CRC Industries (CRC Industries New Zealand)

Chemwatch: **4546-12** Version No: **8.1.1.1**

Safety Data Sheet according to WHS and ADG requirements

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

Product Identifier

Product name	CRC 3013 Soft Seal / Panel Seal (Aerosol) (NZ)			
Synonyms	spray-on rust corrosion inhibitor; CRX3013			
Proper shipping name	AEROSOLS			
Other means of identification	Not Available			

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

Relevant identified uses	Temporary rust preventative coating. Application is by spray atomisation from a hand held aerosol pack The use of a quantity of material in an unventilated or confined space may result in increased exposure and an irritating atmosphere developing. Before starting consider control of exposure by mechanical ventilation.
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Details of the supplier of the safety data sheet

Registered company name	CRC Industries (CRC Industries New Zealand)		
Address	10 Highbrook Drive East Tamaki Auckland New Zealand		
Telephone	+64 9 272 2700		
Fax	+64 9 274 9696		
Website	www.crc.co.nz		
Email	customerservices@crc.co.nz		

Emergency telephone number

Association / Organisation	CRC Industries (CRC Industries New Zealand)			
Emergency telephone numbers	NZ Poisons Centre 0800 POISON (0800 764 766)			
Other emergency telephone numbers	111 (NZ Emergency Services)			

SECTION 2 HAZARDS IDENTIFICATION

Poisons Schedule	Not Applicable			
Classification [1]	Flammable Aerosols Category 1, Specific target organ toxicity - single exposure Category 3 (narcotic effects)			
Legend:	1. Classified by Chernwatch; 2. Classification drawn from HCIS; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI			
abel elements				
Hazard pictogram(s)				
SIGNAL WORD	DANGER			
azard statement(s)				
H222	Extremely flammable aerosol.			
H336	May cause drowsiness or dizziness.			
AUH044	Risk of explosion if heated under confinement.			
recautionary statement(s) Pre	vention			
P210	Keep away from heat/sparks/open flames/hot surfaces No smoking.			
P211	Do not spray on an open flame or other ignition source.			
P251	Pressurized container: Do not pierce or burn, even after use.			
P271	Use only outdoors or in a well-ventilated area.			
P261	Avoid breathing mist/vapours/spray.			

Chemwatch Hazard Alert Code: 3

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Precautionary statement(s) Response

P312	P312 Call a POISON CENTER or doctor/physician if you feel unwell.			
P304+P340	INHALED: Remove victim to fresh air and keep at rest in a position comfortable for breathing.			
Precautionary statement(s) Sto				

P405	pre locked up.		
P410+P412	Protect from sunlight. Do not expose to temperatures exceeding 50 °C/122 °F.		
P403+P233	Store in a well-ventilated place. Keep container tightly closed.		

Precautionary statement(s) Disposal

P501 Disp

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			
73513-42-5	30-60	isohexanes			
Various	30-60	liquid hydrocarbons			
Not Available	1-10	surfactant			
68476-85-7.	10-30	LPG (liquefied petroleum gas)			
Not Available		NOTE: Manufacturer has supplied full ingredient			
Not Available		information to allow CHEMWATCH assessment.			

SECTION 4 FIRST AID MEASURES

Description of first aid measures

Eye Contact	 If aerosols come in contact with the eyes: Immediately hold the eyelids apart and flush the eye continuously for at least 15 minutes 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. 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 solids or aerosol mists are deposited upon the skin: Flush skin and hair with running water (and soap if available). Remove any adhering solids with industrial skin cleansing cream. DO NOT use solvents. Seek medical attention in the event of irritation.
Inhalation	 If aerosols, fumes or combustion products are inhaled: Remove to fresh air. 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. If breathing is shallow or has stopped, ensure clear airway and apply resuscitation, 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	 Not considered a normal route of entry. Avoid giving milk or oils. Avoid giving alcohol.

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.

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

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

SECTION 5 FIREFIGHTING MEASURES

SMALL FIRE:
Water spray, dry chemical or CO2
LARGE FIRE:
Water spray or fog.

Special hazards arising from the substrate or mixture

Avoid contamination with oxidising agents i.e. nitrates, oxidising acids, chlorine bleaches, pool chlorine etc. as ignition may result				
 Alert Fire Brigade and tell them location and nature of hazard. May be violently or explosively reactive. Wear breathing apparatus plus protective gloves. Prevent, by any means available, spillage from entering drains or water course. If safe, switch off electrical equipment until vapour fire hazard removed. Use water delivered as a fine spray to control fire and cool adjacent area. 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. Equipment should be thoroughly decontaminated after use. 				
 Liquid and vapour are highly flammable. Severe fire hazard when exposed to heat or flame. Vapour forms an explosive mixture with air. Severe explosion hazard, in the form of vapour, when exposed to flame or spark. Vapour may travel a considerable distance to source of ignition. Heating may cause expansion or decomposition with violent container rupture. Aerosol cans may explode on exposure to naked flames. Rupturing containers may rocket and scatter burning materials. Hazards may not be restricted to pressure effects. May emit acrid, poisonous or corrosive fumes. On combustion, may emit toxic fumes of carbon monoxide (CO). 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). Kupturing containers of carbon monoxide (CO). May emit acrid smoke. May emit acrid smoke. May emit acrid smoke. Mists containing combustible materials may be explosive. Combustion products include: carbon dioxide (CO2) other pyrolysis products typical of burning organic material. Contains low boiling substance: Closed containers may rupture due to pressure buildup under fire conditions. 				
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

Slippery when spilt. Clean up all spills immediate Avoid breathing vapours and					
Minor Spills Wear protective clothing, im Shut off all possible sources Wipe up. If safe, damaged cans should be Undamaged cans should be 	pervious gloves of ignition and ld be placed in	s and safety glass increase ventilation a container outdoor	on.	gnition sources, unt	il pressure has dissipated.
Chemical Class: aliphatic hydrod For release onto land: recomme		listed in order of p	oriority.		
SORBENT TYPE	RANK	APPLICATION	COLLECTION	LIMITATIONS	
LAND SPILL - SMALL					
cross-linked polymer - particu	late 1	shovel	shovel	R, W, SS	
cross-linked polymer - pillow	1	throw	pitchfork	R, DGC, RT	
wood fiber - pillow	2	throw	pitchfork	R, P, DGC, RT	
Major Spills treated wood fibre- pillow	2	throw	pitchfork	DGC, RT	
sorbent clay - particulate	3	shovel	shovel	R, I, P	
foamed glass - pillow	3	throw	pitchfork	R, P, DGC, RT	
LAND SPILL - MEDIUM					
cross-linked polymer - particul	ate 1	blower	skiploader	R,W, SS	
cross-linked polymer - pillow	2	throw	skiploader	R, DGC, RT	
sorbent clay - particulate	3	blower	skiploader	R, I, P	

polypropylene - particulate	3	blower	skiploader	W, SS, DGC				
expanded mineral - particulate	4	blower	skiploader	R, I, W, P, DGC				
polypropylene - mat	4	throw	skiploader	DGC, RT				
Legend								
DGC: Not effective where ground cover is dense R; Not reusable								
								I: Not incinerable
P: Effectiveness reduced when rainy								
RT:Not effective where terrain is rugge								
SS: Not for use within environmentally								
W: Effectiveness reduced when windy								
Reference: Sorbents for Liquid Hazar	Reference: Sorbents for Liquid Hazardous Substance Cleanup and Control;							
R.W Melvold et al: Pollution Technolo	R.W Melvold et al: Pollution Technology Review No. 150: Noyes Data Corporation 1988 Slippery when spilt.							
Slippery when spilt.								
Clear area of personnel and move								
 Alert Fire Brigade and tell them Ic 	 Alert Fire Brigade and tell them location and nature of hazard. 							
May be violently or explosively re-	active.							
 Wear breathing apparatus plus pr 	rotective of	gloves.						
 Prevent, by any means available, 	spillage	from entering drai	ns or water cours	es				
No smoking, naked lights or igniti-	 No smoking, naked lights or ignition sources. Increase ventilation. Stop leak if safe to do so. 							
Increase ventilation.								
Stop leak if safe to do so.								
Water spray or fog may be used t	o dispers	e / absorb vapour						
 Absorb or cover spill with sand, ex 	arth, inert	materials or vern	niculite.					
			oors, away from ig	nition sources, until pressure has dissip				
 Undamaged cans should be gath 	 Undamaged cans should be gathered and stowed safely. 							
 Collect residues and seal in label 	 Collect residues and seal in labelled drums for disposal. 							

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

SECTION 7 HANDLING AND STORAGE

Safe handling	 Avoid all personal contact, including inhalation. Wear protective clothing when risk of exposure occurs. Use in a well-ventilated area. Prevent concentration in hollows and sumps. DO NOT enter confined spaces until atmosphere has been checked. Avoid smoking, naked lights or ignition sources. Avoid contact with incompatible materials. When handling, DO NOT eat, drink or smoke. DO NOT incinerate or puncture aerosol cans. DO NOT spray directly on humans, exposed food or food utensils. Avoid physical damage to containers. Always wash hands with soap and water after handling. Work clothes should be laundered separately. Use good occupational work practice. Observe manufacturer's storage and handling recommendations contained within this SDS. Atmosphere should be regularly checked against established exposure standards to ensure safe working conditions are maintained. DO NOT allow clothing wet with material to stay in contact with skin
Other information	 Keep dry to avoid corrosion of cans. Corrosion may result in container perforation and internal pressure may eject contents of can Store in original containers in approved flammable liquid storage area. DO NOT store in pits, depressions, basements or areas where vapours may be trapped. No smoking, naked lights, heat or ignition sources. Keep containers securely sealed. Contents under pressure. Store away from incompatible materials. Store in a cool, dry, well ventilated area. Avoid storage at temperatures higher than 40 deg C. Store in an upright position. Protect containers against physical damage. Check regularly for spills and leaks. Observe manufacturer's storage and handling recommendations contained within this SDS.

Conditions for safe storage, including any incompatibilities

Suitable container	 Aerosol dispenser. Check that containers are clearly labelled.
Storage incompatibility	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	isohexanes	Hexane, other isomers	500 ppm / 1760 mg/m3	3500 mg/m3 / 1000 ppm	Not Available	Not Available
Australia Exposure Standards	LPG (liquefied petroleum gas)	LPG (liquified petroleum gas)	1000 ppm / 1800 mg/m3	Not Available	Not Available	Not Available

EMERGENCY LIMITS

Ingredient	Material name	TEEL-1		TEEL-2	TEEL-3	
LPG (liquefied petroleum gas)	Liquified petroleum gas; (L.P.G.) 65,000 ppm			2.30E+05 ppm	4.00E+05 ppm	
Ingredient	Original IDLH		Revised IDLH			
isohexanes	Not Available		Not Availa	Not Available		
liquid hydrocarbons	Not Available		Not Available			
LPG (liquefied petroleum gas)	2,000 ppm		Not Availa	able		

MATERIAL DATA

Odour Safety Factor(OSF) OSF=0.16 (LPG (liquefied petroleum gas)) None assigned. Refer to individual constituents.

Exposed individuals are NOT reasonably expected to be warned, by smell, that the Exposure Standard is being exceeded.

Odour Safety Factor (OSF) is determined to fall into either Class C, D or E.

The Odour Safety Factor (OSF) is defined as:

OSF= Exposure Standard (TWA) ppm/ Odour Threshold Value (OTV) ppm

Classification into classes follows:

ClassOSF Description

A 550 Over 90% of exposed individuals are aware by smell that the Exposure Standard (TLV-TWA for example) is being reached, even when distracted by working activities B 26-550As "A" for 50-90% of persons being distracted

- C 1-26 As "A" for less than 50% of persons being distracted
- D 0.18-1 10-50% of persons aware of being tested perceive by smell that the Exposure Standard is being reached
- E <0.18 As "D" for less than 10% of persons aware of being tested

NOTE K: The classification as a carcinogen need not apply if it can be shown that the substance contains less than 0.1%w/w 1,3-butadiene (EINECS No 203-450-8). - European Union (EU) List of harmonised classification and labelling hazardous substances, Table 3.1, Annex VI, Regulation (EC) No 1272/2008 (CLP) - up to the latest ATP

Exposure controls

	 Engineering controls are used to remove a hazard or place a barrier between the worker and the hazard. Well-designed engineering control be highly effective in protecting workers and will typically be independent of worker interactions to provide this high level of protection. 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 ventilation that strateg "adds" and "removes" air in the work environment. Ventilation can remove or dilute an air contaminant if designed properly. The design of a ventilation system must match the particular process and chemical or contaminant in use. Employers may need to use multiple types of controls to prevent employee overexposure. General exhaust is adequate under normal conditions. If risk of overexposure exists, wear SAA approved respirator. Correct fit is essential obtain adequate protection. Provide adequate ventilation in warehouse or closed storage areas. Air contaminants generated in the workplace possess varying "escape" velocities which, in turn, determine the "capture velocities" of fresh circulating air required to effectively remove the contaminant. 				
	Type of Contaminant:		Speed:		
Appropriate engineering	aerosols, (released at low velocity into zone of active generation	ration)	0.5-1 m/s		
controls	direct spray, spray painting in shallow booths, gas discharg	e (active generation into zone of rapid air motion)	1-2.5 m/s (200-500 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	r 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 de with the square of distance from the extraction point (in simple cases). Therefore the air speed at the extraction point should be adjust accordingly, after reference to distance from the contaminating source. The air velocity at the extraction fan, for example, should be an 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 multi factors of 10 or more when extraction systems are installed or used.				
Personal protection					
Eye and face protection	No special equipment for minor exposure i.e. when handling small quantities. OTHERWISE: For potentially moderate or heavy exposures: F Safety glasses with side shields. F NOTE: Contact lenses pose a special hazard; soft lenses may absorb irritants and ALL lenses concentrate them.				
Skin protection	See Hand protection below				
Hands/feet protection	 No special equipment needed when handling small quan OTHERWISE: For potentially moderate exposures: Wear general protective gloves, eg. light weight rubber g For potentially heavy exposures: 				

	► Wear chemical protective gloves, eg. PVC. and safety footwear.
Body protection	See Other protection below
Other protection	No special equipment needed when handling small quantities. OTHERWISE: • Overalls. • Skin cleansing cream. • Eyewash unit. • Do not spray on hot surfaces.

Respiratory protection

Type AX-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	Air-line*	AX-2 P2	AX-PAPR-2 P2 ^
up to 20 x ES	-	AX-3 P2	-
20+ x ES	-	Air-line**	-

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

^ - Full-face

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

SECTION 9 PHYSICAL AND CHEMICAL PROPERTIES

Information on basic physical and chemical properties

Appearance	Highly flammable liquid; not miscible with water. Supplied as an aerosol pack. Contents under PRESSURE . Contains highly flammable hydrocarbon propellant.		
Physical state	Liquid	Relative density (Water = 1)	Not Available
Odour	Not Available	Partition coefficient n-octanol / water	Not Available
Odour threshold	Not Available	Auto-ignition temperature (°C)	Not Available
pH (as supplied)	Not Applicable	Decomposition temperature	Not Available
Melting point / freezing point (°C)	Not Available	Viscosity (cSt)	Not Available
Initial boiling point and boiling range (°C)	Not Available	Molecular weight (g/mol)	Not Applicable
Flash point (°C)	<23	Taste	Not Available
Evaporation rate	Not Available	Explosive properties	Not Available
Flammability	HIGHLY FLAMMABLE.	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 (1%)	Not Applicable
Vapour density (Air = 1)	>1	VOC g/L	Not Available

SECTION 10 STABILITY AND REACTIVITY

Reactivity	See section 7
Chemical stability	 Elevated temperatures. Presence of open flame. 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

Inhaled

Information on toxicological effects

Inhalation of aerosols (mists, fumes), generated by the material during the course of normal handling, may be damaging to the health of the individual.

Limited evidence or practical experience suggests that the material may produce irritation of the respiratory system, in a significant number of

	individuals, following inhalation. In contrast to most organs, the lung is able to respond to a chemical insult by first removing or neutralising the irritant and then repairing the damage. The repair process, which initially evolved to protect mammalian lungs from foreign matter and antigens, may however, produce further lung damage resulting in the impairment of gas exchange, the primary function of the lungs. Respiratory tract irritation often results in an inflammatory response involving the recruitment and activation of many cell types, mainly derived from the vascular system. WARNING:Intentional misuse by concentrating/inhaling contents may be lethal . If exposure to highly concentrated solvent atmosphere is prolonged this may lead to narcosis, unconsciousness, even coma and possible death. Hydrocarbons may sensitise the heart to adrenalin and other circulatory catecholamines; as a result cardiac arrhythmias and ventricular fibrillation may occur. Abrupt collapse may produce traumatic injury. Central nervous system (CNS) depression may be evident early. Symptoms of moderate poisoning may include giddiness, headache, dizziness and nausea. Serious poisonings may result in respiratory depression and may be fatal. The paraffin gases C1-4 are practically non-toxic below their lower flammability limits (18000-50000 ppm). Above this level, incidental effects include CNS depression and irritation but these are reversible upon cessation of the exposure. The C3 and iso-C5 hydrocarbons show increasing narcotic properties; branching of the chain also enhances the effect. The C4 hydrocarbons appear to be more highly neurotoxic than the C3 and C5 members. Several fatalities due to voluntary inhalation of butane have been reported, possibly due to central, respiratory and circulatory effects resulting from anaesthesia, laryngeal oedema, chemical pneumonia or the combined effects of cardiac toxicity and increased sympathomimetic effects. Inhalation of petroleum gases may produce narcosis, due in part to olefinic im
Ingestion	Not normally a hazard due to physical form of product. Considered an unlikely route of entry in commercial/industrial environments Accidental ingestion of the material may be damaging to the health of the individual. Central nervous system (CNS) depression may include nonspecific 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. Ingestion of petroleum hydrocarbons may produce irritation of the pharynx, oesophagus, stomach and small intestine with oedema and mucosal ulceration resulting; symptoms include a burning sensation in the mouth and throat. Large amounts may produce narcosis with nausea and vomiting, weakness or dizziness, slow and shallow respiration, swelling of the abdomen, unconsciousness and convulsions. Myocardial injury may produce arrhythmias, ventricular fibrillation and electrocardiographic changes. Central nervous system depression may also occur. Light aromatic hydrocarbons produce a warm, sharp, tingling sensation on contact with taste buds and may anaesthetise the tongue. Aspiration into the lungs may produce coughing, gagging and a chemical pneumonitis with pulmonary oedema and haemorrhage.
Skin Contact	Limited evidence exists, or practical experience predicts, that the material either produces inflammation of the skin in a substantial number of individuals following direct contact, and/or produces significant inflammation when applied to the healthy intact skin of animals, for up to four hours, such inflammation being present twenty-four hours or more after the end of the exposure period. Skin irritation may also be present after prolonged or repeated exposure; this may result in a form of contact dermatitis (nonallergic). The dermatitis is often characterised by skin redness (erythema) and swelling (oedema) which may progress to blistering (vesiculation), scaling and thickening of the epidermis. At the microscopic level there may be intercellular oedema of the spongy layer of the skin (spongiosis) and intracellular oedema of the epidermis. Spray mist may produce discomfort Entry into the blood-stream through, for example, cuts, abrasions, puncture wounds 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 skin irritation, vasodilation with erythema and changes in endothelial cell permeability. Systemic intorease percutaneous absorption. In acute skin irritation tests, C12-C18 alpha and internal alkenes (but not C20-24 branched and linear alkenes) caused prolonged slight or slight to moderate skin irritation associated with desquamation. Repeat dermal applications of C12-C16, C16-C18 or C16 alone caused severe skin irritation (Chevron Chemical)
Eye	Limited evidence exists, or practical experience suggests, that the material may cause eye irritation in a substantial number of individuals and/or is expected to produce significant ocular lesions which are present twenty-four hours or more after instillation into the eye(s) of experimental animals. Repeated or prolonged eye contact may cause inflammation characterised by temporary redness (similar to windburn) of the conjunctiva (conjunctivitis); temporary impairment of vision and/or other transient eye damage/ulceration may occur. Petroleum hydrocarbons may produce pain after direct contact with the eyes. Slight, but transient disturbances of the corneal epithelium may also
	result. The aromatic fraction may produce irritation and lachrymation. Principal route of occupational exposure to the gas is by inhalation.
Chronic	Limited evidence suggests that repeated or long-term occupational exposure may produce cumulative health effects involving organs or biochemical systems. Repeated or prolonged exposure to mixed hydrocarbons may produce narcosis with dizziness, weakness, irritability, concentration and/or memory loss, tremor in the fingers and tongue, vertigo, olfactory disorders, constriction of visual field, paraesthesias of the extremities, weight loss and anaemia and degenerative changes in the liver and kidney. Chronic exposure by petroleum workers, to the lighter hydrocarbons, has been associated with visual disturbances, damage to the central nervous system, peripheral neuropathies (including numbness and paraesthesias), psychological and neurophysiological deficits, bone marrow toxicities (including hypoplasia possibly due to benzene) and hepatic and renal involvement. Chronic dermal exposure to petroleum hydrocarbons may result in defatting which produces localised dermatoses. Surface cracking and erosion may also increase susceptibility to infection by microorganisms. One epidemiological study of petroleum refinery workers has reported elevations in standard mortality ratios for skin cancer along with a dose-response relationship indicating an association between routine workplace exposure to petroleum or one of its constituents and skin cancer, particularly melanoma. Other studies have been unable to confirm this finding. Hydrocarbon solvents are liquid hydrocarbon fractions derived from petroleum processing streams, containing only carbon and hydrogen atoms, with carbon numbers ranging from approximately C5-C20 and boiling between approximately 35-370 deg C. Many of the hydrocarbon solvents have complex and variable compositions with constituents of 4 types, alkanes (normal paraffins, isoparaffins, and cycloparaffins) and aromatics (primarily alkylated one- and two-ring species). Despite the compositional complexity, most hydrocarbon solvents cause chemical pneumonitis if aspirated into the lung, and those that are

response is likely related to chronic irritation and not to dose. The mutagenic potential of naphthas has been reported to be largely negative in a variety of mutagenicity tests. The exact relationship between these results and human health is not known. Some components of this product

have been shown to produce a species specific, sex hormonal dependent kidney lesion in male rats from repeated oral or inhalation exposure. Subsequent research has shown that the kidney damage develops via the formation of a alpha-2u-globulin, a mechanism unique to the male rat. Humans do not form alpha-2u-globulin, therefore, the kidney effects resulting from this mechanism are not relevant in human.

Principal route of exposure is by skin contact; lesser exposures include inhalation of fumes from hot oils, oil mists or droplets. Prolonged contact with mineral oils carries with it the risk of skin conditions such as oil folliculitis, eczematous dermatitis, pigmentation of the face (melanosis) and warts on the sole of the foot (plantar warts). With highly refined mineral oils no appreciable systemic effects appear to result through skin absorption.

Exposure to oil mists frequently elicits respiratory conditions, such as asthma; the provoking agent is probably an additive. High oil mist concentrations may produce lipoid pneumonia although clinical evidence is equivocal. In animals exposed to concentrations of 100 mg/m3 oil mist, for periods of 12 to 26 months, the activity of lung and serum alkaline phosphatase enzyme was raised; 5 mg/m3 oil mist did not produce this response. These enzyme changes are sensitive early indicators of lung damage. Workers exposed to vapours of mineral oil and kerosene for 5 to 35 years showed an increased prevalence of slight basal lung fibrosis.

Many studies have linked cancers of the skin and scrotum with mineral oil exposure. Contaminants in the form of additives and the polycyclic aromatic hydrocarbons (PAHs - as in the crude base stock) are probably responsible. PAH levels are higher in aromatic process oils/used /reclaimed motor oils. Subchronic 90-day feeding studies conducted on male and female rats on highly refined white mineral oils and waxes found that higher molecular-weight hydrocarbons (microcrystalline waxes and the higher viscosity oils) were without biological effects. Paraffin waxes and low- to mid viscosity oils produced biological effects that were inversely proportional to molecular weight, viscosity and melting point: oil-type and processing did not appear to be determinants. Biological effects were more pronounced in females than in males. Effects occurred mainly in the liver and mesenteric lymph nodes and included increased organ weights, microscopic inflammatory changes, and evidence for the presence of saturated mineral hydrocarbons in affected tissues. Inflammation of the cardiac mitral valve was also observed at high doses in rats treated with paraffin waxes.

Smith J.H., et al: Toxicologic Pathology: 24, 2, 214-230, 1996

Chronic solvent inhalation exposures may result in nervous system impairment and liver and blood changes. [PATTYS]

CRC 3013 Soft Seal / Panel	ΤΟΧΙΟΙΤΥ	IRRITATION	
Seal (Aerosol) (NZ)	Not Available	Not Available	
isohexanes	ΤΟΧΙΟΙΤΥ	IRRITATION	
	Not Available	Not Available	
liquid hydrocarbons	ΤΟΧΙΟΙΤΥ	IRRITATION	
	Not Available	Not Available	
LPG (liquefied petroleum gas)	ΤΟΧΙCΙΤΥ	IRRITATION	
	Not Available	Not Available	
Legend:	1. Value obtained from Europe ECHA Registered S specified data extracted from RTECS - Register of	Substances - Acute toxicity 2.* Value obtained from manufacturer's SE Toxic Effect of chemical Substances	S. Unless otherwise

LIQUID HYDROCARBONS	For olefins: Studies have shown that normal alpha olefins have little or no toxic effect on animals except in very severe inhalation conditions and that they may produce minimal skin and eye irritation, but are not skin sensitisers. Laboratory exposures to very high airborne concentrations of C6-C16 normal alpha olefin vapors or mists produced central nervous system effects including anesthesia. If C20+ products are heated, fumes may produce nausea and irritation of the upper respiratory tract. Although not all products have been tested in genetic toxicity assays, the available data indicate normal alpha olefins are not mutagenic. Acute toxicity : The weight of evidence indicates alpha and internal olefins with carbon numbers between C6 and C54 have a similar and low level of mammalian toxicity, and the toxicity profile is not affected by changes in the location of the double bond or the addition of branching to the structure. These materials are not eye irritants or skin sensitisers. Prolonged exposure of the skin for many hours may cause skin irritation. Olefins (alkenes) ranging in carbon number from C6 to C42 alpha (linear) and internal (linear and branched), and C24-54 alpha (linear and branched) demonstrate low acute toxicity by the oral, inhalation and dermal routes of exposure: Rat oral LD50 > 5 g/kg; rat 4-hr inhalation LC50 range = 110 mg/L (32,000 ppm) to 6.4 mg/L (683 ppm) for C6 to C16; and ratrabibit dermal LD50 > highest doses tested (1.43-10 g/kg). Repeated dose toxicity : Studies, using the inhalation (C6 alpha), dermal (C12-16 alpha), or oral (C6 alpha and internal linear/branched; c8 and C14 alpha; and C16/18, C18 and C20-24 internal linear/branched) routes of exposure, have shown comparable levels of low toxicity in rats. In females, alterations in body and organ weights, changes in certain clinical chemistry/heematology values, and liver effects were seen in oral studies with C14 alpha olefins (minimal-to-mild hepatocyte cytoplasmic vacuolation with increased liver weight
LPG (LIQUEFIED PETROLEUM GAS)	for Petroleum Hydrocarbon Gases: In many cases, there is more than one potentially toxic constituent in a refinery gas. In those cases, the constituent that is most toxic for a particular endpoint in an individual refinery stream is used to characterize the endpoint hazard for that stream. The hazard potential for each mammalian endpoint for each of the petroleum hydrocarbon gases is dependent upon each petroleum hydrocarbon gas constituent endpoint toxicity values (LC50, LOAEL, etc.) and the relative concentration of the constituent present in that gas. It should also be noted that for an individual petroleum hydrocarbon gas, the constituent characterizing toxicity may be different for different mammalian endpoints, again, being dependent upon the concentration of the different constituents in each, distinct petroleum hydrocarbon gas. All Hydrocarbon Gases Category members contain primarily hydrocarbons (i.e., alkanes and alkenes) and occasionally asphyxiant gases like hydrogen. The inorganic components of the petroleum hydrocarbon gases are less toxic than the C1 - C4 and C5 - C6 hydrocarbon components to both mammalian and aquatic organisms. Unlike other petroleum product categories (<i>e.g.</i> gasoline, diesel fuel, lubricating oils, etc.), the inorganic and hydrocarbon constituents of hydrocarbon gases can be evaluated for hazard individually to then predict the screening level hazard of the Category members
	Destinued

Continued...

CRC 3013 Soft Seal / Panel Seal (Aerosol) (NZ)

	petroleum hydrocarbon gas constituents from most to C5-C6 HCs (LC50 > 1063 ppm) > C1-C4 HCs (LC50 asphyxiant gases (hydrogen, carbon dioxide, nitroger Repeat dose toxicity: With the exception of the asp hydrocarbon gas constituents. Based upon LOAEL v the least toxic is: Benzene (LOAEL .>=10 ppm) >C1-C4 HCs (LOAEL butadiene (LOAEL = 8,000 ppm) > asphyxiant gases Genotoxicity: <i>In vitro</i> : The majority of the Petroleum Hydrocarbon benzene and 1,3-butadiene, which are genotoxic in to <i>In vivo</i> : The majority of the Petroleum Hydrocarbon exceptions are benzene and 1,3-butadiene, which are Developmental toxicity : Bvelopmental effects wer hydrocarbon fraction. No developmental toxicity was	 (- 5 mg/l) for these petroleum hydroc to least toxic is:) > 10,000 ppm) > benzene (LC50 = an). ohyxiant gases, repeated dose toxicity values, the order of order of repeated = 5,000 ppm; assumed to be 100% 2 is (hydrogen, carbon dioxide, nitrogen carbon di dioxide, nitrogen carbon dio	y has been observed in individual selected petroleum dose toxicity of these constituents from most toxic to 2-butene) > C5-C6 HCs (LOAEL = 6,625 ppm) >). gative for <i>in vitro</i> genotoxicity. The exceptions are: systems. gative for <i>in vivo</i> genotoxicity. The ydrocarbon gas constituents, benzene and the C5 -C6 vels tested for the other petroleum hydrocarbon gas omental toxicity. Based on LOAEL and NOAEL values = 3,463 ppm) > C1-C4 HCs (NOAEL >=5,000 ppm;
	Reproductive toxicity: Reproductive effects were in constituent of the the C1-C4 hydrocarbon fraction). N petroleum hydrocarbon gas constituents tested for th LOAEL and NOAEL values, the order of reproductive	nduced by only two petroleum hydroc No reproductive toxicity was observed his effect. The asphyxiant gases have e toxicity of these constituents from n	arbon gas constituents, benzene and isobutane (a d at the highest exposure levels tested for the other e not been tested for reproductive toxicity. Based on nost to least toxic is:
	Reproductive toxicity: Reproductive effects were in constituent of the the C1-C4 hydrocarbon fraction). N petroleum hydrocarbon gas constituents tested for th LOAEL and NOAEL values, the order of reproductive	duced by only two petroleum hydroc No reproductive toxicity was observed his effect. The asphyxiant gases have e toxicity of these constituents from n .>=6,000 ppm) > C5-C6 HCs (NOAE	arbon gas constituents, benzene and isobutane (a d at the highest exposure levels tested for the other e not been tested for reproductive toxicity. Based on nost to least toxic is: L .>=6,521 ppm) > C1-C4 HCs (LOAEL = 9,000 ppm;
ISOHEXANES & LIQUID HYDROCARBONS & LPG (LIQUEFIED PETROLEUM GAS)	Reproductive toxicity: Reproductive effects were in constituent of the the C1-C4 hydrocarbon fraction). N petroleum hydrocarbon gas constituents tested for the LOAEL and NOAEL values, the order of reproductive Benzene (LOAEL = 300 ppm) > butadiene (NOAEL .	duced by only two petroleum hydroc No reproductive toxicity was observed his effect. The asphyxiant gases have e toxicity of these constituents from n .>=6,000 ppm) > C5-C6 HCs (NOAEI s (hydrogen, carbon dioxide, nitrogen	arbon gas constituents, benzene and isobutane (a d at the highest exposure levels tested for the other e not been tested for reproductive toxicity. Based on nost to least toxic is: L .>=6,521 ppm) > C1-C4 HCs (LOAEL = 9,000 ppm;
HYDROCARBONS & LPG (LIQUEFIED PETROLEUM	Reproductive toxicity: Reproductive effects were in constituent of the the C1-C4 hydrocarbon fraction). N petroleum hydrocarbon gas constituents tested for th LOAEL and NOAEL values, the order of reproductive Benzene (LOAEL = 300 ppm) > butadiene (NOAEL . assumed to be 100% isobutane) > asphyxiant gases	nduced by only two petroleum hydroc No reproductive toxicity was observed nis effect. The asphyxiant gases have e toxicity of these constituents from n .>=6,000 ppm) > C5-C6 HCs (NOAEI s (hydrogen, carbon dioxide, nitrogen erature search.	arbon gas constituents, benzene and isobutane (a d at the highest exposure levels tested for the other e not been tested for reproductive toxicity. Based on nost to least toxic is: L .>=6,521 ppm) > C1-C4 HCs (LOAEL = 9,000 ppm;
HYDROCARBONS & LPG (LIQUEFIED PETROLEUM GAS)	Reproductive toxicity: Reproductive effects were in constituent of the the C1-C4 hydrocarbon fraction). N petroleum hydrocarbon gas constituents tested for th LOAEL and NOAEL values, the order of reproductive Benzene (LOAEL = 300 ppm) > butadiene (NOAEL . assumed to be 100% isobutane) > asphyxiant gases No significant acute toxicological data identified in lite	duced by only two petroleum hydroc No reproductive toxicity was observed his effect. The asphyxiant gases have e toxicity of these constituents from n .>=6,000 ppm) > C5-C6 HCs (NOAEI s (hydrogen, carbon dioxide, nitrogen	arbon gas constituents, benzene and isobutane (a d at the highest exposure levels tested for the other e not been tested for reproductive toxicity. Based on nost to least toxic is: L.>=6,521 ppm) > C1-C4 HCs (LOAEL = 9,000 ppm;)
HYDROCARBONS & LPG (LIQUEFIED PETROLEUM GAS) Acute Toxicity Skin Irritation/Corrosion	Reproductive toxicity: Reproductive effects were in constituent of the the C1-C4 hydrocarbon fraction). N petroleum hydrocarbon gas constituents tested for th LOAEL and NOAEL values, the order of reproductive Benzene (LOAEL = 300 ppm) > butadiene (NOAEL . assumed to be 100% isobutane) > asphyxiant gases No significant acute toxicological data identified in lite	hduced by only two petroleum hydroc No reproductive toxicity was observed is effect. The asphyxiant gases have e toxicity of these constituents from m >=6,000 ppm) > C5-C6 HCs (NOAE s (hydrogen, carbon dioxide, nitrogen erature search.	arbon gas constituents, benzene and isobutane (a d at the highest exposure levels tested for the other e not been tested for reproductive toxicity. Based on nost to least toxic is: L.>=6,521 ppm) > C1-C4 HCs (LOAEL = 9,000 ppm;)
HYDROCARBONS & LPG (LIQUEFIED PETROLEUM GAS) Acute Toxicity	Reproductive toxicity: Reproductive effects were in constituent of the the C1-C4 hydrocarbon fraction). N petroleum hydrocarbon gas constituents tested for th LOAEL and NOAEL values, the order of reproductive Benzene (LOAEL = 300 ppm) > butadiene (NOAEL . assumed to be 100% isobutane) > asphyxiant gases No significant acute toxicological data identified in lite	hduced by only two petroleum hydrod No reproductive toxicity was observed is effect. The asphyxiant gases have e toxicity of these constituents from n >=6,000 ppm) > C5-C6 HCs (NOAE s (hydrogen, carbon dioxide, nitrogen erature search.	arbon gas constituents, benzene and isobutane (a d at the highest exposure levels tested for the other e not been tested for reproductive toxicity. Based on nost to least toxic is: L.>=6,521 ppm) > C1-C4 HCs (LOAEL = 9,000 ppm;)

SECTION 12 ECOLOGICAL INFORMATION

Toxicity

CRC 3013 Soft Seal / Panel	ENDPOINT	TEST DURATION (HR) SPECIES		VALUE	SOURCE
Seal (Aerosol) (NZ)	Not Available	Not Available	Not Available	Not Available	Not Available
	ENDPOINT	TEST DURATION (HR)	SPECIES	VALUE	SOURCE
isohexanes	LC50	96	Fish	0.961mg/L	3
	EC50	96	Algae or other aquatic plants	1.526mg/L	3
liquid hydrocarbons	ENDPOINT	TEST DURATION (HR)	SPECIES	VALUE	SOURCE
	Not Available	Not Available	Not Available	Not Available	Not Available
	ENDPOINT	TEST DURATION (HR)	SPECIES	VALUE	SOURCI
LPG (liquefied petroleum gas)	LC50	96	Fish	24.11mg/L	2
	EC50	96	Algae or other aquatic plants	7.71mg/L	2
Legend:	V3.12 (QSAR) -	Aquatic Toxicity Data (Estimated) 4. US	A Registered Substances - Ecotoxicological Informat S EPA, Ecotox database - Aquatic Toxicity Data 5. Ec Japan) - Bioconcentration Data 8. Vendor Data	, ,	

Drinking Water Standards: hydrocarbon total: 10 ug/l (UK max.).

DO NOT discharge into sewer or waterways.

Persistence and degradability

Ingredient	Persistence: Water/Soil	Persistence: Air
isohexanes	LOW	LOW

Bioaccumulative potential

Ingredient	Bioaccumulation
isohexanes	LOW (LogKOW = 3.7056)

Mobility in soil

Ingredient	Mobility
isohexanes	LOW (KOC = 230.3)

SECTION 13 DISPOSAL CONSIDERATIONS

Waste treatment methods Product / Packaging disposal Discharge contents of damaged aerosol cans at an approved site. Allow small quantities to evaporate. DO NOT incinerate or puncture aerosol cans. Bury residues and emptied aerosol cans at an approved site.

SECTION 14 TRANSPORT INFORMATION

Labels Required

Marine Pollutant	NO
HAZCHEM	Not Applicable

Land transport (ADG)

UN number	1950	
UN proper shipping name	AEROSOLS	
Transport hazard class(es)	Class 2.1 Subrisk Not Applicable	
Packing group	Not Applicable	
Environmental hazard	Not Applicable	
Special precautions for user	Special provisions 63 190 277 327 344 381 Limited quantity 1000ml	

Air transport (ICAO-IATA / DGR)

UN number	1950			
UN proper shipping name	Aerosols, flammable			
Transport hazard class(es)	ICAO/IATA Class ICAO / IATA Subrisk ERG Code	2.1 Not Applicable 10L		
Packing group	Not Applicable			
Environmental hazard	Not Applicable			
	Special provisions Cargo Only Packing Ir Cargo Only Maximum		A145 A167 A802 203 150 kg	
Special precautions for user	Passenger and Cargo Packing Instructions		203	
	Passenger and Cargo Maximum Qty / Pack		75 kg	
	Passenger and Cargo	Limited Quantity Packing Instructions	Y203	
	Passenger and Cargo Limited Maximum Qty / Pack		30 kg G	

Sea transport (IMDG-Code / GGVSee)

UN number	1950
UN proper shipping name	AEROSOLS
Transport hazard class(es)	IMDG Class 2.1 IMDG Subrisk Not Applicable
Packing group	Not Applicable
Environmental hazard	Not Applicable

Issue Date: 01/11/2019 Print Date: 03/04/2020

CRC 3013 Soft Seal / Panel Seal (Aerosol) (NZ)

	EMS Number	F-D , S-U
Special precautions for user	Special provisions	63 190 277 327 344 381 959
	Limited Quantities	1000 ml

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

ISOHEXANES IS FOUND ON THE FOLLOWING REGULATORY LISTS

Not Applicable

LIQUID HYDROCARBONS IS FOUND ON THE FOLLOWING REGULATORY LISTS

Not Applicable

LPG (LIQUEFIED PETROLEUM GAS) IS FOUND ON THE FOLLOWING REGULATORY LISTS

Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals Australia Inventory of Chemical Substances (AICS) Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) -Schedule 5 Chemical Footprint Project - Chemicals of High Concern List

National Inventory Status

National Inventory	Status
Australia - AICS	No (isohexanes)
Canada - DSL	No (isohexanes)
Canada - NDSL	No (isohexanes; LPG (liquefied petroleum gas))
China - IECSC	Yes
Europe - EINEC / ELINCS / NLP	Yes
Japan - ENCS	Yes
Korea - KECI	No (isohexanes)
New Zealand - NZIoC	Yes
Philippines - PICCS	Yes
USA - TSCA	No (isohexanes)
Taiwan - TCSI	Yes
Mexico - INSQ	No (isohexanes)
Vietnam - NCI	Yes
Russia - ARIPS	No (isohexanes)
Legend:	Yes = All CAS declared ingredients are on the inventory No = One or more of the CAS listed ingredients are not on the inventory and are not exempt from listing(see specific ingredients in brackets)

SECTION 16 OTHER INFORMATION

Revision Date	01/11/2019
Initial Date	01/11/2009

SDS Version Summary

Version	Issue Date	Sections Updated
7.1.1.1	21/06/2018	Physical Properties
8.1.1.1	01/11/2019	One-off system update. NOTE: This may or may not change the GHS classification

Other information

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

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

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