

Trouble Shooter Acr.-Neutral Base 45195

ICP Construction

Version No: **1.3**Safety Data Sheet according to OSHA HazCom Standard (2012) requirements

Issue Date: **02/07/2017** Print Date: **02/07/2017** S.GHS.USA.EN

SECTION 1 IDENTIFICATION

Product Identifier

Product name	Trouble Shooter AcrNeutral Base 45195	
Synonyms	Not Available	
Proper shipping name	Environmentally hazardous substance, liquid, n.o.s.	
Other means of identification	Not Available	

Recommended use of the chemical and restrictions on use

Relevant identified uses	Primer for Exterior wood clapboards, siding, shingles, trim, shakes and hardboard
--------------------------	---

Name, address, and telephone number of the chemical manufacturer, importer, or other responsible party

Registered company name	ICP Construction	
Address	Address 150 Dascomb Road Massachusetts Andover United States	
Telephone	978-623-9980	
Fax	Not Available	
Website	Not Available	
Email	Not Available	

Emergency phone number

Association / Organisation	Chemtel
Emergency telephone numbers	1-800-255-3924
Other emergency telephone numbers	1-813-248-0585

SECTION 2 HAZARD(S) IDENTIFICATION

Classification of the substance or mixture

Classification

Eye Irritation Category 2A, Skin Sensitizer Category 1, Carcinogenicity Category 2, Acute Aquatic Hazard Category 2, Chronic Aquatic Hazard Category 2, Specific target organ toxicity - repeated exposure Category 1

Label elements

GHS label elements







SIGNAL WORD

DANGER

Hazard statement(s)

H319	Causes serious eye irritation.	
H317 May cause an allergic skin reaction.		
H351	Suspected of causing cancer.	
H411	H411 Toxic to aquatic life with long lasting effects.	
H372 Causes damage to organs through prolonged or repeated exposure.		

Hazard(s) not otherwise specified

Chemwatch: 9-339815 Page 2 of 12 Issue Date: 02/07/2017 Version No: 1.3 Print Date: 02/07/2017

Trouble Shooter Acr.-Neutral Base 45195

Precautionary statement(s) Prevention

P201 Obtain special instructions before use.	
P260 Do not breathe dust/fume/gas/mist/vapours/spray.	
P280 Wear protective gloves/protective clothing/eye protection/face protection.	

Precautionary statement(s) Response

P308+P313	IF exposed or concerned: Get medical advice/attention.	
P363	P363 Wash contaminated clothing before reuse.	
P302+P352 IF ON SKIN: Wash with plenty of soap and water.		

Precautionary statement(s) Storage

P405 Store locked up.

Precautionary statement(s) Disposal

P501 Dispose of contents/container in accordance with local regulations.

SECTION 3 COMPOSITION / INFORMATION ON INGREDIENTS

Substances

See section below for composition of Mixtures

Mixtures

CAS No	%[weight]	Name
57-55-6	3-7	propylene glycol
1897-45-6	0.1-1	<u>chlorothalonil</u>
26172-55-4	0.1-1	5-chloro-2-methyl-4-isothiazolin-3-one
1314-13-2	3-10	zinc oxide

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. 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, without delay.
Ingestion	 Immediately give a glass of water. First aid is not generally required. If in doubt, contact a Poisons Information Centre or a doctor.

Most important symptoms and effects, both acute and delayed

See Section 11

Indication of any immediate medical attention and special treatment needed

Treat symptomatically.

As in all cases of suspected poisoning, follow the ABCDEs of emergency medicine (airway, breathing, circulation, disability, exposure), then the ABCDEs of toxicology (antidotes, basics, change absorption, change distribution, change elimination).

For poisons (where specific treatment regime is absent):

BASIC TREATMENT

- Establish a patent airway with suction where necessary.
- Watch for signs of respiratory insufficiency and assist ventilation as necessary.
- Administer oxygen by non-rebreather mask at 10 to 15 L/min.
- Monitor and treat, where necessary, for pulmonary oedema.
- Monitor and treat, where necessary, for shock.
- ▶ Anticipate seizures.
- DO NOT use emetics. Where ingestion is suspected rinse mouth and give up to 200 ml water (5 ml/kg recommended) for dilution where patient is able to swallow, has a strong gag reflex and does not drool.

Chemwatch: 9-339815 Page 3 of 12 Issue Date: 02/07/2017

Version No: 1.3 Trouble Shooter Acr.-Neutral Base 45195 Print Date: 02/07/2017

ADVANCED TREATMENT

▶ Consider orotracheal or nasotracheal intubation for airway control in unconscious patient or where respiratory arrest has occurred.

- ▶ Positive-pressure ventilation using a bag-valve mask might be of use.
- ▶ Monitor and treat, where necessary, for arrhythmias.
- ▶ Start an IV D5W TKO. If signs of hypovolaemia are present use lactated Ringers solution. Fluid overload might create complications.
- ▶ Drug therapy should be considered for pulmonary oedema.
- ▶ Hypotension with signs of hypovolaemia requires the cautious administration of fluids. Fluid overload might create complications.
- Treat seizures with diazepam.
- ▶ Proparacaine hydrochloride should be used to assist eye irrigation.

BRONSTEIN, A.C. and CURRANCE, P.L.

EMERGENCY CARE FOR HAZARDOUS MATERIALS EXPOSURE: 2nd Ed. 1994

SECTION 5 FIRE-FIGHTING MEASURES

Extinguishing media

- ▶ There is no restriction on the type of extinguisher which may be used.
- Use extinguishing media suitable for surrounding area.

Special hazards arising from the substrate or mixture

Fire Incompatibility	ty None known.	
Special protective equipm	ent and precautions for fire-fighters	
Fire Fighting	 Alert Fire Brigade and tell them location and nature of hazard. Wear breathing apparatus plus protective gloves in the event of a fire. Prevent, by any means available, spillage from entering drains or water courses. Use fire fighting procedures suitable for surrounding 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. 	
Fire/Explosion Hazard	 Non combustible. Not considered a significant fire risk, however containers may burn. May emit poisonous fumes. 	

SECTION 6 ACCIDENTAL RELEASE MEASURES

Personal precautions, protective equipment and emergency procedures

See section 8

Environmental precautions

See section 12

Methods and material for containment and cleaning up

Minor Spills	Environmental hazard - contain spillage. 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.
Major Spills	 Environmental hazard - contain spillage. Absorb or contain isothiazolinone liquid spills with sand, earth, inert material or vermiculite. The absorbent (and surface soil to a depth sufficient to remove all of the biocide) should be shovelled into a drum and treated with an 11% solution of sodium metabisulfite (Na2S2O5) or sodium bisulfite (NaHSO3), or 12% sodium sulfite (Na2SO3) and 8% hydrochloric acid (HCl). Glutathione has also been used to inactivate the isothiazolinones. Use 20 volumes of decontaminating solution for each volume of biocide, and let containers stand for at least 30 minutes to deactivate microbicide before disposal. If contamination of drains or waterways occurs, advise emergency services. After clean up operations, decontaminate and launder all protective clothing and equipment before storing and re-using.

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

SECTION 7 HANDLING AND STORAGE

Precautions for 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.
Safe handling	▶ DO NOT allow material to contact humans, exposed food or food utensils.
	Avoid contact with incompatible materials.
	When handling, DO NOT eat, drink or smoke.
	 Keep containers securely sealed when not in use.

Avoid physical damage to containers.

Chemwatch: 9-339815 Page 4 of 12 Issue Date: 02/07/2017
Version No: 1.3 Print Date: 02/07/2017

Trouble Shooter Acr.-Neutral Base 45195

Always wash hands with soap and water after handling.

Work clothes should be laundered separately. Launder contaminated clothing before re-use.

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

Store in original containers.

Keep containers securely sealed.

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

- ▶ Lined metal can, lined metal pail/ can.
- ▶ Plastic pail.
- Polyliner drum.
- ▶ Packing as recommended by manufacturer.
- Check all containers are clearly labelled and free from leaks.

For low viscosity materials

- ▶ Drums and jerricans must be of the non-removable head type.
- ▶ Where a can is to be used as an inner package, the can must have a screwed enclosure.

For materials with a viscosity of at least 2680 cSt. (23 deg. C) and solids (between 15 C deg. and 40 deg C.):

- Removable head packaging:
- Cans with friction closures and
- ► low pressure tubes and cartridges

may be used.

Where combination packages are used, and the inner packages are of glass, there must be sufficient inert cushioning material in contact with inner and outer packages *.

In addition, where inner packagings are glass and contain liquids of packing group I and II there must be sufficient inert absorbent to absorb any spillage *.

* unless the outer packaging is a close fitting moulded plastic box and the substances are not incompatible with the plastic.

All inner and sole packagings for substances that have been assigned to Packaging Groups I or II on the basis of inhalation toxicity criteria, must be hermetically sealed.

Storage incompatibility

Suitable container

- Glycols and their ethers undergo violent decomposition in contact with 70% perchloric acid. This seems likely to involve formation of the glycol perchlorate esters (after scission of ethers) which are explosive, those of ethylene glycol and 3-chloro-1,2-propanediol being more powerful than glyceryl nitrate, and the former so sensitive that it explodes on addition of water.
- Zinc oxide:
- slowly absorbs carbon dioxide from the air.
- ▶ may react, explosively with magnesium and chlorinated rubber when heated
- ► is incompatible with linseed oil (may cause ignition)

None known

SECTION 8 EXPOSURE CONTROLS / PERSONAL PROTECTION

Control parameters

OCCUPATIONAL EXPOSURE LIMITS (OEL)

INGREDIENT DATA

Source	Ingredient	Material name	TWA	STEL	Peak	Notes
US OSHA Permissible Exposure Levels (PELs) - Table Z1	zinc oxide	Zinc oxide fume / Zinc oxide / Zinc oxide - Respirable fraction	5 mg/m3 / 15 mg/m3	Not Available	Not Available	Total dust
US OSHA Permissible Exposure Levels (PELs) - Table Z3	zinc oxide	Inert or Nuisance Dust	5 mg/m3 / 15 mg/m3 / 15 mppcf / 50 mppcf	Not Available	Not Available	Respirable fraction;All inert or nuisance dusts, whether mineral, inorganic, or organic, not listed specifically by substance name are covered by this limit, which is the same as the Particulates Not Otherwise Regulated (PNOR) limit in Table Z-1. / Total dust;All inert or nuisance dusts, whether mineral, inorganic, or organic, not listed specifically by substance name are covered by this limit, which is the same as the Particulates Not Otherwise Regulated (PNOR) limit in Table Z-1.
US ACGIH Threshold Limit Values (TLV)	zinc oxide	Zinc oxide	2 mg/m3	10 mg/m3	Not Available	TLV® Basis: Metal fume fever
US NIOSH Recommended Exposure Limits (RELs)	zinc oxide	Zinc peroxide	Dust: 5 ,Fume: 5 mg/m3	Fume: 10 mg/m3	Dust: 15 mg/m3	Not Available

EMERGENCY LIMITS

Ingredient	Material name	TEEL-1	TEEL-2	TEEL-3
propylene glycol	Propylene glycol; (1,2-Propanediol)	30 mg/m3	1,300 mg/m3	7,900 mg/m3
chlorothalonil	Chlorothalonil; (Tetrachloroisophthalonitrile)	0.13 mg/m3	1.4 mg/m3	8.6 mg/m3
5-chloro-2-methyl- 4-isothiazolin-3-one	Chloro-2-methyl-4-isothiazolin-3-one, 5-	0.6 mg/m3	6.6 mg/m3	40 mg/m3
zinc oxide	Zinc oxide	10 mg/m3	15 mg/m3	2,500 mg/m3

Chemwatch: 9-339815 Page 5 of 12 Issue Date: 02/07/2017
Version No: 1.3 Print Date: 02/07/2017

Trouble Shooter Acr.-Neutral Base 45195

Ingredient	Original IDLH	Revised IDLH
propylene glycol	Not Available	Not Available
chlorothalonil	Not Available	Not Available
5-chloro-2-methyl- 4-isothiazolin-3-one	Not Available	Not Available
zinc oxide	2,500 mg/m3	500 mg/m3

Exposure controls

Engineering controls are used to remove a hazard or place a barrier between the worker and the hazard. Well-designed engineering controls can be highly effective in protecting workers and will typically be independent of worker interactions to provide this high level of protection.

The basic 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 strategically "adds" and "removes" air in the work environment. Ventilation can remove or dilute an air contaminant if designed properly. The design of a ventilation system must match the particular process and chemical or contaminant in use.

Employers may need to use multiple types of controls to prevent employee overexposure.

Local exhaust ventilation usually required. If risk of overexposure exists, wear approved respirator. Correct fit is essential to obtain adequate protection. Supplied-air type respirator may be required in special circumstances. Correct fit is essential to ensure adequate protection.

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 varying "escape" velocities which, in turn, determine the "capture velocities" of fresh circulating air required to effectively remove the contaminant.

Appropriate engineering controls

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 generation into zone of rapid air motion)	1-2.5 m/s (200-500 f/min.)
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

- ► Wear chemical protective gloves, e.g. PVC.
- ▶ Wear safety 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, belts 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.

Hands/feet protection

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 moisturizer is recommended.

Suitability and durability of glove type is dependent on usage. Important factors in the selection of gloves include:

- illty and durability of glove type is dependent on usage. Important
 frequency and duration of contact,
 - chemical resistance of glove material,
 - glove thickness and
 - dexterity

Chemwatch: 9-339815 Page 6 of 12 Issue Date: 02/07/2017 Version No: 1.3

Trouble Shooter Acr.-Neutral Base 45195

Print Date: 02/07/2017

Select gloves tested to a relevant standard (e.g. Europe EN 374, US F739, AS/NZS 2161.1 or national equivalent).

- When prolonged or frequently repeated contact may occur, a glove with a protection class of 5 or higher (breakthrough time greater than 240 minutes according to EN 374, AS/NZS 2161.10.1 or national equivalent) is recommended.
- When only brief contact is expected, a glove with a protection class of 3 or higher (breakthrough time greater than 60 minutes according to EN 374. AS/NZS 2161.10.1 or national equivalent) is recommended.
- 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.

For general applications, gloves with a thickness typically greater than 0.35 mm, are recommended.

It should be emphasised that glove thickness is not necessarily a good predictor of glove resistance to a specific chemical, as the permeation efficiency of the glove will be dependent on the exact composition of the glove material. Therefore, glove selection should also be based on consideration of the task requirements and knowledge of breakthrough times.

Glove thickness may also vary depending on the glove manufacturer, the glove type and the glove model. Therefore, the manufacturers' technical data should always be taken into account to ensure selection of the most appropriate glove for the task.

Note: Depending on the activity being conducted, gloves of varying thickness may be required for specific tasks. For example:

- Thinner gloves (down to 0.1 mm or less) may be required where a high degree of manual dexterity is needed. However, these gloves are only likely to give short duration protection and would normally be just for single use applications, then disposed of.
- Thicker gloves (up to 3 mm or more) may be required where there is a mechanical (as well as a chemical) risk i.e. where there is abrasion or puncture potential

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.

- ▶ Butyl rubber gloves
- ▶ Nitrile rubber gloves

Body protection

See Other protection below

- Overalls.
- Eyewash unit. Other protection
 - ► Barrier cream.
 - ► Skin cleansing cream.

Thermal hazards

Not Available

Respiratory protection

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

SECTION 9 PHYSICAL AND CHEMICAL PROPERTIES

Information on basic physical and chemical properties

Appearance	Text		
			,
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)	8.5	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 Available
Flash point (°C)	Not Available	Taste	Not Available
Evaporation rate	Not Available	Explosive properties	Not Available
Flammability	Not Available	Oxidising properties	Not Available
Upper Explosive Limit (%)	Not Available	Surface Tension (dyn/cm or mN/m)	Not Available
Lower Explosive Limit (%)	Not Available	Volatile Component (%vol)	Not Available
Vapour pressure (kPa)	Not Available	Gas group	Not Available
Solubility in water (g/L)	Immiscible	pH as a solution (1%)	Not Available
Vapour density (Air = 1)	Not Available	VOC g/L	Not Available

SECTION 10 STABILITY AND REACTIVITY

Reactivity	See section 7
Chemical stability	 Unstable in the presence of incompatible materials. Product is considered stable. Hazardous polymerisation will not occur.
Possibility of hazardous reactions	See section 7
Conditions to avoid	See section 7
Incompatible materials	See section 7
Hazardous decomposition products	See section 5

SECTION 11 TOXICOLOGICAL INFORMATION

Page **7** of **12** Issue Date: 02/07/2017 Version No: 1.3 Print Date: 02/07/2017 **Trouble Shooter Acr.-Neutral Base 45195**

Information on toxicological effects

ormation on toxicologic					
Inhaled	Inhalation of vapours or aerosols (mists, fumes), generated by the material during the course of normal handling, may produce toxic effects. The material is not thought to produce respiratory irritation (as classified by EC Directives using animal models). Nevertheless inhalation of vapours, fumes or aerosols, especially for prolonged periods, may produce respiratory discomfort and occasionally, distress.				
	The material is not thought to produce adverse health effects fo systemic effects have been produced following exposure of anim minimum.	• • •	,		
	Ingestion of propylene glycol produced reversible central nervou heart-rate (tachycardia), excessive sweating (diaphoresis) and giant of city items are protein	, ,	, ·		
Ingestion	ingredient of vitamin preparation. Excessive repeated ingestions may cause hypoglycaemia (low l	evels of glucose in the blood stream) among susce	eptible individuals; this may result in muscul		
	weakness, incoordination and mental confusion. Very high doses given during feeding studies to rats and dogs produce central nervous system depression (although one-third of that produced by ethanol),				
	haemolysis and insignificant kidney changes. In humans propylene glycol is partly excreted unchanged in the	urine and partly metabolised as lactic and pyruvic a	acid. Lactic acidosis may result.		
	Taken by mouth, isothiazolinones have moderate to high toxicity.	Taken by mouth, isothiazolinones have moderate to high toxicity. The major signs of toxicity are severe stomach irritation, lethargy, and inco-ordination.			
	The liquid may be miscible with fats or oils and may degrease the skin, producing a skin reaction described as non-allergic contact dermatitis. The material is unlikely to produce an irritant dermatitis as described in EC Directives.				
Skin Contact	Solutions of isothiazolinones may be irritating or even damaging		ntration of over 0.1% can irritate, and over		
	0.5% can cause severe irritation. Open cuts, abraded or irritated skin should not be exposed to thi	s material			
	Entry into the blood-stream, through, for example, cuts, abrasion of the material and ensure that any external damage is suitably p		mful effects. Examine the skin prior to the us		
	Although the liquid is not thought to be an irritant (as classified to by tearing or conjunctival redness (as with windburn).	by EC Directives), direct contact with the eye may	produce transient discomfort characterised		
Eye	Solutions containing isothiazolinones may damage the mucous i cause irritation, while higher levels (3-5.5%) produced severe in	ũ .	ery low concentrations (under 0.1%) did not		
	There has been concern that this material can cause cancer or n Skin contact with the material is more likely to cause a sensitisat	,			
	Toxic: danger of serious damage to health by prolonged exposu This material can cause serious damage if one is exposed to it f	re through inhalation.			
	defects.				
	Substance accumulation, in the human body, may occur and may The isothiazolinones are known contact sensitisers. Sensitisation	9 ,	·		
	The isothiazolinones are known contact sensitisers. Sensitisation is more likely with the chlorinated species as opposed to the non-chlorinated species. Propylene glycol is though, by some, to be a sensitising principal following the regular use of topical creams by eczema patients. A study of 866 persons using a formulation containing propylene glycol in a patch test indicated that propylene glycol caused primary irritation in 16% of exposed individuals probably caused				
	a formulation containing propylene glycol in a patch test indicate		16% of exposed individuals probably cause		
	by dehydration. Undiluted propylene glycol was tested on 1556 p	d that propylene glycol caused primary irritation in persons in a 24 hour patch test. 12.5% showed rear	ctions which were largely toxic (70%) or		
Chronic		d that propylene glycol caused primary irritation in persons in a 24 hour patch test. 12.5% showed rear mum on the second day or later. Reactions were	ctions which were largely toxic (70%) or seasonal in nature ranging from 17.8% in		
Chronic	by dehydration. Undiluted propylene glycol was tested on 1556 pallergic in nature (30%). Reaction responses reached their maximiter to 9.2% in other seasons. In a patch-test using 25 standaresponse. 84 subjects were patch tested using 100% propylene	d that propylene glycol caused primary irritation in bersons in a 24 hour patch test. 12.5% showed real mum on the second day or later. Reactions were rd allergens conducted on 500 individuals, propyle glycol. as well as 2% and 5% in water. With undilu	ctions which were largely toxic (70%) or seasonal in nature ranging from 17.8% in one glycol ranked fourth in sensitising ted material, 15% demonstrated a reaction		
Chronic	by dehydration. Undiluted propylene glycol was tested on 1556 p allergic in nature (30%). Reaction responses reached their maxi winter to 9.2% in other seasons. In a patch-test using 25 standa response. 84 subjects were patch tested using 100% propylene with 40% of the reactions being allergic in nature and 60% being Undiluted propylene glycol tested on the skin of man produced n	d that propylene glycol caused primary irritation in versons in a 24 hour patch test. 12.5% showed real mum on the second day or later. Reactions were tright allergens conducted on 500 individuals, propyle glycol. as well as 2% and 5% in water. With undiful g irritant. In dilute solutions 5 of 248 subjects exhib o irritation under open conditions but when applied	ctions which were largely toxic (70%) or seasonal in nature ranging from 17.8% in one glycol ranked fourth in sensitising ted material, 15% demonstrated a reaction, bited a reaction.		
Chronic	by dehydration. Undiluted propylene glycol was tested on 1556 p allergic in nature (30%). Reaction responses reached their maxi winter to 9.2% in other seasons. In a patch-test using 25 standa response. 84 subjects were patch tested using 100% propylene with 40% of the reactions being allergic in nature and 60% being	d that propylene glycol caused primary irritation in bersons in a 24 hour patch test. 12.5% showed rear mum on the second day or later. Reactions were a rd allergens conducted on 500 individuals, propyle glycol. as well as 2% and 5% in water. With undilu g irritant. In dilute solutions 5 of 248 subjects exhit o irritation under open conditions but when applied to sweat retention and weak primary irritation.	ctions which were largely toxic (70%) or seasonal in nature ranging from 17.8% in ene glycol ranked fourth in sensitising ted material, 15% demonstrated a reaction, pited a reaction. I under occlusive conditions, for 2 weeks, it		
Chronic	by dehydration. Undiluted propylene glycol was tested on 1556 pallergic in nature (30%). Reaction responses reached their maximiner to 9.2% in other seasons. In a patch-test using 25 standaresponse. 84 subjects were patch tested using 100% propylene with 40% of the reactions being allergic in nature and 60% being Undiluted propylene glycol tested on the skin of man produced no produced severe erythema, oedema and vesicles, probably due 1 Predictive contact skin sensitisation tests indicate that propylene Groups of cats fed 5 gm/kg/day of propylene glycol for 14 week	d that propylene glycol caused primary irritation in versons in a 24 hour patch test. 12.5% showed rear mum on the second day or later. Reactions were rd allergens conducted on 500 individuals, propyle glycol. as well as 2% and 5% in water. With undilu g irritant. In dilute solutions 5 of 248 subjects exhib o irritation under open conditions but when applied o sweat retention and weak primary irritation. e glycol is an intermediate grade sensitiser with an s showed a significant dose-related increase in red	ctions which were largely toxic (70%) or seasonal in nature ranging from 17.8% in ene glycol ranked fourth in sensitising ted material, 15% demonstrated a reaction, bited a reaction. I under occlusive conditions, for 2 weeks, it index of 1% of tested subjects. I blood cell Heinz body formation without an		
Chronic	by dehydration. Undiluted propylene glycol was tested on 1556 pallergic in nature (30%). Reaction responses reached their maximiner to 9.2% in other seasons. In a patch-test using 25 standaresponse. 84 subjects were patch tested using 100% propylene with 40% of the reactions being allergic in nature and 60% bein. Undiluted propylene glycol tested on the skin of man produced no produced severe erythema, oedema and vesicles, probably due to Predictive contact skin sensitisation tests indicate that propylene Groups of cats fed 5 gm/kg/day of propylene glycol for 14 week marked signs of haemolytic anaemia. The no-effect-level for cats degenerative change. Groups of rats dosed orally with 0.5 or 10 in 10	d that propylene glycol caused primary irritation in bersons in a 24 hour patch test. 12.5% showed real mum on the second day or later. Reactions were at dialergens conducted on 500 individuals, propyle glycol. as well as 2% and 5% in water. With undiful g irritant. In dilute solutions 5 of 248 subjects exhib o irritation under open conditions but when applied os weat retention and weak primary irritation. g glycol is an intermediate grade sensitiser with an s showed a significant dose-related increase in red is without formation of Heinz bodies is 100-500 ml/k ng/kg/day for 12 weeks had lowered food intake bu	ctions which were largely toxic (70%) or seasonal in nature ranging from 17.8% in one glycol ranked fourth in sensitising ted material, 15% demonstrated a reaction, bited a reaction. I under occlusive conditions, for 2 weeks, it index of 1% of tested subjects. I blood cell Heinz body formation without an up. There is no evidence of anaemia or		
Chronic	by dehydration. Undiluted propylene glycol was tested on 1556 pallergic in nature (30%). Reaction responses reached their maximiner to 9.2% in other seasons. In a patch-test using 25 standaresponse. 84 subjects were patch tested using 100% propylene with 40% of the reactions being allergic in nature and 60% bein. Undiluted propylene glycol tested on the skin of man produced no produced severe erythema, oedema and vesicles, probably due to Predictive contact skin sensitisation tests indicate that propylene Groups of cats fed 5 gm/kg/day of propylene glycol for 14 week marked signs of haemolytic anaemia. The no-effect-level for cats	d that propylene glycol caused primary irritation in bersons in a 24 hour patch test. 12.5% showed real mum on the second day or later. Reactions were at dialergens conducted on 500 individuals, propyle glycol. as well as 2% and 5% in water. With undiful g irritant. In dilute solutions 5 of 248 subjects exhib o irritation under open conditions but when applied os weat retention and weak primary irritation. g glycol is an intermediate grade sensitiser with an s showed a significant dose-related increase in red is without formation of Heinz bodies is 100-500 ml/k ng/kg/day for 12 weeks had lowered food intake bu	ctions which were largely toxic (70%) or seasonal in nature ranging from 17.8% in one glycol ranked fourth in sensitising ted material, 15% demonstrated a reaction, bited a reaction. I under occlusive conditions, for 2 weeks, it index of 1% of tested subjects. I blood cell Heinz body formation without an up. There is no evidence of anaemia or		
Chronic Trouble Shooter	by dehydration. Undiluted propylene glycol was tested on 1556 pallergic in nature (30%). Reaction responses reached their maximiner to 9.2% in other seasons. In a patch-test using 25 standaresponse. 84 subjects were patch tested using 100% propylene with 40% of the reactions being allergic in nature and 60% bein. Undiluted propylene glycol tested on the skin of man produced no produced severe erythema, oedema and vesicles, probably due to Predictive contact skin sensitisation tests indicate that propylene Groups of cats fed 5 gm/kg/day of propylene glycol for 14 week marked signs of haemolytic anaemia. The no-effect-level for cats degenerative change. Groups of rats dosed orally with 0.5 or 10 in 10	d that propylene glycol caused primary irritation in bersons in a 24 hour patch test. 12.5% showed real mum on the second day or later. Reactions were at dialergens conducted on 500 individuals, propyle glycol. as well as 2% and 5% in water. With undiful g irritant. In dilute solutions 5 of 248 subjects exhib o irritation under open conditions but when applied os weat retention and weak primary irritation. g glycol is an intermediate grade sensitiser with an s showed a significant dose-related increase in red is without formation of Heinz bodies is 100-500 ml/k ng/kg/day for 12 weeks had lowered food intake bu	ctions which were largely toxic (70%) or seasonal in nature ranging from 17.8% in one glycol ranked fourth in sensitising ted material, 15% demonstrated a reaction, bited a reaction. I under occlusive conditions, for 2 weeks, it index of 1% of tested subjects. I blood cell Heinz body formation without an u.g. There is no evidence of anaemia or		
	by dehydration. Undiluted propylene glycol was tested on 1556 pallergic in nature (30%). Reaction responses reached their maximinter to 9.2% in other seasons. In a patch-test using 25 standa response. 84 subjects were patch tested using 100% propylene with 40% of the reactions being allergic in nature and 60% being Undiluted propylene glycol tested on the skin of man produced no produced severe erythema, oedema and vesicles, probably due to Predictive contact skin sensitisation tests indicate that propylene Groups of cats fed 5 gm/kg/day of propylene glycol for 14 week marked signs of haemolytic anaemia. The no-effect-level for cats degenerative change. Groups of rats dosed orally with 0.5 or 10 rerythrocytes were more fragile. Heinz bodies were not apparent	d that propylene glycol caused primary irritation in persons in a 24 hour patch test. 12.5% showed rear mum on the second day or later. Reactions were a rd allergens conducted on 500 individuals, propyle glycol. as well as 2% and 5% in water. With undilu g irritant. In dilute solutions 5 of 248 subjects exhib o irritation under open conditions but when applied o sweat retention and weak primary irritation. glycol is an intermediate grade sensitiser with an s showed a significant dose-related increase in red is without formation of Heinz bodies is 100-500 ml/k ng/kg/day for 12 weeks had lowered food intake bu	ctions which were largely toxic (70%) or seasonal in nature ranging from 17.8% in one glycol ranked fourth in sensitising ted material, 15% demonstrated a reaction, bited a reaction. I under occlusive conditions, for 2 weeks, it index of 1% of tested subjects. I blood cell Heinz body formation without an g. There is no evidence of anaemia or		
Trouble Shooter	by dehydration. Undiluted propylene glycol was tested on 1556 pallergic in nature (30%). Reaction responses reached their maximinter to 9.2% in other seasons. In a patch-test using 25 standa response. 84 subjects were patch tested using 100% propylene with 40% of the reactions being allergic in nature and 60% being Undiluted propylene glycol tested on the skin of man produced not produced severe erythema, oedema and vesicles, probably due to Predictive contact skin sensitisation tests indicate that propylene Groups of cats fed 5 gm/kg/day of propylene glycol for 14 week marked signs of haemolytic anaemia. The no-effect-level for cats degenerative change. Groups of rats dosed orally with 0.5 or 10 rerythrocytes were more fragile. Heinz bodies were not apparent	d that propylene glycol caused primary irritation in persons in a 24 hour patch test. 12.5% showed rear mum on the second day or later. Reactions were a rear dallergens conducted on 500 individuals, propyle glycol. as well as 2% and 5% in water. With undilu g irritant. In dilute solutions 5 of 248 subjects exhib o irritation under open conditions but when applied o sweat retention and weak primary irritation. glycol is an intermediate grade sensitiser with an s showed a significant dose-related increase in red is without formation of Heinz bodies is 100-500 ml/k ng/kg/day for 12 weeks had lowered food intake bu IRRITATION Not Available	ctions which were largely toxic (70%) or seasonal in nature ranging from 17.8% in one glycol ranked fourth in sensitising ted material, 15% demonstrated a reaction, bited a reaction. I under occlusive conditions, for 2 weeks, it index of 1% of tested subjects. I blood cell Heinz body formation without an g. There is no evidence of anaemia or		
Trouble Shooter	by dehydration. Undiluted propylene glycol was tested on 1556 p allergic in nature (30%). Reaction responses reached their maximinter to 9.2% in other seasons. In a patch-test using 25 standa response. 84 subjects were patch tested using 100% propylene with 40% of the reactions being allergic in nature and 60% being Undiluted propylene glycol tested on the skin of man produced no produced severe erythema, oedema and vesicles, probably due to Predictive contact skin sensitisation tests indicate that propylene Groups of cats fed 5 gm/kg/day of propylene glycol for 14 week marked signs of haemolytic anaemia. The no-effect-level for cats degenerative change. Groups of rats dosed orally with 0.5 or 10 or Erythrocytes were more fragile. Heinz bodies were not apparent TOXICITY Not Available TOXICITY	d that propylene glycol caused primary irritation in persons in a 24 hour patch test. 12.5% showed rear mum on the second day or later. Reactions were dallergens conducted on 500 individuals, propyle glycol. as well as 2% and 5% in water. With undifu g irritant. In dilute solutions 5 of 248 subjects exhib o irritation under open conditions but when applied os sweat retention and weak primary irritation. glycol is an intermediate grade sensitiser with an s showed a significant dose-related increase in red is without formation of Heinz bodies is 100-500 ml/k mg/kg/day for 12 weeks had lowered food intake bu IRRITATION Not Available IRRITATION	ctions which were largely toxic (70%) or seasonal in nature ranging from 17.8% in one glycol ranked fourth in sensitising ted material, 15% demonstrated a reaction, bited a reaction. I under occlusive conditions, for 2 weeks, it index of 1% of tested subjects. I blood cell Heinz body formation without an g. There is no evidence of anaemia or		
Trouble Shooter AcrNeutral Base 45195	by dehydration. Undiluted propylene glycol was tested on 1556 p allergic in nature (30%). Reaction responses reached their maximinter to 9.2% in other seasons. In a patch-test using 25 standa response. 84 subjects were patch tested using 100% propylene with 40% of the reactions being allergic in nature and 60% being Undiluted propylene glycol tested on the skin of man produced no produced severe erythema, oedema and vesicles, probably due to Predictive contact skin sensitisation tests indicate that propylene Groups of cats fed 5 gm/kg/day of propylene glycol for 14 week marked signs of haemolytic anaemia. The no-effect-level for cats degenerative change. Groups of rats dosed orally with 0.5 or 10 in Erythrocytes were more fragile. Heinz bodies were not apparent TOXICITY Not Available TOXICITY Dermal (rabbit) LD50: >2000 mg/kg ^[1]	d that propylene glycol caused primary irritation in persons in a 24 hour patch test. 12.5% showed rear mum on the second day or later. Reactions were a rd allergens conducted on 500 individuals, propyle glycol. as well as 2% and 5% in water. With undilu g irritant. In dilute solutions 5 of 248 subjects exhit o irritation under open conditions but when applied o sweat retention and weak primary irritation. g glycol is an intermediate grade sensitiser with an s showed a significant dose-related increase in red s without formation of Heinz bodies is 100-500 ml/k ng/kg/day for 12 weeks had lowered food intake bu IRRITATION Not Available IRRITATION Eye (rabbit): 100 mg - mild	ctions which were largely toxic (70%) or seasonal in nature ranging from 17.8% in ene glycol ranked fourth in sensitising ted material, 15% demonstrated a reaction, pited a reaction. I under occlusive conditions, for 2 weeks, it index of 1% of tested subjects. I blood cell Heinz body formation without an eg. There is no evidence of anaemia or at no adverse effects on body weights.		
Trouble Shooter	by dehydration. Undiluted propylene glycol was tested on 1556 p allergic in nature (30%). Reaction responses reached their maximinter to 9.2% in other seasons. In a patch-test using 25 standa response. 84 subjects were patch tested using 100% propylene with 40% of the reactions being allergic in nature and 60% being Undiluted propylene glycol tested on the skin of man produced no produced severe erythema, oedema and vesicles, probably due to Predictive contact skin sensitisation tests indicate that propylene Groups of cats fed 5 gm/kg/day of propylene glycol for 14 week marked signs of haemolytic anaemia. The no-effect-level for cats degenerative change. Groups of rats dosed orally with 0.5 or 10 or Erythrocytes were more fragile. Heinz bodies were not apparent TOXICITY Not Available TOXICITY	d that propylene glycol caused primary irritation in persons in a 24 hour patch test. 12.5% showed rear mum on the second day or later. Reactions were a rear dallergens conducted on 500 individuals, propyle glycol. as well as 2% and 5% in water. With undilu g irritant. In dilute solutions 5 of 248 subjects exhib o irritation under open conditions but when applied o sweat retention and weak primary irritation. glycol is an intermediate grade sensitiser with an s showed a significant dose-related increase in red s without formation of Heinz bodies is 100-500 ml/k ng/kg/day for 12 weeks had lowered food intake bu IRRITATION Not Available IRRITATION Eye (rabbit): 100 mg - mild Eye (rabbit): 500 mg/24h - mild	ctions which were largely toxic (70%) or seasonal in nature ranging from 17.8% in ene glycol ranked fourth in sensitising ted material, 15% demonstrated a reaction, ited a reaction. I under occlusive conditions, for 2 weeks, it index of 1% of tested subjects. I blood cell Heinz body formation without an g. There is no evidence of anaemia or it no adverse effects on body weights.		
Trouble Shooter AcrNeutral Base 45195	by dehydration. Undiluted propylene glycol was tested on 1556 p allergic in nature (30%). Reaction responses reached their maximinter to 9.2% in other seasons. In a patch-test using 25 standa response. 84 subjects were patch tested using 100% propylene with 40% of the reactions being allergic in nature and 60% being Undiluted propylene glycol tested on the skin of man produced no produced severe erythema, oedema and vesicles, probably due to Predictive contact skin sensitisation tests indicate that propylene Groups of cats fed 5 gm/kg/day of propylene glycol for 14 week marked signs of haemolytic anaemia. The no-effect-level for cats degenerative change. Groups of rats dosed orally with 0.5 or 10 in Erythrocytes were more fragile. Heinz bodies were not apparent TOXICITY Not Available TOXICITY Dermal (rabbit) LD50: >2000 mg/kg ^[1]	d that propylene glycol caused primary irritation in persons in a 24 hour patch test. 12.5% showed rear mum on the second day or later. Reactions were d allergens conducted on 500 individuals, propyle glycol. as well as 2% and 5% in water. With undifu g irritant. In dilute solutions 5 of 248 subjects exhib o irritation under open conditions but when applied os sweat retention and weak primary irritation. glycol is an intermediate grade sensitiser with an s showed a significant dose-related increase in red s without formation of Heinz bodies is 100-500 ml/k mg/kg/day for 12 weeks had lowered food intake bu IRRITATION Not Available IRRITATION Eye (rabbit): 100 mg - mild Eye (rabbit): 500 mg/24h - mild Skin(human):104 mg/3d Intermi	ctions which were largely toxic (70%) or seasonal in nature ranging from 17.8% in see glycol ranked fourth in sensitising ted material, 15% demonstrated a reaction, litted a reaction. I under occlusive conditions, for 2 weeks, it index of 1% of tested subjects. I blood cell Heinz body formation without an g. There is no evidence of anaemia or at no adverse effects on body weights.		
Trouble Shooter AcrNeutral Base 45195	by dehydration. Undiluted propylene glycol was tested on 1556 p allergic in nature (30%). Reaction responses reached their maximinter to 9.2% in other seasons. In a patch-test using 25 standa response. 84 subjects were patch tested using 100% propylene with 40% of the reactions being allergic in nature and 60% being Undiluted propylene glycol tested on the skin of man produced no produced severe erythema, oedema and vesicles, probably due to Predictive contact skin sensitisation tests indicate that propylene Groups of cats fed 5 gm/kg/day of propylene glycol for 14 week marked signs of haemolytic anaemia. The no-effect-level for cats degenerative change. Groups of rats dosed orally with 0.5 or 10 in Erythrocytes were more fragile. Heinz bodies were not apparent TOXICITY Not Available TOXICITY Dermal (rabbit) LD50: >2000 mg/kg ^[1]	d that propylene glycol caused primary irritation in persons in a 24 hour patch test. 12.5% showed rear mum on the second day or later. Reactions were a rear dallergens conducted on 500 individuals, propyle glycol. as well as 2% and 5% in water. With undilu g irritant. In dilute solutions 5 of 248 subjects exhib o irritation under open conditions but when applied o sweat retention and weak primary irritation. glycol is an intermediate grade sensitiser with an s showed a significant dose-related increase in red s without formation of Heinz bodies is 100-500 ml/k ng/kg/day for 12 weeks had lowered food intake bu IRRITATION Not Available IRRITATION Eye (rabbit): 100 mg - mild Eye (rabbit): 500 mg/24h - mild	ctions which were largely toxic (70%) or seasonal in nature ranging from 17.8% in see glycol ranked fourth in sensitising ted material, 15% demonstrated a reaction, litted a reaction. I under occlusive conditions, for 2 weeks, it index of 1% of tested subjects. I blood cell Heinz body formation without an g. There is no evidence of anaemia or at no adverse effects on body weights.		
Trouble Shooter AcrNeutral Base 45195	by dehydration. Undiluted propylene glycol was tested on 1556 p allergic in nature (30%). Reaction responses reached their maximinter to 9.2% in other seasons. In a patch-test using 25 standa response. 84 subjects were patch tested using 100% propylene with 40% of the reactions being allergic in nature and 60% being Undiluted propylene glycol tested on the skin of man produced no produced severe erythema, oedema and vesicles, probably due to Predictive contact skin sensitisation tests indicate that propylene Groups of cats fed 5 gm/kg/day of propylene glycol for 14 week marked signs of haemolytic anaemia. The no-effect-level for cats degenerative change. Groups of rats dosed orally with 0.5 or 10 in Erythrocytes were more fragile. Heinz bodies were not apparent TOXICITY Not Available TOXICITY Dermal (rabbit) LD50: >2000 mg/kg ^[1]	d that propylene glycol caused primary irritation in persons in a 24 hour patch test. 12.5% showed rear mum on the second day or later. Reactions were d allergens conducted on 500 individuals, propyle glycol. as well as 2% and 5% in water. With undifu g irritant. In dilute solutions 5 of 248 subjects exhib o irritation under open conditions but when applied os sweat retention and weak primary irritation. glycol is an intermediate grade sensitiser with an s showed a significant dose-related increase in red s without formation of Heinz bodies is 100-500 ml/k mg/kg/day for 12 weeks had lowered food intake bu IRRITATION Not Available IRRITATION Eye (rabbit): 100 mg - mild Eye (rabbit): 500 mg/24h - mild Skin(human):104 mg/3d Intermi	ctions which were largely toxic (70%) or seasonal in nature ranging from 17.8% in see glycol ranked fourth in sensitising ted material, 15% demonstrated a reaction, litted a reaction. I under occlusive conditions, for 2 weeks, it index of 1% of tested subjects. I blood cell Heinz body formation without an g. There is no evidence of anaemia or at no adverse effects on body weights.		
Trouble Shooter AcrNeutral Base 45195	by dehydration. Undiluted propylene glycol was tested on 1556 p allergic in nature (30%). Reaction responses reached their maximinter to 9.2% in other seasons. In a patch-test using 25 standa response. 84 subjects were patch tested using 100% propylene with 40% of the reactions being allergic in nature and 60% being Undiluted propylene glycol tested on the skin of man produced no produced severe erythema, oedema and vesicles, probably due to Predictive contact skin sensitisation tests indicate that propylene Groups of cats fed 5 gm/kg/day of propylene glycol for 14 week marked signs of haemolytic anaemia. The no-effect-level for cats degenerative change. Groups of rats dosed orally with 0.5 or 10 m Erythrocytes were more fragile. Heinz bodies were not apparent TOXICITY Not Available TOXICITY Dermal (rabbit) LD50: >2000 mg/kg ^[1] Oral (rat) LD50: 20000 mg/kg ^[2]	d that propylene glycol caused primary irritation in persons in a 24 hour patch test. 12.5% showed rear mum on the second day or later. Reactions were d allergens conducted on 500 individuals, propyle glycol. as well as 2% and 5% in water. With undifu g irritant. In dilute solutions 5 of 248 subjects exhib o irritation under open conditions but when applied os sweat retention and weak primary irritation. glycol is an intermediate grade sensitiser with an s showed a significant dose-related increase in red s without formation of Heinz bodies is 100-500 ml/k mg/kg/day for 12 weeks had lowered food intake bu IRRITATION Not Available IRRITATION Eye (rabbit): 100 mg - mild Eye (rabbit): 500 mg/24h - mild Skin(human):104 mg/3d Intermi	ctions which were largely toxic (70%) or seasonal in nature ranging from 17.8% in sea glycol ranked fourth in sensitising ted material, 15% demonstrated a reaction, littled a reaction. I under occlusive conditions, for 2 weeks, it index of 1% of tested subjects. I blood cell Heinz body formation without an g. There is no evidence of anaemia or it no adverse effects on body weights.		
Trouble Shooter AcrNeutral Base 45195	by dehydration. Undiluted propylene glycol was tested on 1556 pallergic in nature (30%). Reaction responses reached their maximinter to 9.2% in other seasons. In a patch-test using 25 standa response. 84 subjects were patch tested using 100% propylene with 40% of the reactions being allergic in nature and 60% being Undiluted propylene glycol tested on the skin of man produced no produced severe erythema, oedema and vesicles, probably due to Predictive contact skin sensitisation tests indicate that propylene Groups of cats fed 5 gm/kg/day of propylene glycol for 14 week marked signs of haemolytic anaemia. The no-effect-level for cats degenerative change. Groups of rats dosed orally with 0.5 or 10 regythrocytes were more fragile. Heinz bodies were not apparent TOXICITY Not Available TOXICITY Dermal (rabbit) LD50: >2000 mg/kg ^[1] Oral (rat) LD50: 20000 mg/kg ^[2]	d that propylene glycol caused primary irritation in persons in a 24 hour patch test. 12.5% showed rear mum on the second day or later. Reactions were d allergens conducted on 500 individuals, propyle glycol. as well as 2% and 5% in water. With undifu g irritant. In dilute solutions 5 of 248 subjects exhib o irritation under open conditions but when applied os sweat retention and weak primary irritation. glycol is an intermediate grade sensitiser with an s showed a significant dose-related increase in red s without formation of Heinz bodies is 100-500 ml/k mg/kg/day for 12 weeks had lowered food intake bu IRRITATION Not Available IRRITATION Eye (rabbit): 100 mg - mild Eye (rabbit): 500 mg/24h - mild Skin(human):104 mg/3d Intermi	ctions which were largely toxic (70%) or seasonal in nature ranging from 17.8% in see glycol ranked fourth in sensitising ted material, 15% demonstrated a reaction litted a reaction. I under occlusive conditions, for 2 weeks, it index of 1% of tested subjects. I blood cell Heinz body formation without arrig. There is no evidence of anaemia or at no adverse effects on body weights.		
Trouble Shooter AcrNeutral Base 45195 propylene glycol	by dehydration. Undiluted propylene glycol was tested on 1556 pallergic in nature (30%). Reaction responses reached their maximinter to 9.2% in other seasons. In a patch-test using 25 standa response. 84 subjects were patch tested using 100% propylene with 40% of the reactions being allergic in nature and 60% being Undiluted propylene glycol tested on the skin of man produced no produced severe erythema, oedema and vesicles, probably due to Predictive contact skin sensitisation tests indicate that propylene Groups of cats fed 5 gm/kg/day of propylene glycol for 14 week marked signs of haemolytic anaemia. The no-effect-level for cats degenerative change. Groups of rats dosed orally with 0.5 or 10 regythrocytes were more fragile. Heinz bodies were not apparent TOXICITY Not Available TOXICITY Dermal (rabbit) LD50: >2000 mg/kg ^[1] Oral (rat) LD50: 20000 mg/kg ^[2] Inhalation (rat) LC50: 0.1 mg/l/4h. ^[2]	d that propylene glycol caused primary irritation in persons in a 24 hour patch test. 12.5% showed rear mum on the second day or later. Reactions were d allergens conducted on 500 individuals, propyle glycol. as well as 2% and 5% in water. With undifu g irritant. In dilute solutions 5 of 248 subjects exhib o irritation under open conditions but when applied os sweat retention and weak primary irritation. glycol is an intermediate grade sensitiser with an s showed a significant dose-related increase in red s without formation of Heinz bodies is 100-500 ml/k mg/kg/day for 12 weeks had lowered food intake bu IRRITATION Not Available IRRITATION Eye (rabbit): 100 mg - mild Eye (rabbit): 500 mg/24h - mild Skin(human):104 mg/3d Intermi	ctions which were largely toxic (70%) or seasonal in nature ranging from 17.8% in one glycol ranked fourth in sensitising ted material, 15% demonstrated a reaction, litted a reaction. I under occlusive conditions, for 2 weeks, it index of 1% of tested subjects. I blood cell Heinz body formation without an g. There is no evidence of anaemia or at no adverse effects on body weights.		
Trouble Shooter AcrNeutral Base 45195 propylene glycol	by dehydration. Undiluted propylene glycol was tested on 1556 p allergic in nature (30%). Reaction responses reached their maximinter to 9.2% in other seasons. In a patch-test using 25 standa response. 84 subjects were patch tested using 100% propylene with 40% of the reactions being allergic in nature and 60% being Undiluted propylene glycol tested on the skin of man produced no produced severe erythema, oedema and vesicles, probably due to Predictive contact skin sensitisation tests indicate that propylene Groups of cats fed 5 gm/kg/day of propylene glycol for 14 week marked signs of haemolytic anaemia. The no-effect-level for cats degenerative change. Groups of rats dosed orally with 0.5 or 10 r. Erythrocytes were more fragile. Heinz bodies were not apparent TOXICITY Not Available TOXICITY Dermal (rabbit) LD50: >20000 mg/kg ^[2] TOXICITY dermal (rat) LD50: >2500 mg/kg ^[2]	d that propylene glycol caused primary irritation in persons in a 24 hour patch test. 12.5% showed rear mum on the second day or later. Reactions were d allergens conducted on 500 individuals, propyle glycol. as well as 2% and 5% in water. With undifu g irritant. In dilute solutions 5 of 248 subjects exhib o irritation under open conditions but when applied os sweat retention and weak primary irritation. glycol is an intermediate grade sensitiser with an s showed a significant dose-related increase in red s without formation of Heinz bodies is 100-500 ml/k mg/kg/day for 12 weeks had lowered food intake bu IRRITATION Not Available IRRITATION Eye (rabbit): 100 mg - mild Eye (rabbit): 500 mg/24h - mild Skin(human):104 mg/3d Intermi	ctions which were largely toxic (70%) or seasonal in nature ranging from 17.8% in one glycol ranked fourth in sensitising ted material, 15% demonstrated a reaction, litted a reaction. I under occlusive conditions, for 2 weeks, it index of 1% of tested subjects. I blood cell Heinz body formation without an g. There is no evidence of anaemia or at no adverse effects on body weights.		
Trouble Shooter AcrNeutral Base 45195 propylene glycol	by dehydration. Undiluted propylene glycol was tested on 1556 pallergic in nature (30%). Reaction responses reached their maximinter to 9.2% in other seasons. In a patch-test using 25 standa response. 84 subjects were patch tested using 100% propylene with 40% of the reactions being allergic in nature and 60% bein Undiluted propylene glycol tested on the skin of man produced norduced severe erythema, oedema and vesicles, probably due to Predictive contact skin sensitisation tests indicate that propylene Groups of cats fed 5 gm/kg/day of propylene glycol for 14 week marked signs of haemolytic anaemia. The no-effect-level for cats degenerative change. Groups of rats dosed orally with 0.5 or 10 negretory to the context of the context	d that propylene glycol caused primary irritation in persons in a 24 hour patch test. 12.5% showed rearmum on the second day or later. Reactions were at allergens conducted on 500 individuals, propyle glycol. as well as 2% and 5% in water. With undilug irritant. In dilute solutions 5 of 248 subjects exhib o irritation under open conditions but when applied os sweat retention and weak primary irritation. In glycol is an intermediate grade sensitiser with an as showed a significant dose-related increase in red is without formation of Heinz bodies is 100-500 ml/k ing/kg/day for 12 weeks had lowered food intake but in the properties of th	ctions which were largely toxic (70%) or seasonal in nature ranging from 17.8% in one glycol ranked fourth in sensitising ted material, 15% demonstrated a reaction, litted a reaction. I under occlusive conditions, for 2 weeks, it index of 1% of tested subjects. I blood cell Heinz body formation without an g. There is no evidence of anaemia or at no adverse effects on body weights.		
Trouble Shooter AcrNeutral Base 45195 propylene glycol chlorothalonil	by dehydration. Undiluted propylene glycol was tested on 1556 pallergic in nature (30%). Reaction responses reached their maximinter to 9.2% in other seasons. In a patch-test using 25 standa response. 84 subjects were patch tested using 100% propylene with 40% of the reactions being allergic in nature and 60% being Undiluted propylene glycol tested on the skin of man produced no produced severe erythema, oedema and vesicles, probably due to Predictive contact skin sensitisation tests indicate that propylene Groups of cats fed 5 gm/kg/day of propylene glycol for 14 week marked signs of haemolytic anaemia. The no-effect-level for cats degenerative change. Groups of rats dosed orally with 0.5 or 10 rerythrocytes were more fragile. Heinz bodies were not apparent TOXICITY Not Available TOXICITY Dermal (rabbit) LD50: >20000 mg/kg ^[2] TOXICITY dermal (rat) LD50: >25000 mg/kg ^[2] Inhalation (rat) LC50: 0.1 mg/l/4h. [2] Inhalation (rat) LC50: 0.31 mg/L/1hr ^[2] Oral (rat) LD50: 10000 mg/kg ^[2]	d that propylene glycol caused primary irritation in ersons in a 24 hour patch test. 12.5% showed reamum on the second day or later. Reactions were at allergens conducted on 500 individuals, propyle glycol. as well as 2% and 5% in water. With undilug irritant. In dilute solutions 5 of 248 subjects exhib o irritation under open conditions but when applied os sweat retention and weak primary irritation. If glycol is an intermediate grade sensitiser with an as showed a significant dose-related increase in red is without formation of Heinz bodies is 100-500 ml/k mg/kg/day for 12 weeks had lowered food intake but a significant was a showed a significant dose-related increase in red is without formation of Heinz bodies is 100-500 ml/k mg/kg/day for 12 weeks had lowered food intake but a significant was a showed a significant dose-related increase in red is without formation of Heinz bodies is 100-500 ml/k mg/kg/day for 12 weeks had lowered food intake but a skinch of the significant of the si	ctions which were largely toxic (70%) or seasonal in nature ranging from 17.8% in one glycol ranked fourth in sensitising ted material, 15% demonstrated a reaction, litted a reaction. I under occlusive conditions, for 2 weeks, it index of 1% of tested subjects. I blood cell Heinz body formation without an g. There is no evidence of anaemia or at no adverse effects on body weights.		
Trouble Shooter AcrNeutral Base 45195 propylene glycol	by dehydration. Undiluted propylene glycol was tested on 1556 pallergic in nature (30%). Reaction responses reached their maximinter to 9.2% in other seasons. In a patch-test using 25 standa response. 84 subjects were patch tested using 100% propylene with 40% of the reactions being allergic in nature and 60% bein Undiluted propylene glycol tested on the skin of man produced norduced severe erythema, oedema and vesicles, probably due to Predictive contact skin sensitisation tests indicate that propylene Groups of cats fed 5 gm/kg/day of propylene glycol for 14 week marked signs of haemolytic anaemia. The no-effect-level for cats degenerative change. Groups of rats dosed orally with 0.5 or 10 negretory to the context of the context	d that propylene glycol caused primary irritation in persons in a 24 hour patch test. 12.5% showed rearmum on the second day or later. Reactions were at allergens conducted on 500 individuals, propyle glycol. as well as 2% and 5% in water. With undilug irritant. In dilute solutions 5 of 248 subjects exhib o irritation under open conditions but when applied os sweat retention and weak primary irritation. In glycol is an intermediate grade sensitiser with an as showed a significant dose-related increase in red is without formation of Heinz bodies is 100-500 ml/k ing/kg/day for 12 weeks had lowered food intake but in the properties of th	ctions which were largely toxic (70%) or seasonal in nature ranging from 17.8% in one glycol ranked fourth in sensitising ted material, 15% demonstrated a reaction, litted a reaction. I under occlusive conditions, for 2 weeks, it index of 1% of tested subjects. I blood cell Heinz body formation without an g. There is no evidence of anaemia or at no adverse effects on body weights.		
Trouble Shooter AcrNeutral Base 45195 propylene glycol chlorothalonil	by dehydration. Undiluted propylene glycol was tested on 1556 pallergic in nature (30%). Reaction responses reached their maximinter to 9.2% in other seasons. In a patch-test using 25 standa response. 84 subjects were patch tested using 100% propylene with 40% of the reactions being allergic in nature and 60% bein Undiluted propylene glycol tested on the skin of man produced norduced severe erythema, oedema and vesicles, probably due to Predictive contact skin sensitisation tests indicate that propylene Groups of cats fed 5 gm/kg/day of propylene glycol for 14 week marked signs of haemolytic anaemia. The no-effect-level for cats degenerative change. Groups of rats dosed orally with 0.5 or 10 negretory to the context of the context	d that propylene glycol caused primary irritation in persons in a 24 hour patch test. 12.5% showed rearmum on the second day or later. Reactions were at allergens conducted on 500 individuals, propyle glycol. as well as 2% and 5% in water. With undilug irritant. In dilute solutions 5 of 248 subjects exhib o irritation under open conditions but when applied os sweat retention and weak primary irritation. In glycol is an intermediate grade sensitiser with an as showed a significant dose-related increase in red is without formation of Heinz bodies is 100-500 ml/k ing/kg/day for 12 weeks had lowered food intake but a significant dose-related increase in red is without formation of Heinz bodies is 100-500 ml/k ing/kg/day for 12 weeks had lowered food intake but in the second state of the second state	ctions which were largely toxic (70%) or seasonal in nature ranging from 17.8% in one glycol ranked fourth in sensitising ted material, 15% demonstrated a reaction, litted a reaction. I under occlusive conditions, for 2 weeks, it index of 1% of tested subjects. I blood cell Heinz body formation without an g. There is no evidence of anaemia or at no adverse effects on body weights.		
Trouble Shooter AcrNeutral Base 45195 propylene glycol chlorothalonil	by dehydration. Undiluted propylene glycol was tested on 1556 pallergic in nature (30%). Reaction responses reached their maximinter to 9.2% in other seasons. In a patch-test using 25 standa response. 84 subjects were patch tested using 100% propylene with 40% of the reactions being allergic in nature and 60% being Undiluted propylene glycol tested on the skin of man produced no produced severe erythema, oedema and vesicles, probably due to Predictive contact skin sensitisation tests indicate that propylene Groups of cats fed 5 gm/kg/day of propylene glycol for 14 week marked signs of haemolytic anaemia. The no-effect-level for cats degenerative change. Groups of rats dosed orally with 0.5 or 10 rerythrocytes were more fragile. Heinz bodies were not apparent TOXICITY Not Available TOXICITY Dermal (rabbit) LD50: >20000 mg/kg ^[2] TOXICITY dermal (rat) LD50: >25000 mg/kg ^[2] Inhalation (rat) LC50: 0.1 mg/l/4h. [2] Inhalation (rat) LC50: 0.31 mg/L/1hr ^[2] Oral (rat) LD50: 10000 mg/kg ^[2]	d that propylene glycol caused primary irritation in ersons in a 24 hour patch test. 12.5% showed reamum on the second day or later. Reactions were at allergens conducted on 500 individuals, propyle glycol. as well as 2% and 5% in water. With undilug irritant. In dilute solutions 5 of 248 subjects exhib o irritation under open conditions but when applied os sweat retention and weak primary irritation. If glycol is an intermediate grade sensitiser with an as showed a significant dose-related increase in red is without formation of Heinz bodies is 100-500 ml/k mg/kg/day for 12 weeks had lowered food intake but a significant was a showed a significant dose-related increase in red is without formation of Heinz bodies is 100-500 ml/k mg/kg/day for 12 weeks had lowered food intake but a significant was a showed a significant dose-related increase in red is without formation of Heinz bodies is 100-500 ml/k mg/kg/day for 12 weeks had lowered food intake but a skinch of the significant of the si	ctions which were largely toxic (70%) or seasonal in nature ranging from 17.8% in ene glycol ranked fourth in sensitising ted material, 15% demonstrated a reaction, ited a reaction. I under occlusive conditions, for 2 weeks, it index of 1% of tested subjects. I blood cell Heinz body formation without an g. There is no evidence of anaemia or it no adverse effects on body weights. It Mod I IRRITATION Not Available		

Chemwatch: 9-339815 Page 8 of 12 Issue Date: 02/07/2017
Version No: 1.3 Print Date: 02/07/2017

Trouble Shooter Acr.-Neutral Base 45195

Skin (rabbit): 500 mg/24 h- mild 1. Value obtained from Europe ECHA Registered Substances - Acute toxicity 2.* Value obtained from manufacturer's SDS. Unless otherwise specified data Leaend: extracted from RTECS - Register of Toxic Effect of chemical Substances The acute oral toxicity of propylene glycol is very low, and large quantities are required to cause perceptible health damage in humans. Serious toxicity PROPYLENE GLYCOL generally occurs only at plasma concentrations over 1 g/L, which requires extremely high intake over a relatively short period of time. It would be nearly impossible to reach toxic levels by consuming foods or supplements, which contain at most 1 g/kg of PG. Chlorothalonil has low toxicity, according to animal testing. It irritates the skin and eye. Animal testing suggests that at sufficient doses it can cause cancer of the kidney and forestomach. CHLOROTHALONIL WARNING: This substance has been classified by the IARC as Group 2B: Possibly Carcinogenic to Humans. ADI: 0.01 mg/kg/day NOEL: 1.5 mg/kg/day No significant acute toxicological data identified in literature search. 5-CHLORO-2-METHYL-The material may be irritating to the eye, with prolonged contact causing inflammation. Repeated or prolonged exposure to irritants may produce conjunctivitis. 4-ISOTHIAZOLIN-3-ONE NOTE: Substance has been shown to be mutagenic in at least one assay, or belongs to a family of chemicals producing damage or change to cellular DNA. Considered to be the major sensitiser in Kathon CG (1) (1). Bruze et al - Contact Dermatitis 20: 219-39, 1989 PROPYLENE GLYCOL & 5-CHLORO-2-METHYL-The material may cause skin irritation after prolonged or repeated exposure and may produce on contact skin redness, swelling, the production of vesicles, 4-ISOTHIAZOLIN-3-ONE & scaling and thickening of the skin. ZINC OXIDE **CHLOROTHALONIL &** The following information refers to contact allergens as a group and may not be specific to this product. 5-CHLORO-2-METHYL-Contact allergies quickly manifest themselves as contact eczema, more rarely as urticaria or Quincke's oedema. The pathogenesis of contact eczema involves 4-ISOTHIAZOLIN-3-ONE a cell-mediated (T lymphocytes) immune reaction of the delayed type. CHLOROTHALONIL & Asthma-like symptoms may continue for months or even years after exposure to the material ceases. This may be due to a non-allergenic condition known as 5-CHLORO-2-METHYLreactive airways dysfunction syndrome (RADS) which can occur following exposure to high levels of highly irritating compound. Key criteria for the diagnosis 4-ISOTHIAZOLIN-3-ONE of RADS include the absence of preceding respiratory disease, in a non-atopic individual, with abrupt onset of persistent asthma-like symptoms within minutes to hours of a documented exposure to the irritant. **Acute Toxicity** 0 Carcinogenicity Skin Irritation/Corrosion 0 Reproductivity 0 Serious Eye STOT - Single Exposure 0 Damage/Irritation Respiratory or Skin STOT - Repeated Exposure sensitisation

Aspiration Hazard

Legend:

X - Data available but does not fill the criteria for classification

Data available to make classification

Data Not Available to make classification

SECTION 12 ECOLOGICAL INFORMATION

0

Mutagenicity

Toxicity

Oxicity					
Ingredient	Endpoint	Test Duration (hr)	Species	Value	Source
propylene glycol	LC50	96	Fish	710mg/L	4
propylene glycol	EC50	48	Crustacea	>1000mg/L	4
propylene glycol	EC50	96	Algae or other aquatic plants	10905.921mg/L	3
propylene glycol	EC50	384	Crustacea	311.145mg/L	3
propylene glycol	NOEC	168	Fish	98mg/L	4
chlorothalonil	LC50	96	Fish	0.0076mg/L	4
chlorothalonil	EC50	48	Crustacea	0.0066475mg/L	4
chlorothalonil	EC50	72	Algae or other aquatic plants	0.0068mg/L	4
chlorothalonil	BCF	336	Algae or other aquatic plants	0.02mg/L	4
chlorothalonil	EC10	48	Crustacea	0.00055839mg/L	4
chlorothalonil	NOEC	240	Crustacea	0.0003mg/L	4
5-chloro-2-methyl- 4-isothiazolin-3-one	LC50	96	Fish	0.19mg/L	4
5-chloro-2-methyl- 4-isothiazolin-3-one	EC50	48	Crustacea	0.028mg/L	4
5-chloro-2-methyl- 4-isothiazolin-3-one	EC50	72	Algae or other aquatic plants	0.021mg/L	4
5-chloro-2-methyl- 4-isothiazolin-3-one	EC50	120	Algae or other aquatic plants	0.022mg/L	4
5-chloro-2-methyl- 4-isothiazolin-3-one	NOEC	504	Crustacea	0.172mg/L	1
zinc oxide	LC50	96	Fish	0.439mg/L	2
zinc oxide	EC50	48	Crustacea	0.105mg/L	2

Chemwatch: 9-339815 Page 9 of 12 Issue Date: 02/07/2017
Version No: 1.3 Print Date: 02/07/2017

Trouble Shooter Acr.-Neutral Base 45195

zinc oxide	EC50	72	Algae or other aquatic plants	0.042mg/L	4
zinc oxide	BCF	336	Fish	4376.673mg/L	4
zinc oxide	EC20	72	Algae or other aquatic plants	0.023mg/L	4
zinc oxide	NOEC	72	Algae or other aquatic plants	0.0049mg/L	2
	Extracted from 1. IUCLID Toxicity Data 2. Europe ECHA Registered Substances - Ecotoxicological Information - Aquatic Toxicity 3. EPIWIN Suite V3.12 - Aquatic Toxicity Data (Estimated) 4. US EPA, Ecotox database - Aquatic Toxicity Data 5. ECETOC Aquatic Hazard Assessment Data 6. NITE (Japan) - Bioconcentration Data 7. METI (Japan) - Bioconcentration Data 8. Vendor Data				

Toxic to aquatic organisms, may cause long-term adverse effects in the aquatic environment.

Do NOT allow product to come in contact with surface waters or to intertidal areas below the mean high water mark. Do not contaminate water when cleaning equipment or disposing of equipment wash-waters.

Wastes resulting from use of the product must be disposed of on site or at approved waste sites.

Propylene glycol is known to exert high levels of biochemical oxygen demand (BOD) during degradation in surface waters. This process can adversely affect aquatic life by consuming oxygen needed by aquatic organisms for survival. Large quantities of dissolved oxygen (DO) in the water column are consumed when microbial populations decompose propylene glycol. Sufficient dissolved oxygen levels in surface waters are critical for the survival of fish, macro-invertebrates, and other aquatic organisms. If oxygen concentrations drop below a minimum level, organisms emigrate, if able and possible, to areas with higher oxygen levels or eventually die. This effect can drastically reduce the amount of usable aquatic habitat. Reductions in DO levels can reduce or eliminate bottom-feeder populations, create conditions that favour a change in a community's species profile, or after critical food-web interactions.

log Kow: -1.41--0.3 Half-life (hr) air: 32 Henry's atm m3 /mol: 1.20E-08 BOD 5: 0.995,2.2% ThOD: 1.685 BCF: <1

Bioaccumulation: not sig processes Abiotic: photoxid

Environmental Fate: Isothiazolinones are antimicrobials used to control bacteria, fungi, and for wood preservation and antifouling agents. They are frequently used in personal care products such as shampoos and other hair care products, as well as certain paint formulations. The most common isothiazolinone combinations are 5-chloro-2-methyl-4-isothiazolin-3-one, (CMI), and 2-methyl-4-isothiazolin-3-one, (MI).

Aquatic Fate: 5-chloro-2-methyl-4-isothiazolin-3-one, (CMI), and 2-methyl-4-isothiazolin-3-one, (MI), undergo primary biological breakdown with half-lives of less than 24 hours in both oxygenated and low oxygen sediments with >55% breakdown occurring within 29 days.

Ecotoxicity: The isothiazolinones are very toxic to marine organisms, (fish, Daphnia magna water fleas, and algae), and have low potential for accumulation in aquatic species. The proposed metabolites of MI and CMI are considered to have a low aquatic toxicity, based partially on data for the structurally related N-(n-octyl) malonamic acid.

DO NOT discharge into sewer or waterways

Persistence and degradability

Ingredient	Persistence: Water/Soil	Persistence: Air
propylene glycol	LOW	LOW
chlorothalonil	HIGH	HIGH
5-chloro-2-methyl- 4-isothiazolin-3-one	HIGH	HIGH

Bioaccumulative potential

Ingredient	Bioaccumulation
propylene glycol	LOW (BCF = 1)
chlorothalonil	LOW (BCF = 125)
5-chloro-2-methyl- 4-isothiazolin-3-one	LOW (LogKOW = 0.0444)
zinc oxide	LOW (BCF = 217)

Mobility in soil

Ingredient	Mobility
propylene glycol	HIGH (KOC = 1)
chlorothalonil	LOW (KOC = 2392)
5-chloro-2-methyl- 4-isothiazolin-3-one	LOW (KOC = 45.15)

SECTION 13 DISPOSAL CONSIDERATIONS

Waste treatment methods

- ► 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.
 - $\blacksquare \ \ \, \text{Where possible retain label warnings and SDS and observe all notices pertaining to the product.}$

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

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

- ► Reduction
- ► Reuse
- Recycling
- Disposal (if all else fails)

Continued...

Product / Packaging disposal

Chemwatch: 9-339815 Page **10** of **12**

Version No: 1.3

Trouble Shooter Acr.-Neutral Base 45195

Issue Date: 02/07/2017 Print Date: 02/07/2017

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

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

- ▶ 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.

SECTION 14 TRANSPORT INFORMATION

Labels Required



Marine Pollutant



Land transport (DOT)

UN number	3082		
UN proper shipping name	Environmentally hazardous substance, liquid, n.o.s.		
Transport hazard class(es)	Class 9 Subrisk Not Applicable		
Packing group			
Environmental hazard	Not Applicable		
Special precautions for user	Hazard Label 9 Special provisions 8, 146, 173, 335, IB3, T4, TP1, TP29		

Air transport (ICAO-IATA / DGR)

	,		
UN number	3082		
UN proper shipping name	Environmentally hazardous substance, liquid, n.o.s. *		
		_	
Transport hazard class(es)	ICAO/IATA Class	9	
	ICAO / IATA Subrisk	Not Applicable	
	ERG Code	9L	
		·	
Packing group	III		
Environmental hazard	Not Applicable		
	Special provisions		A97 A158 A197
	Cargo Only Packing Instructions		964
	Cargo Only Maximum Qty / Pack		450 L
Special precautions for user	Passenger and Cargo	Packing Instructions	964
	Passenger and Cargo	Maximum Qty / Pack	450 L
	Passenger and Cargo Limited Quantity Packing Instructions		Y964
	Passenger and Cargo	Limited Maximum Qty / Pack	30 kg G
		-	

Sea transport (IMDG-Code / GGVSee)

UN number	3082		
UN proper shipping name	ENVIRONMENTALLY HAZARDOUS SUBSTANCE, LIQUID, N.O.S.		
Transport hazard class(es)	IMDG Class 9 IMDG Subrisk Not Applicable		
Packing group	Ш		
Environmental hazard	Marine Pollutant		

Trouble Shooter Acr.-Neutral Base 45195

Special precautions for user

EMS Number	F-A, S-F
Special provisions	274 335 969
Limited Quantities	5 L

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

Not Applicable

SECTION 15 REGULATORY INFORMATION

US AIHA Workplace Environmental Exposure Levels (WEELs)

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

PROPYLENE GLYCOL(57-55-6) IS FOUND ON THE FOLLOWING REGULATORY LISTS	
US - Pennsylvania - Hazardous Substance List	US ATSDR Minimal Risk Levels for Hazardous Substances (MRLs)
US - Rhode Island Hazardous Substance List	US Spacecraft Maximum Allowable Concentrations (SMACs) for Airborne Contaminants
US - Washington Toxic air pollutants and their ASIL, SQER and de minimis emission values	US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory

CHLOROTHALONIL(1897-45-6) IS FOUND ON THE FOLLOWING REGULATORY LISTS	
International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Monographs	US - New Jersey Right to Know - Special Health Hazard Substance List (SHHSL): Carcinogens
US - California - Proposition 65 - Priority List for the Development of MADLs for Chemicals	US - Pennsylvania - Hazardous Substance List
Causing Reproductive Toxicity	US - Washington Toxic air pollutants and their ASIL, SQER and de minimis emission values
US - California Proposition 65 - Carcinogens	US EPCRA Section 313 Chemical List
US - California Proposition 65 - No Significant Risk Levels (NSRLs) for Carcinogens	US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory
LIS - Massachusetts - Right To Know Listed Chemicals	·

5-CHLORO-2-METHYL-4-ISOTHIAZOLIN-3-ONE(26172-55-4) IS FOUND ON THE FOLLOWING REGULATORY LISTS

US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory	US TSCA Section 5(a)(2) - Significant New Use Rules (SNURs)

ZINC OXIDE(1314-13-2) IS FOUND ON THE FOLLOWING REGULATORY LISTS

, ,	
International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Monographs	US - Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air Contaminants
US - Alaska Limits for Air Contaminants	US - Washington Permissible exposure limits of air contaminants
US - California OEHHA/ARB - Chronic Reference Exposure Levels and Target Organs	US - Wyoming Toxic and Hazardous Substances Table Z1 Limits for Air Contaminants
(CRELs)	US ACGIH Threshold Limit Values (TLV)
US - California Permissible Exposure Limits for Chemical Contaminants	US CWA (Clean Water Act) - Priority Pollutants
US - Hawaii Air Contaminant Limits	US CWA (Clean Water Act) - Toxic Pollutants
US - Idaho - Limits for Air Contaminants	US EPA Carcinogens Listing
US - Massachusetts - Right To Know Listed Chemicals	US EPCRA Section 313 Chemical List
US - Michigan Exposure Limits for Air Contaminants	US National Toxicology Program (NTP) 14th Report Part B. Reasonably Anticipated to be a
US - Minnesota Permissible Exposure Limits (PELs)	Human Carcinogen
US - Oregon Permissible Exposure Limits (Z-1)	US NIOSH Recommended Exposure Limits (RELs)
US - Pennsylvania - Hazardous Substance List	US OSHA Permissible Exposure Levels (PELs) - Table Z1
US - Rhode Island Hazardous Substance List	US OSHA Permissible Exposure Levels (PELs) - Table Z3
US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants	US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory
US - Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants	

Federal Regulations

Superfund Amendments and Reauthorization Act of 1986 (SARA)

SECTION 311/312 HAZARD CATEGORIES

Immediate (acute) health hazard	Yes
Delayed (chronic) health hazard	Yes
Fire hazard	No
Pressure hazard	No
Reactivity hazard	No

US. EPA CERCLA HAZARDOUS SUBSTANCES AND REPORTABLE QUANTITIES (40 CFR 302.4)

None Reported

State Regulations

US. CALIFORNIA PROPOSITION 65

WARNING: This product contains a chemical known to the State of California to cause cancer and birth defects or other reproductive harm

US - CALIFORNIA PREPOSITION 65 - CARCINOGENS & REPRODUCTIVE TOXICITY (CRT): LISTED SUBSTANCE

Chlorothalonil Listed

National Inventory	Status
Australia - AICS	Y
Canada - DSL	Υ

Chemwatch: 9-339815 Page 12 of 12 Issue Date: 02/07/2017 Version No: 1.3 Print Date: 02/07/2017

Trouble Shooter Acr.-Neutral Base 45195

Canada - NDSL	N (chlorothalonil; 5-chloro-2-methyl-4-isothiazolin-3-one; propylene glycol)
China - IECSC	Y
Europe - EINEC / ELINCS / NLP	Y
Japan - ENCS	Y
Korea - KECI	Y
New Zealand - NZIoC	Y
Philippines - PICCS	Υ
USA - TSCA	Υ
Legend:	Y = All ingredients are on the inventory N = Not determined or one or more ingredients are not on the inventory and are not exempt from listing(see specific ingredients in brackets)

SECTION 16 OTHER INFORMATION

CONTACT POINT

PLEASE NOTE THAT TITANIUM DIOXIDE IS NOT PRESENT IN CLEAR OR NEUTRAL BASES

Other information

Ingredients with multiple cas numbers

Name	CAS No
zinc oxide	1314-13-2, 175449-32-8

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

This document is copyright.

Apart from any fair dealing for the purposes of private study, research, review or criticism, as permitted under the Copyright Act, no part may be reproduced by any process without written permission from CHEMWATCH.

TEL (+61 3) 9572 4700.