Animal studies indicate that normal, branched and cyclic paraffins are absorbed from the 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 than iso- or cyclo-paraffins. The major classes of hydrocarbons are well absorbed into the gastrointestinal tract in various species. In many cases, the hydrophobic hydrocarbons are ingested in association with fats in the diet. Some hydrocarbons may appear unchanged as in the lipoprotein particles in the gut lymph, but most hydrocarbons partly separate from fats and undergo metabolism in the gut cell. The gut cell may play a major role in determining the proportion of hydrocarbon that becomes available to be deposited unchanged in peripheral tissues such as in the body fat stores or the liver.		
NAPHTHA PETROLEUM, ISOPARAFFIN, HYDROTREATED & STODDARD SOLVENT & SOLVENT NAPHTHA PETROLEUM, HEAVY AROMATIC	For petroleum: This product contains benzene, which can cause acute myeloid leukaemia, and n-hexane, which can be metabolized to compounds which are toxic to the nervous system. This product contains toluene, and animal studies suggest high concentrations of toluene lead to hearing loss. This product contains ethyl benzene and naphthalene, from which animal testing shows evidence of tumour formation. Cancer-causing potential: Animal testing shows inhaling petroleum causes tumours of the liver and kidney; these are however not considered to be relevant in humans. Mutation-causing potential: Most studies involving gasoline have returned negative results regarding the potential to cause mutations, including all recent studies in living human subjects (such as in petrol service station attendants). Reproductive toxicity: Animal studies show that high concentrations of toluene (>0.1%) can cause developmental effects such as lower birth weight and developmental toxicity to the nervous system of the foetus. Other studies show no adverse effects on the foetus. Human effects: Prolonged or repeated contact may cause defatting of the skin which can lead to skin inflammation and may make the skin more susceptible to irritation and penetration by other materials.		
MANGANESE 2-METHYLCYCLOPENTADIENYL TRICARBONYL & NAPHTHALENE	The material may cause skin irritation after prolonge of vesicles, scaling and thickening of the skin.	d or repeated exposure and may pro	duce on contact skin redness, swelling, the production
Acute Toxicity	×	Carcinogenicity	×
Skin Irritation/Corrosion	×	Reproductivity	×
Serious Eye Damage/Irritation	×	STOT - Single Exposure	¥
Respiratory or Skin sensitisation	v	STOT - Repeated Exposure	×

SECTION 12 Ecological information

Mutagenicity

X

	Endpoint	Test Duration (hr)	Species	Value	Source
21696 Octane Booster	Not Available	Not Available	Not Available	Not Available	Not Available
	Endpoint	Test Duration (hr)	Species	Value	Source
naphtha petroleum,	EC50(ECx)	96h	Algae or other aquatic plants	64mg/l	2
soparanni, nyurotreateu	EC50	96h	Algae or other aquatic plants	64mg/l	2
	Endpoint	Test Duration (hr)	Species	Value	Source
	LC50	96h	Fish	33.25mg/l	2
camphor	EC50	72h	Algae or other aquatic plants	0.3mg/l	2
	EC50	48h	Crustacea	4.23mg/l	2
	5050	001		6 0E1mg/	2

Aspiration Hazard

Legend:

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Continued...

21696 Octane Booster

	NOEC(ECx)	72h	Algae or other aquatic plants	0.032mg/l	2
	Endpoint	Test Duration (hr)	Species	Value	Source
	NOEC(ECx)	3072h	Fish	1mg/l	1
Stoddard Solvent	NOEC(ECx)	720h	Crustacea	0.024mg/l	2
	LC50	96h	Fish	0.14mg/l	2
	EC50	96h	Algae or other aquatic plants	0.277mg/l	2
	Endpoint	Test Duration (hr)	Species	Value	Source
	EC50(ECx)	48h	Crustacea	0.95mg/l	1
solvent naphtha petroleum.	LC50	96h	Fish	0.58mg/l	2
heavy aromatic	EC50	72h	Algae or other aquatic plants	<1mg/l	1
	EC50	48h	Crustacea	0.95mg/l	1
	EC50	96h	Algae or other aquatic plants	1mg/l	2
		Test Describer (Le)	0		Course
	Endpoint	lest Duration (nr)	Species	Value	Source
manganese	Endpoint LC50	96h	Fish	0.21mg/l	2
manganese 2-methylcyclopentadienyl	Endpoint LC50 EC50	96h 48h	Species Fish Crustacea	0.21mg/l 0.83mg/l	2 2
manganese 2-methylcyclopentadienyl tricarbonyl	Endpoint LC50 EC50 EC50	96h 96h 96h	Species Fish Crustacea Algae or other aquatic plants	Value 0.21mg/l 0.83mg/l >0.46mg/l	2 2 2 2
manganese 2-methylcyclopentadienyl tricarbonyl	Endpoint LC50 EC50 EC50 NOEC(ECx)	Pest Duration (hr) 96h 48h 96h 48h	Species Fish Crustacea Algae or other aquatic plants Algae or other aquatic plants	Value 0.21mg/l 0.83mg/l >0.46mg/l 0.07mg/l	2 2 2 2 2
manganese 2-methylcyclopentadienyl tricarbonyl	Endpoint LC50 EC50 EC50 NOEC(ECx) Endpoint	1est Duration (hr) 96h 48h 96h 48h Test Duration (hr)	Species Fish Crustacea Algae or other aquatic plants Algae or other aquatic plants Species	Value 0.21mg/l 0.83mg/l >0.46mg/l 0.07mg/l	2 2 2 2 2 Source
manganese 2-methylcyclopentadienyl tricarbonyl	Endpoint LC50 EC50 EC50 NOEC(ECx) Endpoint BCF	1est Duration (hr) 96h 48h 96h 48h Test Duration (hr) 1344h	Species Fish Crustacea Algae or other aquatic plants Algae or other aquatic plants Species Fish	Value 0.21mg/l 0.83mg/l >0.46mg/l 0.07mg/l Value 23-146	2 2 2 2 2 2 5 5 0 0 7
manganese 2-methylcyclopentadienyl tricarbonyl	Endpoint LC50 EC50 EC50 NOEC(ECx) Endpoint BCF NOEC(ECx)	Test Duration (hr) 96h 48h 96h 48h 1000 mm	Species Fish Crustacea Algae or other aquatic plants Algae or other aquatic plants Species Fish Fish Fish	Value 0.21mg/l 0.83mg/l >0.46mg/l 0.07mg/l Value 23-146 0.013mg/L	2 2 2 2 2 2 5 5 5 0 0 7 4
manganese 2-methylcyclopentadienyl tricarbonyl naphthalene	Endpoint LC50 EC50 EC50 NOEC(ECx) Endpoint BCF NOEC(ECx) LC50	Test Duration (hr) 96h 48h 96h 48h 1344h 48h 96h	Species Fish Crustacea Algae or other aquatic plants Algae or other aquatic plants Species Fish Fish Fish Fish Fish Fish	Value 0.21mg/l 0.83mg/l >0.46mg/l 0.07mg/l 23-146 0.013mg/L 0.51mg/l	Source 2 2 2 2 2 2 2 2 2 2 2 2 2 4 4
manganese 2-methylcyclopentadienyl tricarbonyl naphthalene	Endpoint LC50 EC50 EC50 NOEC(ECx) Endpoint BCF NOEC(ECx) LC50 EC50	Test Duration (hr) 96h 48h 96h 48h 7Est Duration (hr) 1344h 48h 96h 72h	Species Fish Crustacea Algae or other aquatic plants Algae or other aquatic plants Fish Fish Fish Fish Algae or other aquatic plants	Value 0.21mg/l 0.83mg/l >0.46mg/l 0.07mg/l 23-146 0.013mg/L 0.51mg/l ~0.4~0.5mg/l	Source 2 2 2 2 2 2 7 4 4 2 2
manganese 2-methylcyclopentadienyl tricarbonyl naphthalene	Endpoint LC50 EC50 NOEC(ECx) Endpoint BCF NOEC(ECx) LC50 EC50 EC50	Test Duration (hr) 96h 48h 96h 48h Test Duration (hr) 1344h 48h 96h 48h 96h 48h	Species Fish Crustacea Algae or other aquatic plants Algae or other aquatic plants Species Fish Fish Fish Algae or other aquatic plants Crustacea	Value 0.21mg/l 0.83mg/l >0.46mg/l 0.07mg/l 23-146 0.013mg/L 0.51mg/l -0.4~0.5mg/l 1.09-3.4mg/l	Source 2 2 2 2 2 2 3 Source 7 4 4 2 4 4 2 4 4

Harmful to aquatic organisms, may cause long-term adverse effects in the aquatic environment. **DO NOT** discharge into sewer or waterways.

Persistence and degradability

Ingredient	Persistence: Water/Soil	Persistence: Air
camphor	HIGH	HIGH
naphthalene	HIGH (Half-life = 258 days)	LOW (Half-life = 1.23 days)

Bioaccumulative potential

Ingredient	Bioaccumulation
camphor	LOW (LogKOW = 2.74)
Stoddard Solvent	LOW (BCF = 159)
solvent naphtha petroleum, heavy aromatic	LOW (BCF = 159)
naphthalene	HIGH (BCF = 18000)

Mobility in soil

Ingredient	Mobility
camphor	LOW (KOC = 106)
naphthalene	LOW (KOC = 1837)

SECTION 13 Disposal considerations

Waste treatment methods

Product / Packaging disposal	 Containers may still present a chemical hazard/ danger when empty. Return to supplier for reuse/ recycling if possible. Otherwise: If container can not be cleaned sufficiently well to ensure that residuals do not remain or if the container cannot be used to store the same product, then puncture containers, to prevent re-use, and bury at an authorised landfill. Where possible retain label warnings and SDS and observe all notices pertaining to the product. 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. 	

 Recycle wherever possible or consult manufacturer for recycling options. 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 14 Transport information

Labels Required	
	COMBUSTIBLE LIQUID, regulated for storage purposes only
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 / GGVSee): NOT REGULATED FOR TRANSPORT OF DANGEROUS GOODS

Transport in bulk according to Annex II of MARPOL and the IBC code

Not Applicable

Transport in bulk in accordance with MARPOL Annex V and the IMSBC Code

Product name	Group
naphtha petroleum, isoparaffin, hydrotreated	Not Available
camphor	Not Available
Stoddard Solvent	Not Available
solvent naphtha petroleum, heavy aromatic	Not Available
manganese 2-methylcyclopentadienyl tricarbonyl	Not Available
naphthalene	Not Available

Transport in bulk in accordance with the ICG Code

Product name	Ship Type
naphtha petroleum, isoparaffin, hydrotreated	Not Available
camphor	Not Available
Stoddard Solvent	Not Available
solvent naphtha petroleum, heavy aromatic	Not Available
manganese 2-methylcyclopentadienyl tricarbonyl	Not Available
naphthalene	Not Available

SECTION 15 Regulatory information

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

l	naphtha petroleum, isoparaffin, hydrotreated is found on the following regulatory lists		
	Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals	Chemical Footprint Project - Chemicals of High Concern List	
	Australian Inventory of Industrial Chemicals (AIIC)	International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Monographs	
l	camphor is found on the following regulatory lists		
	Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) -	Australian Inventory of Industrial Chemicals (AIIC)	
	Schedule 5	International WHO List of Proposed Occupational Exposure Limit (OEL) Values for	
	Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Schedule 6	Manufactured Nanomaterials (MNMS)	
l	Stoddard Solvent is found on the following regulatory lists		
	Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals	International Agency for Research on Cancer (IARC) - Agents Classified by the IARC	
	Australian Inventory of Industrial Chemicals (AIIC)	Monographs	
	Chemical Footprint Project - Chemicals of High Concern List	International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Monographs - Group 1: Carcinogenic to humans	
l	solvent naphtha petroleum, heavy aromatic is found on the following regulatory lists		
	Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals	International Agency for Research on Cancer (IARC) - Agents Classified by the IARC	
	Australian Inventory of Industrial Chemicals (AIIC)	Monographs	

manganese 2-methylcyclopentadienyl tricarbonyl is found on the following regulatory lists

Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) -Schedule 6

naphthalene is found on the following regulatory lists

Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals Australia Model Work Health and Safety Regulations - Hazardous chemicals (other than lead) requiring health monitoring

Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Schedule 10 / Appendix C

Australian Inventory of Industrial Chemicals (AIIC)

Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) -Schedule 7 Australian Inventory of Industrial Chemicals (AIIC)

Chemical Footprint Project - Chemicals of High Concern List International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Monographs International Agency for Research on Cancer (IARC) - Agents Classified by the IARC

Monographs - Group 2B: Possibly carcinogenic to humans International WHO List of Proposed Occupational Exposure Limit (OEL) Values for Manufactured Nanomaterials (MNMS)

National Inventory Status

National Inventory	Status	
Australia - AIIC / Australia Non-Industrial Use	Yes	
Canada - DSL Yes		
Canada - NDSL No (naphtha petroleum, isoparaffin, hydrotreated; Stoddard Solvent; solvent naphtha petroleum, heavy aromatic; manganese 2-methylcyclopentadienyl tricarbonyl; naphthalene)		
China - IECSC	Yes	
Europe - EINEC / ELINCS / NLP	Yes	
Japan - ENCS	No (naphtha petroleum, isoparaffin, hydrotreated; solvent naphtha petroleum, heavy aromatic)	
Korea - KECI	Yes	
New Zealand - NZIoC	Yes	
Philippines - PICCS	Yes	
USA - TSCA	Yes	
Taiwan - TCSI	Yes	
Mexico - INSQ	Yes	
Vietnam - NCI	No (manganese 2-methylcyclopentadienyl tricarbonyl)	
Russia - FBEPH	Yes	
Legend:	Yes = All CAS declared ingredients are on the inventory No = One or more of the CAS listed ingredients are not on the inventory. These ingredients may be exempt or will require registration.	

SECTION 16 Other information

Revision Date	22/02/2022
Initial Date	22/02/2022

SDS Version Summary

Version	Date of Update	Sections Updated
2.1	22/02/2022	Acute Health (eye), Acute Health (inhaled), Acute Health (skin), Acute Health (swallowed), Advice to Doctor, Appearance, Chronic Health, Classification, Disposal, Environmental, Exposure Standard, Fire Fighter (extinguishing media), Fire Fighter (fire/explosion hazard), Spills (major), Spills (minor), Storage (storage incompatibility), Toxicity and Irritation (Toxicity Figure), Toxicity and Irritation (Other), Use

Other information

Ingredients with multiple cas numbers

Name	CAS No	
naphtha petroleum, isoparaffin, hydrotreated	64742-48-9., 101795-02-2., 64771-72-8.	
camphor	76-22-2, 21368-68-3, 464-49-3, 464-48-2	
Stoddard Solvent	8052-41-3., 64742-47-8	
solvent naphtha petroleum, heavy aromatic	64742-94-5, 1189173-42-9	

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

ES: Exposure Standard 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 AIIC: Australian Inventory of Industrial Chemicals DSL: Domestic Substances List NDSL: Non-Domestic Substances List IECSC: Inventory of Existing Chemical Substance in China EINECS: European INventory of Existing Commercial chemical Substances ELINCS: European List of Notified Chemical Substances NLP: No-Longer Polymers ENCS: Existing and New Chemical Substances Inventory KECI: Korea Existing Chemicals Inventory NZIOC: New Zealand Inventory of Chemicals PICCS: Philippine Inventory of Chemicals and Chemical Substances TSCA: Toxic Substances Control Act TCSI: Taiwan Chemical Substance Inventory INSQ: Inventario Nacional de Sustancias Químicas NCI: National Chemical Inventory FBEPH: Russian Register of Potentially Hazardous Chemical and Biological Substances

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