EME215/40A POLYOL



Version 1 Erapol Co. Material Safety Data Sheet (Conforms to Reg. (EC) No 1907/2006, Reg. (EC) No 1272/2008 and their amendments) Erapol Co.9-31399 Print Date: 20-Jan-2012 Revision Date: 20-Jan-2012 Issue Date: 20-Jan-2012

SAFETY DATA SHEET

	of the substance (minture and of the company) (undertaking			
	of the substance / mixture and of the company / undertaking			
.1. Product Identifier				
Product name:	EME215/40A POLYOL			
Chemical product name:	No data available			
Synonyms:	No data available			
roper shipping name:	None			
chemical formula:	No data available			
Other means of identification:	No data available			
ndex number:	No data available			
D number:	No data available			
AS number:	No data available			
REACH registration number:	No data available			
C number:	Not Available			
.2. Relevant identified us	ses of the substance or mixture and uses advised against			
Relevant identified uses:	Used according to manufacturer's directions. Polyol component			
Jses advised against:	No data available			
.3. Details of the supplier	r of the safety data sheet			
Registered company name:	Era Polymers Pty Ltd			
Address:	25-27 Green Street, Banksmeadow, NSW 2019, Australia			
elephone:	+61 2 9666 3788			
ax:	+61 2 9666 4805			
mail:	erapol@erapol.com.au			
Vebsite:				
.4. Emergency telephone	e number			
Association / Organisation:				
Other emergency telephone	1800 039 008 (AUS)			
umbers: Other emergency telephone Jumbers:	+80024362255 (INTL)			
SECTION 2: Hazards ident	tification			
2.1. Classification of the s				
DSD classification:				
SD classification	In case of mixtures, classification has been prepared by following DPD (Directive 1999/45/EC) or CLP (Regulation (EC) No 1272/2008) regulations No data available			
additional): DPD classification:				
FD classification.	R20/21/22 • Harmful by inhalation, in contact with skin and if swallowed.			
	R33 • Danger of cumulative effects.			
	 May cause long-term adverse effects in the aquatic environment. 			
CLP classification:	Chronic Aquatic Hazard Category 4			
	Acute Toxicity (Oral) Category 4			
	Acute Toxicity (Dermal) Category 4			
	Acute Toxicity (Inhalation) Category 4			
	STOT - RE Category 2			
CLP classification additional):	No data available			
2.2. Label elements				
CLP label elements				



Signal word: Hazard statement(s): WARNING H413 H302

May cause long lasting harmful effects to aquatic life. Harmful if swallowed

H332 Harmful if inhaled H373 May cause damage to organs through prolonged or repeated exposure. Determined by Cherwwatch using CLP criteria Additional Statement(s): No data available Supplementary statement(s): No data available Precautionary statement(s): Prevention Code Phrase P260 Do not breathe dust/fume/gas/mist/vapours/spr P261 Avoid breathing dust/fume/gas/mist/vapours/spr P264 Wash thoroughly after handling. P270 Do not eat, drink or smoke when using this prod P271 Use only outdoors or in a well-ventilated area. P273 Avoid release to the environment. P280 Wear protective gloves/protective clothing/eye protective gloves/protective gloves/protective clothing/eye protective gloves/protective gloves/protective gloves/protective gloves/protective gloves/protective gloves/protective gloves/protective gloves/protect	Harmful in contact with skin		
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P330 Rinse mouth. P363 Wash contaminated clothing before reuse. Disposal Code Phrase P501 Dispose of contents/container to	Do not breathe dust/fume/gas/mist/vapours/spray. Avoid breathing dust/fume/gas/mist/vapours/spray. Wash thoroughly after handling. Do not eat, drink or smoke when using this product. Use only outdoors or in a well-ventilated area. Avoid release to the environment. Wear protective gloves/protective clothing/eye protection/face protection. Phrase IF SWALLOWED: Call a POISON CENTER or doctor/physician if you feel unwell. IF ON SKIN: Wash with plenty of soap and water. IF INHALED: Remove to fresh air and keep at rest in a position comfortable for breathing. Call a POISON CENTER or doctor/physician if you feel unwell. Get medical advice/attention if you feel unwell. Rinse mouth. Wash contaminated clothing before reuse. Phrase		

DSD / DPD label elements



Indication(s) of danger:	CONSIDERED A DANGE	EROUS MIXTURE ACCORDING TO DIRECTIVE 1999/45/EC AND ITS AMENDMENTS.
Safety advice:	S23	 Do not breathe gas/fumes/vapour/spray.
	S24	Avoid contact with skin.
	S36	Wear suitable protective clothing.
	S37	Wear suitable gloves.
	S51	Use only in well ventilated areas.
	S09	Keep container in a well ventilated place.
	S401	 To clean the floor and all objects contaminated by this material, use water and detergent.
	S07	Keep container tightly closed.
	S13	 Keep away from food, drink and animal feeding stuffs.
	S46	 If swallowed, IMMEDIATELY contact Doctor or Poisons Information Centre. (show this container or label).

2.3. Other hazards

	No data available
PBT/vPvB criteria	No data available

SECTION 3: Composition / information on ingredients

Section 3. composition / information on ingredients						
3.1. Substances						
See 'Composition on ingre	dients' in section 3.	2				
3.2. Mixtures						
1. CAS No 2. EC No 3. Index No 4. REACH No	%[weight]	Name		cation according to Directive /EC [DPD]	Classification according to (EC) No 1272/2008 [CLP]	
					Acute Tox. 2 *	
	<1				Acute Tox. 1	
1. 27236-65-3			T+		Acute Tox. 2 *	
2. 248-355-2 3. 080-004-00-7 4. No data available		bis(phenylmercury)	N	R50/53	STOT RE 2 *	
		dodecenylsuccinate	_	R26/27/28	Aquatic Acute 1	
				R33	Aquatic Chronic 1	
					CLP classification according to Annex VI of CLP (Regulation (EC) No 1272/2008)	
1. 13674-84-5* 2. 237-158-7 3. 237-158-7 4. No data available	>25	tris(2-chloroisopropyl)phosphate	Xn	R22	Acute Toxicity Category 4	

4.1. Description of f	
General:	No data available
Ingestion:	 IF SWALLOWED, REFER FOR MEDICAL ATTENTION, WHERE POSSIBLE, WITHOUT DELAY. For advice, contact a Poisons Information Centre or a doctor. Urgent hospital treatment is likely to be needed. In the mean time, qualified first-aid personnel should treat the patient following observation and employing supportive measures as indicated be the patient's condition. If the services of a medical officer or medical doctor are readily available, the patient should be placed in his/her care and a copy of the MSD should be provided. Further action will be the responsibility of the medical specialist. If medical attention is not available on the worksite or surroundings send the patient to a hospital together with a copy of the MSDS.
	Where medical attention is not immediately available or where the patient is more than 15 minutes from a hospital or unless instructe otherwise:
	 INDUCE vomiting with fingers down the back of the throat, ONLY IF CONSCIOUS. Lean patient forward or place on left side (head-down position if possible) to maintain open airway and prevent aspiration.
	NOTE: Wear a protective glove when inducing vomiting by mechanical means.
Eye Contact:	If this product comes in contact with the eyes:
	 Wash out immediately with fresh running water. Ensure complete irrigation of the eye by keeping eyelids apart and away from eye and moving the eyelids by occasionally lifting the upper an lower lids. Seek medical attention without delay; if pain persists or recurs seek medical attention. Removal of contact lenses after an eye injury should only be undertaken by skilled personnel.
Skin Contact:	If skin contact occurs:
	 Immediately remove all contaminated clothing, including footwear. Flush skin and hair with running water (and soap if available). Seek medical attention in event of irritation.
Inhalation:	 If fumes or combustion products are inhaled remove from contaminated area. Lay patient down. Keep warm and rested. Prostheses such as false teeth, which may block airway, should be removed, where possible, prior to initiating first aid procedures. Apply artificial respiration if not breathing, preferably with a demand valve resuscitator, bag-valve mask device, or pocket mask as trained. Perfort CPR if necessary. Transport to hospital, or doctor.
4.2. Most important	symptoms and effects, both acute and delayed
Inhaled:	Inhalation of vapours or aerosols (mists, fumes), generated by the material during the course of normal handling, may be harmful.
	• 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.
Ingestion:	Accidental ingestion of the material may be harmful; animal experiments indicate that ingestion of less than 150 gram may be fatal or may product serious damage to the health of the individual.
Skin Contact:	 Skin contact with the material may be harmful; systemic effects may result following absorption. The material is not thought to be a skin irritant (i.e. is unlikely to produce irritant dermatitis as described in EC Directives using animal models). Temporary discomfort, however, may result from prolonged dermal exposures. Good hygiene practice requires that exposure be kept to a minimum and that suitable gloves be used in an occupational setting.
	Open cuts, abraded or irritated skin should not be exposed to this material.
	• 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.
Eye:	Although the liquid is not thought to be an irritant (as classified by EC Directives), direct contact with the eye may produce transient discomfor characterised by tearing or conjunctival redness (as with windburn).
Chronic:	Repeated or long-term occupational exposure is likely to produce cumulative health effects involving organs or biochemical systems.

For acute and short term repeated exposures to anyl and alkylmethoxy compounds of mercury: Absorption proceeds more rapidly than its inorganic counterpart but once inside the body biotransformation releases inorganic mercury. [Ellenhorn and Barceloux: Medical Toxicology]

- Moderate adsorption of inorganic mercury compounds through the gastro-intestinal tract (7-15%) is the principal cause of poisoning. These compounds are highly concentrated (as the mercuric (Hg (2+) form) in the kidney; acute ingestion may lead to oliguric renal failure. Severe mucosal necrosis may also result from ingestion.
 Chronic effects range from proteinuria to nephrotic syndrome. Chronic presentation also involves dermatitis, gingivitis, stomatitis, tremor and neuropsychiatric symptoms of erethism.
 Absorbed inorganic mercury does not significantly cross the blood-brain barrier.
 Emesis and lavage should be initiated following acute ingestion.
 Activated charcoal interrupts absorption; cathartics should be administered when charcoal is given.
 The use of British Anti-Lewisite is indicated in severe inorganic poisoning. Newer derivatives of BAL (e.g. dimercaptosuccinic acid, [DMSA] and 2,3-dimercapto-1-propanesulfate [DMPS]) may prove more effective. [Ellenhorn and Barceloux: Medical Toxicology]

BIOLOGICAL EXPOSURE INDEX - BEI

These represent the determinants observed in specimens from a healthy worker exposed at the Exposure Standard (ES or TLV).					
Determinant	Index	Sampling Time	Comments		
1. Total inorganic mercury in urine	35 ug/gm creatinine	Preshift	В		
2. Total inorganic mercury in blood	15 ug/L	End of shift at end of workweek	В		

B: Background levels occur in specimens collected from subjects NOT exposed.

SECTION 5: Firefighting measures	
5.1. Extinguishing media	

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

	Use extinguishing media suitable for surrounding area.
5.2. Special hazards ari	ising from the substrate or mixture
Fire Incompatibility:	None known.
5.3. Advice for firefight	ers
Fire Fighting:	 Alert Fire Brigade and tell them location and nature of hazard. Wear breathing apparatus plus protective gloves for fire only. 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 from path of fire. 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
6.1. Personal precautio	ns, protective equipment and emergency procedures
Personal Protective	Breathing apparatus.Chemical splash suit.
Equipment: Minor Spills:	
	 Clean up all spills immediately. Avoid breathing vapours and contact with skin and eyes. Control personal contact 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:	 Moderate hazard. Clear area of personnel and move upwind. Alert Fire Brigade and tell them location and nature of hazard. Wear breathing apparatus plus protective gloves. Prevent, by any means available, spillage from entering drains or water course. Stop leak if safe to do so. Contain spill with sand, earth or vermiculite. Collect recoverable product into labelled containers for recycling. Neutralise/decontaminate residue (see Section 13 for specific agent). Collect solid residues and seal in labelled drums for disposal. Wash area and prevent runoff into drains. After clean up operations, decontaminate and launder all protective clothing and equipment before storing and re-using. If contamination of drains or waterways occurs, advise emergency services.
6.2. Environmental pred See section 12	
	rial for containment and cleaning up
6.4. Reference to other	
	nent advice is contained in Section 8 of the MSDS
SECTION 7: Handling a	ind storage
7.1. Precautions for saf	fe handling
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. DO NOT allow material to contact humans, exposed food or food utensils. Avoid contact with incompatible materials. When handling, DO NOT et dirk or smoke. Keep containers securely sealed when not in use. Avoid physical damage to containers. 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 storing and handling recommendations. Atmosphere should be regularly checked against established exposure standards to ensure safe working conditions are maintained.
Fire and explosion protection	

Fire and explosion protection See section 5

Other information

SAFE STORAGE WITH OTHER CLASSIFIED CHEMICALS



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+: May be stored together.

O: May be stored together with specific preventions.

X: Must not be stored together.

7.2. Conditions for safe storage, including any incompatibilities

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	orago, moraling any moonpationatoo
Suitable container:	 Polyethylene or polypropylene container. Packing as recommended by manufacturer. Check all containers are clearly labelled and free from leaks.
Storage incompatibility:	 WARNING: Avoid or control reaction with peroxides. All <i>transition metal</i> peroxides should be considered as potentially explosive. For example transition metal complexes of alkyl hydroperoxides may decompose explosively. The pi-complexes formed between chromium(0), vanadium(0) and other transition metals (haloarene-metal complexes) and mono-or polyfluorobenzene show extreme sensitivity to heat and are explosive. Avoid reaction with borohydrides or cyanoborohydrides

Package Material Incompatibilities:

No data available

7.3. Specific end use(s)

See section 1.2

SECTION 8: Exposure controls / personal protection

8.1. Control parameters					
Derived No Effect Level (DNEL)					
Exposure Pattern	Workers	General Population	Exposure Pattern	Workers	General Population
Long term - dermal, systemic effects	No data available	No data available	Short term - dermal, systemic effects	No data available	No data available
Long term - inhalation, systemic effects	No data available	No data available	Short term - inhalation, systemic effects	No data available	No data available
Long term - oral, systemic effects	No data available	No data available	Short term - oral, systemic effects	No data available	No data available
Long term - dermal, local effects	No data available	No data available	Short term - dermal, local effects	No data available	No data available
Long term - inhalation, local effects	No data available	No data available	Short term - inhalation, local effects	No data available	No data available
Occupational Exposure Limi	its (OFI.)				

The following materials had no OELs on our records		
 bis(phenylmercury) dodecenylsuccinate: 		CAS:27236-65-3
tris(2-chloroisopropyl)phosphate:		CAS:13674-84-5
EMERGENCY EXPOSURE LIMITS		
Material	Revised IDLH Value (mg/m ³)	Revised IDLH Value (ppm)
bis(phenylmercury) dodecenylsuccinate 35331	10	

BIS(PHENYLMERCURY) DODECENYLSUCCINATE: EME215/40A POLYOL:

BIS(PHENYLMERCURY) DODECENYLSUCCINATE:

It is the goal of the AGCIH (and other Agencies) to recommend TLVs (or their equivalent) for all substances for which there is evidence of health effects at airborne concentrations encountered in the workplace.

At this time no TLV has been established, even though this material may produce adverse health effects (as evidenced in animal experiments or clinical experience). Airborne concentrations must be maintained as low as is practically possible and occupational exposure must be kept to a minimum. **NOTE:** The ACGIH occupational exposure standard for Particles Not Otherwise Specified (P.N.O.S) does NOT apply.

Exposure limits with "skin" notation indicate that vapour and liquid may be absorbed through intact skin. Absorption by skin may readily exceed vapour inhalation exposure. Symptoms for skin absorption are the same as for inhalation. Contact with eyes and mucous membranes may also contribute to overall exposure and may also invalidate the exposure standard. TRIS(2-CHLOROISOPROPYL)PHOSPHATE:

Not available

8.2. Exposure controls

8.2.1. Appropriate engineering 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 the design of a ventilation system must match the particular process and chemical or contaminant in the design of a ventilation system must match the particular process and chemical or contaminant in the design of a ventilation system must match the particular process and chemical or contaminant in the design of a ventilation system must match the particular process and chemical or contaminant in the design of a ventilation system must match the particular process and chemical or contaminant in the design of a ventilation system must match the particular process and chemical or contaminant in the design of a ventilation system must match the particular process and chemical or contaminant in the design of a ventilation system must match the particular process and chemical or contaminant in the design of a ventilation system must match the particular process and chemical or contaminant in the design of a ventilation system must match the particular process and chemical or contaminant in the design of a ventilation system must match the particular process and chemical or contaminant in the design of a ventilation system must match the particular process and chemical or contaminant in the design of a ventilation system must match the particular process and chemical or contaminant in the design of a ventilation system must be a ventilation system must 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.

Type of Contaminant:

solvent, vapours, degreasing etc., evaporating from tank (in still air). 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)

Air Speed: 0.25-0.5 m/s (50-100 f/min.) 0.5-1 m/s (100-200 f/min.)

1-2.5 m/s (200-500 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)

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

Within each range the appropriate value depends on: Lower end of the range

- 1: Room air currents minimal or favourable to capture
- 2: Contaminants of low toxicity or of nuisance value only.

3: Intermittent, low production.

4: Large hood or large air mass in motion

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

- Upper end of the range
- Disturbing room air currents
 Contaminants of high toxicity
- 3: High production, heavy use
- 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.

8.2.2. Personal protection

Eye and face protection:



· Safety glasses with side shields

	wearing of lens or restrictions on use, sh adsorption for the class of chemicals in u removal and suitable equipment should b contact lens as soon as practicable. Lens	e readily available. In the event of chemical exposishould be removed at the first signs of eye redness	
Skin protection:	See Hand protection: below		
Hand protection:	 frequency and duration of contact, chemical resistance of glove material, glove thickness and dexterity Select gloves tested to a relevant standard (e.g. E When prolonged or frequently repeated cominutes according to EN 374, AS/NZS 216 When only brief contact is expected, a glo 374, AS/NZS 2161.10.1 or national equival Contaminated gloves should be replaced. Gloves must only be worn on clean hands. After u is recommended. WARNING: Do NOT use latex or PVC gloves In 1997, a researcher (Dr. Karen E. Wett almost a year before. Heavy metals and organic metal compoun she lapsed into a vegetative state, Dr. Wet 	t on usage. Important factors in the selection of glow urope EN 374, US F739, AS/NZS 2161.1 or national ontact may occur, a glove with a protection class of 1.10.1 or national equivalent) is recommended. we with a protection class of 3 or higher (breakthro lent) is recommended. sing gloves, hands should be washed and dried tho erhahn) died from organic mercury poisoning, res ds, in particular, have posed special hazards in wo terhahn asked that her case be made known to othe exposure to dimethylmercury produced the following Thickness in mm* 0.2 0.8 0.33 0.28 0.13	al equivalent). of 5 or higher (breakthrough time greater than 240 ugh time greater than 60 minutes according to EN proughly. Application of a non-perfumed moisturiser sulting from a single exposure to dimethylmercury rker protection. At the time of diagnosis and before ars.
	*Michael B Blayney: Applied Occupational and Environmental Hygiene: * Originally quoted as mil (one mil = 0.001 inches)	0.7 : 16, pp 233-236, 2001	>240 mins.
Body protection:	See Other protection: below		
Other protection:	 Overalls. P.V.C. apron. Barrier cream. Skin cleansing cream. Eye wash unit. 		
Respiratory protection:			
Thermal hazards:	No data available		
Recommended material(s):	Glove selection is based on a modified presentation "Forsberg Clothing Performance Index" . The effect(s) of the following substance(s) are take	n of the:	

The effect(s) of the following substance(s) are taken into account in the *computer-generated* selection: Material CPI * CPI - Chemwatch Performance Index

* CPI - Chemwatch Performance index A: Best Selection B: Satisfactory; may degrade after 4 hours continuous immersion C: Poor to Dangerous Choice for other than short term immersion NOTE: As a series of factors will influence the actual performance of the glove, a final selection must be based on detailed observation. -* Where the glove is to be used on a short term, casual or infrequent basis, factors such as "feel" or convenience (e.g. disposability), may dictate a choice of gloves which might otherwise be unsuitable following long-term or frequent use. A qualified practitioner should be consulted.

8.2.3. Environmental exposure controls

See section 12

SECTION 9: Physical and chemical properties

SECTION 9: Physical and chemical properties	
9.1. Information on basic physical and chemica	l properties
Appearance	No data available
Odour	No data available
Odour threshold	No data available
Taste	No data available
pH (1% solution)	No data available
pH (as supplied)	No data available
Melting point / freezing point (°C)	No data available
Initial boiling point and boiling range (°C)	No data available
Flash point (°C)	No data available
Evaporation rate	No data available
Flammability	No data available
Vapour pressure (kPa)	No data available
Vapour density (Air = 1)	No data available
Relative density (Water = 1)	1.20
Solubility in Water (g/L)	No data available
Partition coefficient: n-octanol / water	No data available
Auto-ignition temperature (°C)	No data available
Critical temperature (°C)	No data available
Viscosity (cSt)	No data available
Explosive properties	No data available
Oxidising properties	No data available
Physical state	Liquid
Upper Explosive Limit (%)	No data available
Lower Explosive Limit (%)	No data available
Surface Tension	No data available
Volatile Component (%vol)	No data available
Gas group	No data available
Molecular weight (g/mol)	No data available
Evaporation Rate (BuAc = 1 EtAc = 1 Ether = 1)	No data available
IUCLID Remarks	No data available

9.2. Other information

No data available

SECTION 10: Stability and reactivity

02011	on To. Stability and	
10.1.	Reactivity	See section 7.2
10.2.	Chemical stability	1
		Presence of incompatible materials.
		Product is considered stable.
		Hazardous polymerisation will not occur.
10.3.	Possibility of	
10.5.	hazardous reactions	See section 7.2
10.4.	Conditions to avoid	See section 7.2
10.5.	Incompatible materials	See section 7.2
10.6.	Hazardous decomposition products	See section 5.3
	<mark>ON 11: Toxicologic</mark> a	

11.1. Information on toxicological effects					
Mutagenicity:	No data available				
Reproductive Toxicity:	No data available				
Carcinogenicity:	No data available				
STOT - single exposure:	No data available				

BIS(PHENYLMERCURY) DODECENYLSUCCINATE:EME215/40A POLYOL: No significant acute toxicological data identified in literature search.EME215/40A POLYOL:-OTHERBIS (PHENYLMERCURY) DODECENYLSUCCINATE:TRIS(2-CHLOROISOPROPYL)PHOSPHATE: None assigned. Refer to individual constituents.

SECTION 12: Ecologi	SECTION 12: Ecological information				
12.1. Toxicity					
Fish:	No data available				
Daphnia Magna:	No data available				
Algae:	No data available				
	No data available				

Toxic to aquatic microorganisms:

TRIS(2-CHLOROISOPROPYL)PHOSPHATE: BIS(PHENYLMERCURY) DODECENYLSUCCINATE:

BIS(PHENYLMERCURY) DODECENYLSUCCINATE:

Yes

Marine Pollutant

Very 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.

equipinient was revenues. Wastes resulting from use of the product must be disposed of on site or at approved waste sites. Mercury may occur in the environment as free mercury, Hg(0), mercury ions in salts and complexes, Hg+ and (Hg2)2+ and as organic mercury compounds. Each species has its own set of physical, chemical and toxicologic properties. In natural systems a dynamic equilibrium between soil and water mercury occurs determined largely by the physicochemical and biological conditions which pertain. Mercury ion is

transported to aquatic ecosystems via surface run-off and from the atmosphere. It is complexed or tightly bound to both inorganic and organic particles, particularly sediments with high sulfur content. Organic acids such as fulvic and humic acids are often associated with mercury not bound to particles. Methyl mercury is produced by sediment micro-organisms, nonbiologically in sediments and by certain species of fish. The methylation of mercury by micro-organisms is the detoxification response that allows the organism to dispose of the heavy metal ions as small organometallic complexes. Methylation occurs only within a narrow pH range in which the micro-organism inght exist and the rate of synthesis depends on the redox potential, composition of the microbial population, availability of Hg2+ and temperature. Vitamin B12 derivatives are thought to be the methylating agents, because they are the only methyl carbanion- or methyl radical-donating coenzymes known. In addition it has been demonstrated that the livers of yellow-fin tuna and albacore produce methyl mercury results in its desorption at relatively high rates thus little methyl mercury is found in sediments. Demethylation by sediment micro-organisms also occurs at a rapid rate compared with methylation. The best conversion rate for inorganic mercury to methyl mercury under ideal conditions is less than 1.5% per month. Methyl mercury released into surface waters may also undergo photodecomposition into mercury.

Methyl mercury can be bioaccumulated by planktonic algae and fish. In fish, the rate of absorption of methyl mercury is faster than that of inorganic mercury and the clearance rate is slower resulting in high concentrations of methyl mercury in muscle tissue. The ratio of organic mercury to total mercury is generally high in fish compared with other aquatic organisms. Selenium which is also present in seawater and other seafoods readily complexes with methyl mercury and is thought to have a protective effect against the toxic action of methyl mercury. The danger of methyl mercury poisoning has been illustrated in Minimata, Japan in the late 1950s following industrial release of mercury into the bay which subsequently resulted in at least 1200 cases of poisoning, some fatal. TRIS(2-CHLOROISOPROPYL)PHOSPHATE:

IngredientPersistence: Water/SoilPersistence: AirEME215/40A POLYOLNo Data AvailableNo Data Availablebis(phenylmercury) dodecenylsuccinateNo Data AvailableNo Data Availabletris(2-chloroisopropyl)phosphateHIGHNo Data AvailableBioaccumulative potentialIngredientBioaccumulationtris(2-chloroisopropyl)phosphateLOWI.4. Mobility in soilIngredientMobilityIngredientIngredientIngredientI.2.S. Results of PBT and vPvB assessmentPBT	12.2. Persistence and deg	radability			
bis(phenylmercury) dodecenylsuccinate No Data Available No Data Available tris(2-chloroisopropyl)phosphate HIGH No Data Available 12.3. Bioaccumulative potential Ingredient Bioaccumulation tris(2-chloroisopropyl)phosphate LOW 12.4. Mobility in soil 12.4. Mobility in soil Mobility Ingredient Ingredient Mobility Mobility tris(2-chloroisopropyl)phosphate MED (ESTIMATED)	Ingredient Persistence: Water/Soil		Persistence: Water/Soil		Persistence: Air
tris(2-chloroisopropyl)phosphate HIGH No Data Available 12.3. Bioaccumulative potential Ingredient Bioaccumulation Ingredient LOW Ingredient 12.4. Mobility in soil Mobility Ingredient Ingredient Mobility Mobility 12.5. Results of PBT and vPvB assessment Ingredient Ingredient	EME215/40A POLYOL		No Data Available		No Data Available
12.3. Bioaccumulative potential Ingredient Bioaccumulation tris(2-chloroisopropyl)phosphate LOW 12.4. Mobility in soil Mobility Ingredient Mobility tris(2-chloroisopropyl)phosphate Mobility 12.5. Results of PBT and vPvB assessment	bis(phenylmercury) dodece	enylsuccinate	No Data Available		No Data Available
Ingredient Bioaccumulation tris(2-chloroisopropyl)phosphate LOW 12.4. Mobility in soil Ingredient Ingredient Mobility tris(2-chloroisopropyl)phosphate MED (ESTIMATED) 12.5. Results of PBT and vPvB assessment	tris(2-chloroisopropyl)phosp	ohate	HIGH		No Data Available
tris(2-chloroisopropyl)phosphate LOW 12.4. Mobility in soil Ingredient Ingredient Mobility tris(2-chloroisopropyl)phosphate MED (ESTIMATED) 12.5. Results of PBT and vPvB assessment	12.3. Bioaccumulative pot	tential			
12.4. Mobility in soil Ingredient Mobility tris(2-chloroisopropyl)phosphate MED (ESTIMATED) 12.5. Results of PBT and vPvB assessment	Ingredient		Bioaccumulation		
Ingredient Mobility tris(2-chloroisopropyl)phosphate MED (ESTIMATED) 12.5. Results of PBT and vPvB assessment Estimate of the second se	tris(2-chloroisopropyl)phosphate		LOW		
tris(2-chloroisopropyl)phosphate MED (ESTIMATED) 12.5. Results of PBT and vPvB assessment	12.4. Mobility in soil				
12.5. Results of PBT and vPvB assessment	Ingredient		M	obility	
	tris(2-chloroisopropyl)phosphate		MED (ESTIMATED)		
P B T	12.5. Results of PBT and vPvB assessment				
		Р	В	T	
Relevant available data No data available No data available No data available	Relevant available data	No data available	No data available	No data available	
PBT and vPvB Criteria fulfilled? No data available No data available No data available		No data available	No data available	No data available	

12.6. Other adverse effects

No data available

SECTION 13: Disposal considerations

13.1. Waste treatment methods

Product / Packaging

disposal:

Containers may still present a chemical hazard/ danger when empty.
Return to supplier for reuse/ recycling if possible.

Otherwise:

- If container can not be cleaned sufficiently well to ensure that residuals do not remain or if the container cannot be used to store the same product, then puncture containers, to prevent re-use, and bury at an authorised landfill.
 Where possible retain label warnings and MSDS 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)

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 pro-
- It may be necessary to collect all wash water for treatment before disposal. In all cases disposal to sewer may be subject to local laws and regulations and these should be considered first.
- · Where in doubt contact the responsible authority.
- Recycle wherever possible
- Consult manufacturer for recycling options or consult local or regional waste management authority for disposal if no suitable treatment or disposal facility can be identified. Dispose of by: burial in a land-fill specifically licenced to accept chemical and / or pharmaceutical wastes or incineration in a licenced apparatus
- (after admixture with suitable combustible material).

 Decontaminate empty container 	. Observe all label safeguards until containers are cleaned and destroy	ea.

Waste treatment options:	
Sewage disposal options:	No relevant data

Other disposal recommendations:

SECTION 14: Transport in	formation					
Labels Required:	No data available					
Land transport (ADR / RID / GO	GVSE)					
No data available 14.1. UN number	No data available			14.4. Packing group	No data available	
14.2. UN proper shipping name				14.5. Environmental hazard	No relevant data	
14.3. Transport hazard class(es)				14.6. Special precautions for user	Hazard identification (Kemler)	No data available
	No data available				Classification Code	No data available
	No data available				Hazard Label	No data available
					Special provisions	No data available
					Add limited quantity	No data available
Air transport (ICAO-IATA / DGI	R)					
No data available 14.1. UN number	No data available			14.4. Packing group	No data available	
14.2. UN proper shipping name				14.5. Environmental hazard	No relevant data	
14.3. Transport hazard class(es)				14.6. Special precautions for user	Special provisions	No data available
					Cargo Only Packing Instructions	No data available
					Cargo Only Maximum Qty / Pack	No data available
	ICAO/IATA Class: ICAO/IATA Subris		data available data available		Passenger and Cargo Packing Instructions	No data available
	ERG Code	No	data available		Passenger and Cargo Maximum Qty / Pack	No data available
					Passenger and Cargo Limited Quantity Packing Instructions	No data available
					Passenger and Cargo Maximum Qty / Pack	No data available
Sea transport (IMDG-Code / G0	GVSee)					
No data available 14.1. UN number				14.4. Packing group		
14.1. UN proper shipping	No data available			14.4. Facking group	No data available	
name	No data available			hazard	No relevant data	
14.3. Transport hazard				14.6. Special	EMS Number	No data available
class(es)	No data available	IMDG Subrisk	No data available	precautions for user	Special provisions	No data available
					Limited Quantities	No data available
Inland waterways transport (A	DNR / River <u>Rhine)</u>					
No data available						
14.1. UN number	No data available			14.4. Packing group	No data available	
14.2. UN proper shipping name	No data available			14.5. Environmental hazard	No relevant data	
14.3. Transport hazard				14.6. Special	Classification code	No data available
class(es)	No data available	ADNR	No data available	precautions for user	Limited quantity	No data available
	No data available	Label	No data available		Equipment required	No data available
					Fire cones number	No data available
14.7. Transport in bulk accordi		01 70 / 70				
14.7. Transport in bulk accord	Ing to Annex II of MARI	OL 13/18	and the IBC code			

No data available

SECTION 15: Regulatory information

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

Regulations for ingredients

bis(phenylmercury) dodecenylsuccinate (CAS: 27236-65-3) is found on the following regulatory lists;

"European Customs Inventory of Chemical Substances (English)", "European Union - European Inventory of Existing Commercial Chemical Substances (EINECS) (English)" tris(2-chloroisopropyl)phosphate (CAS: 13674-84-5) is found on the following regulatory lists;

"European Chemicals Agency (ECHA) List of substances identified for registration in 2010", "European Customs Inventory of Chemical Substances (English)", "European Union -European Inventory of Existing Commercial Chemical Substances (EINECS) (English)", "International Chemical Secretariat (ChemSec) REACH SIN* List (*Substitute It Now!) 1.1"

No data for EME215/40A POLYOL (CW: 9-31399)

This safety data sheet is in compliance with the following EU legislation and its adaptations – as far as applicable - : 67/548/EEC, 1999/45/EC, 98/24/EC, 92/85/EEC, 94/33/EC, 91/689/EEC, 1999/13/EC, Regulation (EU) No 453/2010, Regulation (EC) No 1907/2006, Regulation (EC) No 1272/2008, and their amendments as well as the following British legislation:

- The Control of Substances Hazardous to Hea	Ith Regulations (COSHH) 2002				
- COSHH Essentials					
- The Management of Health and Safety at Wo	rk Regulations 1999				
15.2. Chemical safety assessment					
ANNEX 1					
Ingredient		Annex 1 67/548/EEC			
bis(phenylmercury) dodecenylsuccinate		080-004-00-7			
Annex VI					
Chronic Aquatic Hazard Category 4					
Acute Toxicity (Oral) Category 4					
Acute Toxicity (Dermal) Category 4					
Acute Toxicity (Inhalation) Category 4					
STOT - RE Category 2					
RISK					
Risk Codes		Risk Phrases			
R20/21/22		• •	ntact with skin and if swallowed.		
R33	Danger of cumulative effects.				
R53		May cause long-term adver	May cause long-term adverse effects in the aquatic environment.		
SECTION 16: Other information					
ANNEX 2: Indications of Danger					
N Dangerous for the environ	ment				
T+ Very toxic					
Xn Harmful					
Substance	CAS	Suggested codes			
tris(2- chloroisopropyl)phosphate		13674- 84- 5	Mut3; R68 Rep3;		
			R63 Xn; R22 Xi;		
			R38		
Denmark Advisory list for selfclassification of da	ngerous substances				
OTHER					

Classification of the preparation and its individual components has drawn on official and authoritative sources using available literature references.

The (M)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.

For detailed advice on Personal Protective Equipment, refer to the following EU CEN Standards: EN 16 Personal eye-protection EN 340 Protective clothing EN 374 Protective gloves against chemicals and micro-organisms EN 13832 Footwear protecting against chemicals EN 133 Respiratory protective devices

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Not applicable

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