

MG Chemicals UK Limited

Version No: 1.1

Safety Data Sheet (Conforms to Regulations (EC) No 2015/830)

Chemwatch Hazard Alert Code: 2 Issue Date: 02/04/2016 Print Date: 02/04/2016 Initial Date: 02/04/2016 L.REACH.GBR.EN

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

1.1.Product Identifier

Product name	32HD Black 1:1 Epoxy Potting and Encapsulating Compound (Part A)		
Synonyms	SDS Code: 832HD-Part A; 832HD-25ML, 832HD-50ML, 832HD-400ML, 832HD-1.7L, 832HD-7.4L, 832HD-40L		
Proper shipping name	ENVIRONMENTALLY HAZARDOUS SUBSTANCE, LIQUID, N.O.S. (contains bisphenol a diglycidyl ether resin, solid)		
Other means of identification Not Available			

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

Relevant identified uses	poxy resin for use with hardeners	
Uses advised against	Not Applicable	

1.3. Details of the supplier of the safety data sheet

Registered company name	MG Chemicals UK Limited	MG Chemicals (Head office)
Address	Address Heame House, 23 Bilston Street, Sedgely Dudley DY3 1JA United Kingdom 9347 - 193 Street Surrey V4N 4E7 British Colu	
Telephone	Telephone +(44) 1663 362888 +(1) 800-201-8822	
Fax	Fax Not Available +(1) 800-708-9888	
Website Not Available www.mgchemicals.com		www.mgchemicals.com
Email sales@mgchemicals.com Info@mgchemicals.com		Info@mgchemicals.com

1.4. Emergency telephone number

	Association / Organisation	CHEMTREC	Not Available	
	Emergency telephone numbers	+(44) 870-8200418	Not Available	
Other emergency telephone numbers +(1) 703-527-3887		+(1) 703-527-3887	Not Available	

SECTION 2 HAZARDS IDENTIFICATION

2.1.Classification of the substance or mixture

Classification according to regulation (EC) No 1272/2008 [CLP] ^[1]	Skin Corrosion/Irritation Category 2, Eye Irritation Category 2, Skin Sensitizer Category 1, Chronic Aquatic Hazard Category 2
Legend:	1. Classified by Chemwatch; 2. Classification drawn from EC Directive 67/548/EEC - Annex I ; 3. Classification drawn from EC Directive 1272/2008 - Annex VI

2.2. Label elements

CLP label elements	
SIGNAL WORD	WARNING
Hazard statement(s)	

H315	auses skin irritation.	
H319	H319 Causes serious eye irritation.	
H317	May cause an allergic skin reaction.	
H411	H411 Toxic to aquatic life with long lasting effects.	

Supplementary statement(s)

Not Applicable

Precautionary statement(s) Prevention

P280	Wear protective gloves/protective clothing/eye protection/face protection.	
P261	P261 Avoid breathing dust/fume/gas/mist/vapours/spray.	
P273	Avoid release to the environment.	
P272 Contaminated work clothing should not be allowed out of the workplace.		

Precautionary statement(s) Response

P302+P352	ON SKIN: Wash with plenty of water and soap.		
P305+P351+P338	IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.		
P333+P313	xin irritation or rash occurs: Get medical advice/attention.		
P337+P313	If eye irritation persists: Get medical advice/attention.		
P362+P364	Take off contaminated clothing and wash it before reuse.		
P391	Collect spillage.		

Precautionary statement(s) Storage

Not Applicable

Precautionary statement(s) Disposal

Dispose of contents/container in accordance with local regulations.

2.3. Other hazards

Cumulative effects may result following exposure*.

P501

Limited evidence of a carcinogenic effect*.

REACh - Art.57-59: The mixture does not contain Substances of Very High Concern (SVHC) at the SDS print date.

SECTION 3 COMPOSITION / INFORMATION ON INGREDIENTS

3.1.Substances

See 'Composition on ingredients' in Section 3.2

3.2.Mixtures

1.CAS No 2.EC No 3.Index No 4.REACH No	%[weight]	Name	Classification according to regulation (EC) No 1272/2008 [CLP]	
1.25085-99-8 2.500-033-5 3.603-074-00-8 4.01-2119456619-26-XXXX	89	bisphenol A/ diglycidyl ether resin, liquid	Eye Irritation Category 2, Skin Corrosion/Irritation Category 2, Skin Sensitizer Category 1, Chronic Aquatic Hazard Category 2; H319, H315, H317, H411 ^[3]	
1.17557-23-2 2.241-536-7 3.603-094-00-7 4.Not Available	6	neopentyl glycol diglycidyl ether	Skin Corrosion/Irritation Category 2, Skin Sensitizer Category 1; H315, H317 ^[3]	
1.64741-65-7. 2.265-067-2 3.649-275-00-4 4.01-2119850115-46-XXXX	2	naphtha petroleum, heavy alkylate	Flammable Liquid Category 3, Specific target organ toxicity - single exposure Category 3(narcotic effects), Aspiration Hazard Category 1; H226, H336, H304 ^[1]	
1.25068-38-6 2.500-033-5 3.603-074-00-8 4.01-2119456619-26-XXXX	1	bisphenol A diglycidyl ether resin, solid	Eye Irritation Category 2, Skin Corrosion/Irritation Category 2, Skin Sensitizer Category 1, Chronic Aquatic Hazard Category 2; H319, H315, H317, H411 ^[3]	
1.1333-86-4 2.215-609-9 3.Not Available 4.01-2119384822-32-XXXX, 01-2119489801-30-XXXX, 01-2119475601-40-XXXX	0.4	carbon black	Carcinogenicity Category 2; H351 ^[1]	
1.68609-97-2 2.271-846-8 3.603-103-00-4 4.01-2119485289-22-XXXX	0.3	(C12-14)alkylglycidyl ether	Skin Corrosion/Irritation Category 2, Skin Sensitizer Category 1; H315, H317 ^[3]	
Legend:	1. Classified by Chernwatch; 2. Classification drawn from EC Directive 67/548/EEC - Annex I ; 3. Classification drawn from EC Directive 1272/2008 - Annex VI 4. Classification drawn from C&L			

SECTION 4 FIRST AID MEASURES

4.1. Description of first aid measures

General	 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. If this product comes in contact with the eyes: Wash out immediately with fresh running water. Ensure complete irrigation of the eye by keeping eyelids apart and away from eye and moving the eyelids by occasionally lifting the upper and lower lids. Seek medical attention without delay; if pain persists or recurs seek medical attention. Removal of contact lenses after an eye injury should only be undertaken by skilled personnel. If fumes, aerosols or combustion products are inhaled remove from contaminated area. Other measures are usually unnecessary. Immediately give a glass of water. First aid is not generally required. If in doubt, contact a Poisons Information Centre or a doctor.
Eye Contact	 If this product comes in contact with the eyes: Wash out immediately with fresh running water. Ensure complete irrigation of the eye by keeping eyelids apart and away from eye and moving the eyelids by occasionally lifting the upper and lower lids. Seek medical attention without delay; if pain persists or recurs seek medical attention. Removal of contact lenses after an eye injury should only be undertaken by skilled personnel.
Skin Contact	 If skin contact occurs: Immediately remove all contaminated clothing, including footwear. Flush skin and hair with running water (and soap if available). Seek medical attention in event of irritation.
Inhalation	 If fumes, aerosols or combustion products are inhaled remove from contaminated area. Other measures are usually unnecessary.
Ingestion	 Immediately give a glass of water. First aid is not generally required. If in doubt, contact a Poisons Information Centre or a doctor.

4.2 Most important symptoms and effects, both acute and delayed

See Section 11

4.3. Indication of any immediate medical attention and special treatment needed

Treat symptomatically.

SECTION 5 FIREFIGHTING MEASURES

5.1. Extinguishing media

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

5.2. Special hazards arising from the substrate or mixture

Fire Incompatibility	Fire Incompatibility • Avoid contamination with oxidising agents i.e. nitrates, oxidising acids, chlorine bleaches, pool chlorine etc. as ignition may result				
5.3. Advice for firefighters					
Fire Fighting	 Alert Fire Brigade and tell them location and nature of hazard. Wear full body protective clothing with breathing apparatus. Prevent, by any means available, spillage from entering drains or water course. Use water delivered as a fine spray to control fire and cool adjacent area. Avoid spraying water onto liquid pools. DO NOT approach containers suspected to be hot. Cool fire exposed containers with water spray from a protected location. If safe to do so, remove containers from path of fire. 				
Fire/Explosion Hazard	 Combustible. Slight fire hazard when exposed to heat or flame. Heating may cause expansion or decomposition leading to violent rupture of containers. On combustion, may emit toxic fumes of carbon monoxide (CO). May emit acrid smoke. Mists containing combustible materials may be explosive. Combustion products include; carbon dioxide (CO2) other pyrolysis products typical of burning organic material 				

SECTION 6 ACCIDENTAL RELEASE MEASURES

6.1. Personal precautions, protective equipment and emergency procedures

See section 8

6.2. Environmental precautions

See section 12

6.3. 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. Moderate hazard. Clear area of personnel and move upwind. Alert Fire Brigade and tell them location and nature of hazard. Wear breathing apparatus plus protective gloves. Prevent, by any means available, spillage from entering drains or water course. No smoking, naked lights or ignition sources. Increase ventilation. Stop leak if safe to do so. Contain spill with sand, earth or vermiculite. Collect recoverable product into labelled containers for recycling. Absorb remaining product with sand, earth or vermiculite. Collect solid residues and seal in labelled drums for disposal. Wash area and prevent runoff into drains. If contamination of drains or waterways occurs, advise emergency services.

6.4. Reference to other sections

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

SECTION 7 HANDLING AND STORAGE

7.1. Precautions for safe 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. Avoid smoking, naked lights or ignition sources. 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. Always wash hands with scap and water after handling. Work clothes should be laundered separately. Use good occupational work practice. Observe manufacturer's storage and handling recommendations contained within this SDS. Atmosphere should be regularly checked against established exposure standards to ensure safe working conditions. DO NOT allow clothing wet with material to stay in contact with skin
Fire and explosion protection	See section 5
Other information	 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.

7.2. Conditions for safe storage, including any incompatibilities

Suitable container	 Metal can or drum Packaging as recommended by manufacturer. Check all containers are clearly labelled and free from leaks.
Storage incompatibility	 Avoid cross contamination between the two liquid parts of product (kit). If two part products are mixed or allowed to mix in proportions other than manufacturer's recommendation, polymerisation with gelation and evolution of heat (exotherm) may occur. This excess heat may generate toxic vapour Avoid reaction with amines, mercaptans, strong acids and oxidising agents Glycidyl ethers: may form unstable peroxides on storage in air ,light, sunlight, UV light or other ionising radiation, trace metals - inhibitor should be maintained at adequate levels may polymerise in contact with heat, organic and inorganic free radical producing initiators may polymerise with evolution of heat in contact with oxidisers, strong acids, bases and amines react violently with strong oxidisers, permanganates, peroxides, acyl halides, alkalis, ammonium persulfate, bromine dioxide attack some forms of plastics, coatings, and rubber

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)

Not Available

PREDICTED NO EFFECT LEVEL (PNEC)

Not Available

OCCUPATIONAL EXPOSURE LIMITS (OEL)

Source	Ingredient	Material name	TWA	STEL		Peak	Notes
UK Workplace Exposure Limits (WELs)	carbon black	Carbon black	3.5 mg/m3	7 mg/m	3	Not Available	Not Available
EMERGENCY LIMITS							
Ingredient	Material name			TE	EL-1	TEEL-2	TEEL-3
bisphenol A/ diglycidyl ether resin, liquid	Epoxy resin (EPON 1001)		90	mg/m3	990 mg/m3	5900 mg/m3
bisphenol A/ diglycidyl ether resin, liquid	Epoxy resin (EPON 1007)		90	mg/m3	990 mg/m3	5900 mg/m3
bisphenol A/ diglycidyl ether resin, liquid	Epoxy resin (EPON 820)			41	mg/m3	450 mg/m3	2700 mg/m3
bisphenol A/ diglycidyl ether resin, liquid	Epoxy resin ERL-2795			32	mg/m3	350 mg/m3	2100 mg/m3
bisphenol A diglycidyl ether resin, solid	Epoxy resin (EPON 1001)		90	mg/m3	990 mg/m3	5900 mg/m3
bisphenol A diglycidyl ether resin, solid	Epoxy resin (EPON 1007	Epoxy resin (EPON 1007)			mg/m3	990 mg/m3	5900 mg/m3
bisphenol A diglycidyl ether resin, solid	Epoxy resin (EPON 820)	Epoxy resin (EPON 820)			mg/m3	450 mg/m3	2700 mg/m3
bisphenol A diglycidyl ether resin, solid	Epoxy resin ERL-2795	Epoxy resin ERL-2795 Polypropylene glycol, (chloromethyl) oxirane polymer			mg/m3	350 mg/m3	2100 mg/m3
bisphenol A diglycidyl ether resin, solid	Polypropylene glycol, (chl				6 mg/m3	66 mg/m3	400 mg/m3
carbon black	Carbon black			9 n	ng/m3	99 mg/m3	590 mg/m3
Ingredient	Original IDLH				Revised	IDLH	
bisphenol A/ diglycidyl ether resin, liquid	Not Available				Not Available		
neopentyl glycol diglycidyl ether	Not Available			Not Available			
naphtha petroleum, heavy alkylate	Not Available			Not Available			
bisphenol A diglycidyl ether resin, solid	Not Available				Not Available		
carbon black	N.E. mg/m3 / N.E. ppm	N.E. mg/m3 / N.E. ppm			1,750 mg/m3		
(C12-14)alkylglycidyl ether	Not Available				Not Available		

MATERIAL DATA

Sensory irritants are chemicals that produce temporary and undesirable side-effects on the eyes, nose or throat. Historically occupational exposure standards for these irritants have been based on observation of workers' responses to various airborne concentrations. Present day expectations require that nearly every individual should be protected against even minor sensory irritation and exposure standards are established using uncertainty factors or safety factors of 5 to 10 or more. On occasion animal no-observable-effect-levels (NOEL) are used to determine these limits where human results are unavailable. An additional approach, typically used by the TLV committee (USA) in determining respiratory standards for this group of chemicals, has been to assign ceiling values (TLV C) to rapidly acting irritants and to assign short-term exposure limits (TLV STELs) when the weight of evidence from irritation, bioaccumulation and other endpoints combine to warrant such a limit. In contrast the MAK Commission (Germany) uses a five-category system based on intensive odour, local irritation, and elimination half-life. However this system is being replaced to be consistent with the European Union (EU) Scientific Committee for Occupational Exposure Limits (SCOEL); this is more closely allied to that of the USA. OSHA (USA) concluded that exposure to sensory irritants can:

cause inflammation

cause increased susceptibility to other irritants and infectious agents

lead to permanent injury or dysfunction

permit greater absorption of hazardous substances and

+ acclimate the worker to the irritant warning properties of these substances thus increasing the risk of overexposure.

8.2. Exposure controls

	Engineering controls are used to remove a hazard or place a barrier between the worker and the hazard. Well-designed engineering or 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 strat 'removes' air in the work environment. Ventilation can remove or dilute an air contaminant if designed properly. The design of a ventilation the particular process and chemical or contaminant in use.	tegically 'adds' and
8.2.1. Appropriate		
engineering controls		
General exhaust is adequate under normal operating conditions. Local exhaust ventilation may be required in specific circumstance exists, wear approved respirator. Correct fit is essential to obtain adequate protection. Provide adequate ventilation in warehouse of contaminants generated in the workplace possess varying 'escape' velocities which, in turn, determine the 'capture velocities' of fit to effectively remove the contaminant.		osed storage areas. Air
	Type of Contaminant:	Air Speed:

	solvent, vapours, degreasing etc., evaporating from tank (in still air). 0 aerosols, fumes from pouring operations, intermittent container filling, low speed conveyer transfers, welding, spray drift, plating 0 aerosols, fumes, pickling (released at low velocity into zone of active generation) 0 direct spray, spray painting in shallow booths, drum filling, conveyer loading, crusher dusts, gas discharge (active generation into zone of rapid air motion) 1 grinding, abrasive blasting, tumbling, high speed wheel generated dusts (released at high initial velocity into zone of very high rapid air motion). 2		
	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 extr of distance from the extraction point (in simple cases). Therefore the air speed at the extraction point distance from the contaminating source. The air velocity at the extraction fan, for example, should be a solvents generated in a tank 2 meters distant from the extraction point. Other mechanical considerati apparatus, make it essential that theoretical air velocities are multiplied by factors of 10 or more when	should be adjusted, accordingly, a a minimum of 1-2 m/s (200-400 f/m ons, producing performance deficit	after reference to nin) for extraction of ts within the extraction
8.2.2. Personal protection			
	 Safety glasses with side shields. Chemical goggles. 		
Eye and face protection	 Contact lenses may pose a special hazard; soft contact lenses may absorb and concentrate irrita lenses or restrictions on use, should be created for each workplace or task. This should include a chemicals in use and an account of injury experience. Medical and first-aid personnel should be readily available. In the event of chemical exposure, begin eye irrigation immediately and remove at the first signs of eye redness or irritation - lens should be removed in a clean environment only Current Intelligence Bulletin 59], [AS/NZS 1336 or national equivalent] 	a review of lens absorption and ac trained in their removal and suitab contact lens as soon as practicabl	Isorption for the class of ble equipment should be le. Lens should be remov
Eye and face protection Skin protection	Contact lenses may pose a special hazard; soft contact lenses may absorb and concentrate irritral lenses or restrictions on use, should be created for each workplace or task. This should include a chemicals in use and an account of injury experience. Medical and first-aid personnel should be readily available. In the event of chemical exposure, begin eye irrigation immediately and remove at the first signs of eye redness or irritation - lens should be removed in a clean environment only	a review of lens absorption and ac trained in their removal and suitab contact lens as soon as practicabl	Isorption for the class of ble equipment should be le. Lens should be remov
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Skin protection Hands/feet protection	 Contact lenses may pose a special hazard; soft contact lenses may absorb and concentrate irrital lenses or restrictions on use, should be created for each workplace or task. This should include a chemicals in use and an account of injury experience. Medical and first-aid personnel should be readily available. In the event of chemical exposure, begin eye irrigation immediately and remove at the first signs of eye redness or irritation - lens should be removed in a clean environment only Current Intelligence Bulletin 59], [AS/NZS 1336 or national equivalent] See Hand protection below NOTE: The material may produce skin sensitisation in predisposed individuals. Care must be taken, whe all possible skin contact. Contaminated leather items, such as shoes, belts and watch-bands should be removed and destr The selection of suitable gloves does not only depend on the material, but also on further marks of que the chemical is a preparation of several substances, the resistance of the glove material can not be care to the application. The exact break through time for substances has to be obtained from the manufacturer of the protectiv choice. Suitability and durability of glove type is dependent on usage. Important factors in the selection of glove frequency and duration of contact, chemical resistance of glove material, glove thickness and dexterity Select gloves tested to a relevant standard (e.g. Europe EN 374, US F739, AS/NZS 2161.1 or nation 2161.10.1 or national equivalent) is recommended. When norloged or frequently repeated contact may occur, a glove with a protection class of 5 o 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	a review of lens absorption and ad trained in their removal and suitab contact lens as soon as practicabl after workers have washed hands an removing gloves and other prote oyed. ality which vary from manufacturer alculated in advance and has there re gloves and has to be observed v ves include: hal equivalent). In higher (breakthrough time greated time greater than 60 minutes accor- when considering gloves for long-to- roughly. Application of a non-perfu- butatoluene rubber), boots and ap- per or polyethylene gloves (which a	disorption for the class of ble equipment should be le. Lens should be remove is thoroughly. [CDC NIOS ective equipment, to avoid to manufacturer. Where efore to be checked prior when making a final er than 240 minutes ording to EN 374, AS/NZ erm use. umed moisturiser is prons. absorb the resin).
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Respiratory protection

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

Where the concentration of gas/particulates in the breathing zone, approaches or exceeds the 'Exposure Standard' (or ES), respiratoryprotection is required. Degree of protection varies with both face-piece and Class offilter; the nature of protection varies with Type of filter.

Required Minimum Protection Factor	Half-Face Respirator	Full-Face Respirator	Powered Air Respirator
up to 10 x ES	A-AUS P2	-	A-PAPR-AUS / Class 1 P2

up to 50 x ES	-	A-AUS / Class 1 P2	-
up to 100 x ES	-	A-2 P2	A-PAPR-2 P2 ^

^ - Full-face

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

8.2.3. Environmental exposure controls

See section 12

SECTION 9 PHYSICAL AND CHEMICAL PROPERTIES

9.1. Information on basic physical and chemical properties

Appearance	black		
Physical state	Liquid	Relative density (Water = 1)	1.15
Odour	Slight	Partition coefficient n-octanol / water	Not Available
Odour threshold	Not Available	Auto-ignition temperature (°C)	>235
pH (as supplied)	Not Available	Decomposition temperature	Not Available
Melting point / freezing point (°C)	Not Available	Viscosity (cSt)	5850
Initial boiling point and boiling range (°C)	>150	Molecular weight (g/mol)	Not Available
Flash point (°C)	>150	Taste	Not Available
Evaporation rate	Not Available	Explosive properties	Not Available
Flammability	Not Applicable	Oxidising properties	Not Available
Upper Explosive Limit (%)	Not Available	Surface Tension (dyn/cm or mN/m)	Not Available
Lower Explosive Limit (%)	Not Available	Volatile Component (%vol)	Not Available
Vapour pressure (kPa)	Not Available	Gas group	Not Available
Solubility in water (g/L)	Miscible	pH as a solution (1%)	Not Available
Vapour density (Air = 1)	>1	VOC g/L	Not Available

9.2. Other information

Not Available

SECTION 10 STABILITY AND REACTIVITY

10.1.Reactivity	See section 7.2
10.2.Chemical stability	 Unstable in the presence of incompatible materials. Product is considered stable. Hazardous polymerisation will not occur.
10.3. Possibility of 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

SECTION 11 TOXICOLOGICAL INFORMATION

11.1. Information on toxicological effects

Inhaled	The material is not thought to produce adverse health effects or irritation of the respiratory tract (as classified by EC Directives using animal models). Nevertheless, good hygiene practice requires that exposure be kept to a minimum and that suitable control measures be used in an occupational setting.
Ingestion	The material has NOT been classified by EC Directives or other classification systems as 'harmful by ingestion'. This is because of the lack of corroborating animal or human evidence. The material may still be damaging to the health of the individual, following ingestion, especially where pre-existing organ (e.g liver, kidney) damage is evident. Present definitions of harmful or toxic substances are generally based on doses producing mortality rather than those producing morbidity (disease, ill-health). Gastrointestinal tract discomfort may produce nausea and vomiting. In an occupational setting however, ingestion of insignificant quantities is not thought to be cause for concern.
Skin Contact	Evidence exists, or practical experience predicts, that the material either produces inflammation of the skin in a substantial number of individuals following direct contact, and/or produces significant inflammation when applied to the healthy intact skin of animals, for up to four hours, such inflammation being present twenty-four hours or more after the end of the exposure period. Skin irritation may also be present after prolonged or repeated exposure; this may result in a form of contact dermatitis (nonallergic). The dermatitis is often characterised by skin redness (erythema) and swelling (oedema) which may progress to blistering (vesiculation), scaling and thickening of the epidermis. At the microscopic level there may be intercellular oedema of the spongy layer of the skin (spongiosis) and intracellular oedema of the epidermis. The material may accentuate any pre-existing dermatitis condition

	Skin contact is not thought to have harmful health effects (as classified under EC Directives); the material may still produce health damage following entry through wounds, lesions or abrasions. 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	Evidence exists, or practical experience predicts, that the material may cause eye irritation in a substantial number of individuals and/or may produce significant ocular lesions which are present twenty-four hours or more after instillation into the eye(s) of experimental animals. Repeated or prolonged eye contact may cause inflammation characterised by temporary redness (similar to windburn) of the conjunctiva (conjunctivitis); temporary impairment of vision and/or other transient eye damage/ulceration may occur.			
Chronic	Practical experience shows that skin contact with the material is capable either of inducing a sensitisation reaction in a substantial number of individuals, and/or of producing a positive response in experimental animals. Limited evidence suggests that repeated or long-term occupational exposure may produce cumulative health effects involving organs or biochemical systems. All glycidyl ethers show genotoxic potential due their alkylating properties. Those glycidyl ethers that have been investigated in long term studies exhibit more or less marked carcinogenic potential. Alkylating agents may damage the stem cell which acts as the precursor to components of the blood. Loss of the stem cell may result in pancytopenia (a reduction in the number of red and white blood cells and platelets) with a latency period corresponding to the lifetime of the individual blood cells. Granulocytopenia (a reduction in granular leukocytes) develops within days and thrombocytopenia (a disorder involving platelets), within 1-2 weeks, whilst loss of erythrocytes (red blood cells) need months to become clinically manifest. Aplastic anaemia develops due to complete destruction of the stem cells. Glycidyl ethers have been shown to cause allergic contact dermatitis in humans. Glycidyl ethers generally cause skin sensitization in experimental animals. Necrosis of the mucous membranes of the nasal cavities was induced in mice exposed to allyl glycidyl ether. A study of workers with mixed exposures was inconclusive with regard to the effects of specific glycidyl ethers. Phenyl glycidyl ether, but not <i>n</i> -butyl glycidyl ether did not induce micronuclei or chromosomal aberrations <i>in vivo</i> following intraperitoneal but not oral administration. Phenyl glycidyl ether did not induce DNA damage in cultured human cells or mutation in cultured animal cells. <i>in vivo</i> o following intraperitoneal but not oral administration. Phenyl glycidyl ether did not induce DNA damage in cultured human cells or mutation in cultured animal cells. Allyl glycidyl et			
832HD 1:1 Epoxy	ΤΟΧΙCΙΤΥ	IRRITATI	ON	
Potting and Encapsulating Compound (Part A)	Not Available	Not Availa	-	
bisphenol A/ diglycidyl ether				
resin, liquid	dermal (rat) LD50: >800 mg/kg ^[1] Oral (rat) LD50: 13447 mg/kg ^[1]		ye (rabbit): 100mg -	IVIIG
	ΤΟΧΙCΙΤΥ		IRRITATION	
neopentyl glycol diglycidyl			Skin (human): Se	ensitiser [Shell]
ether	Oral (rat) LD50: 4500 mg/kg] ^[2]			
	ΤΟΧΙCITY			IRRITATION
	Dermal (rabbit) LD50: >5000 mg/kg ^[2]			Not Available
naphtha petroleum, heavy alkylate	Inhalation (rat) LC50: >3.83 mg/L/4H ^[2]			
	Initialation (rat) LC50: >3.83 mg/L/4H* Oral (rat) LD50: >25000 mg/kg ^[2]			
	ΤΟΧΙCΙΤΥ			IRRITATION
bisphenol A diglycidyl ether	dermal (rat) LD50: >800 mg/kg ^[1]		Nil reported	
resin, solid	Oral (rat) LD50: 13447 mg/kg ^[1]			
	ΤΟΧΙΟΙΤΥ			IRRITATION
carbon black	Dermal (rabbit) LD50: >3000 mg/kg ^[2]			Not Available
	Oral (rat) LD50: >8000 mg/kg ^[1]			
	ΤΟΧΙCITY	IRRIT	ATION	
	Oral (rat) LD50: 16896 mg/kg ^[1] Eye (rabbit): mild [Ciba]			
	Skin (guinea pig): sensitiser		er	
(C12-14)alkylglycidyl ether			uman): Irritant	
		,	uman): non- sensitis	ser
			abbit): moderate	
Legend:	1. Value obtained from Europe ECHA Registered Substances - Acute toxicity 2 extracted from RTECS - Register of Toxic Effect of chemical Substances	2.* Value obta	ained from manufact	urer's SDS. Unless otherwise specified data

Continued...

832HD 1:1 Epoxy Potting and Encapsulating Compound (Part A) Sight Bisphe activity hormoo ring ar bridgin acteritylos for 1.2 Ethylo papilla inhalat papilla inhalat observ weeks at wee substat The for endoc Sight Sight Potting and Encapsulating Compound (Part A) The for endoc Sight Sight <tr< th=""><th>Ilowing information refers to contact allergens as a group and may not be specific to this product. t allergies quickly manifest themselves as contact eczema, more rarely as urlicaria or Quincke's oedema. The pathogenesis of contact eczema involves mediated (T lymphocytes) immune reaction of the delayed type. Other allergic skin reactions, e.g. contact urticaria, involve antibody-mediated immune ns. The significance of the contact allergen is not simply determined by its sensitisation potential: the distribution of the substance and the opportunities tact with it are equally important. A weakly sensitising substance which is widely distributed can be a more important allergen than one with stronger sing potential with which few individuals come into contact. From a clinical point of view, substances are noteworthy if they produce an allergic test n in more than 1% of the persons tested. termical structure of hydroxylated diphenylalkanes or bisphenols consists of two phenolic rings joined together through a bridging carbon. This class of fine disruptors that mimic oestrogens is widely used in industry, particularly in plastics nol A (BPA) and some related compounds exhibit oestrogenic activity in human breast cancer cell line MCF-7, but there were remarkable differences in a Several derivatives of BPA exhibited significant thyroid hormonal activity towards rat pituitary cell line GH3, which releases growth hormone in a thyroid the B-phenyl ring of BPA derivatives are required for these hormonal activities, and substituents at the 3,5-positions of the phenyl rings and the g alkyl moiety markedly influence the activities. anots promoted cell proliferation and increased the synthesis and secretion of cell type-specific proteins. When ranked by proliferative potency, the longer y substituent at the bridging carbon, the lower the concentration needed for maximal callyield; the most active compound contained two propyl chains at diging carbon. Bisphenols with two hydroxyl groups in the para position and</th></tr<>	Ilowing information refers to contact allergens as a group and may not be specific to this product. t allergies quickly manifest themselves as contact eczema, more rarely as urlicaria or Quincke's oedema. The pathogenesis of contact eczema involves mediated (T lymphocytes) immune reaction of the delayed type. Other allergic skin reactions, e.g. contact urticaria, involve antibody-mediated immune ns. The significance of the contact allergen is not simply determined by its sensitisation potential: the distribution of the substance and the opportunities tact with it are equally important. A weakly sensitising substance which is widely distributed can be a more important allergen than one with stronger sing potential with which few individuals come into contact. From a clinical point of view, substances are noteworthy if they produce an allergic test n in more than 1% of the persons tested. termical structure of hydroxylated diphenylalkanes or bisphenols consists of two phenolic rings joined together through a bridging carbon. This class of fine disruptors that mimic oestrogens is widely used in industry, particularly in plastics nol A (BPA) and some related compounds exhibit oestrogenic activity in human breast cancer cell line MCF-7, but there were remarkable differences in a Several derivatives of BPA exhibited significant thyroid hormonal activity towards rat pituitary cell line GH3, which releases growth hormone in a thyroid the B-phenyl ring of BPA derivatives are required for these hormonal activities, and substituents at the 3,5-positions of the phenyl rings and the g alkyl moiety markedly influence the activities. anots promoted cell proliferation and increased the synthesis and secretion of cell type-specific proteins. When ranked by proliferative potency, the longer y substituent at the bridging carbon, the lower the concentration needed for maximal callyield; the most active compound contained two propyl chains at diging carbon. Bisphenols with two hydroxyl groups in the para position and
Conta a cell-r reaction for con sensiti reaction The ch endoc Bisphe activity hormoon ring ar bridgin Bisphe the alk the brin accept The st NOT on Evider	nces, oxirane (ethylene oxide) and methyloxirane (propylene oxide), which are also direct-acting alkylating agents, have been classified as carcinogenic
BISPHENOL A/ DIGLYCIDYL ETHER RESIN, LIQUID RESIN, LIQUID In alife oneou etal., 1 weakly expose 1997). Genot in TAse al., 197 host-r immu weekir - Consu BADG Proview ranges negati fromsu weigh/ human	noes, ourse, tempere oxee) and memploatine (prophere oxee), which are also altech-doing alivplang agents, have been classified as calcinopence lively information refers to contact allergens as a group and may not be specific to his product. Ta ellergies quickly manifest themselves as contact eczema, more rarely as uricaria or Quinck's oederna. The pathogenesis of contact eczema involves mediand (T improvplas) immune reaction of the delayed by pc. Other allergic akin reactions, a g. contact uticaris, involve antabody-mediated immune rars. The significance of the contact allergen is not simply determined by its sensitisation potential; the distribution of the subtance and the opportunities tate with its are qualy important. A weakly sensitivitary setsion in the other of the subtances are noteworthy if they produce an allergic test in more than '19' of the parsons tested. The distribution of hydroxylated diphenylakanes or bisphenols consists of two phenolic rings joined together through a bridging carbon. This class of the distructors that minic extremores is subdely used in industry particularly in plastics. The APA and some related compounds exhibit estrogenic activity in human breast cancer cell line KHZ, which neleases growth hormone in a thyrid the dispersion that all BPA charinkes are reguined for these hormonal activities, and substituents at the 3.5-positions of the phenyl rings and the g alg/ mixing immandaly influence the activities. The particular allergin carbon, the lower the concentration needed for maximal call yiek; the most active compound contained two propi chains at ding gatomate process. The activity and the hydroxy group of the Aphenyl dist of the coarbing needee. The classified by IARC as Group 3: assistable to is distance and tay to mytepic is and secretion of cell type-specific proteins. When rarked by profileration to the assistable of substituents and a role mytepic is then advised and the maxima and all yiek; the most active companyl to the aphenyl dig gatoma. Bephenol

Foetoxicity has been observed in animal studies Oral (rabbit, female) NOEL 180 mg/kg (teratogenicity; NOEL (maternal 60 mg/kg

NEOPENTYL GLYCOL DIGLYCIDYL ETHER	* Anchor SDS]		
NAPHTHA PETROLEUM, HEAVY ALKYLATE	Studiesindicate that normal, branched and cyclic paraffins are absorbed from themammalian gastrointestinal tract and that the absorption of n-paraffins isinversely proportional to the carbon chain length, with little absorption aboveC30. With respect to the carbon chain lengths likely to be present in mineraloil, n-paraffins may be absorbed to a greater extent that iso- orcyclo-paraffins. The majorclasses of hydrocarbons have been shown to be well absorbed by thegastrointestinal tract in various species. In many cases, the hydrophobichydrocarbons are ingested in association with dietary lipids. The dependence ofhydrocarbon and sorption on concomitant triglyceride digestion and absorption jakinown as the hydrocarbon continuum hypothesis', and asserts that aseries of solubilising phases in the intestinal lumen, created by dietaryfigiverides and their digestion products, afford hydrocarbons aroute to thelipid phase of the intestinal absorptive cell (enterocyte) membrane. While somehydrocarbons partially separate from nutrient lipids and undergo metabolictransformation in the enterocyte. The enterocyte may play a major role indetermining the proportion of an absorbed hydrocarbon that, by escaping initialbiotransformation in the enterocyte. The enterocyte may play a major role indetermining the proportion of an absorbed nydrocarbon that, by escaping initialbiotransformation, becomes available for deposition in its unchanged form nurrent lipids. The dependence of tumours in rodents. The renterocyte may play a major role indetermining the proportion of an absorbed nydrocarbon that, by escaping initialbiotransformation in the enterocyte. The enterocyte may lead to hearing loss. This product contains toluene. There are indications from animal studies that prolonged exposure to high concentrations of toluene may lead to hearing loss. This product contains toluene. There are indications from animal studies on gasoline and gasoline blending streams, which use a wide variety of endpoints and give predunitary to hydrocarbo.		
BISPHENOL A DIGLYCIDYL ETHER RESIN, SOLID	alpha2-microglobulin is produced under the influence of hormonal controls in male rats but not in females and, more importantly, not in humans. The following information refers to contact allergens as a group and may not be specific to this product. Contact allergies quickly manifest themselves as contact eczema, more rarely as urticaria or Quincke's oedema. The pathogenesis of contact eczema involves a cell-mediated (T lymphocytes) immune reaction of the delayed type. Other allergic skin reactions, e.g. contact uticaria, involve antibody-mediated immune reactions. The significance of the contact allergen is not simply determined by its sensitisation potential: the distribution of the substance and the opportunities for contact with it are equally important. A weakly sensitising substance which is widely distributed can be a more important allergen than one with stronger sensitising potential with which few individuals come into contact. From a clinical point of view, substances are noteworthy if they produce an allergic test reaction in more than 1% of the persons tested. The chemical structure of hydroxylated diphenylalkanes or bisphenols consists of two phenolic rings joined together through a bridging carbon. This class of endocrine disruptors that mimic oestrogens is widely used in industry, particularly in plastics Bisphenol A (BPA) and some related compounds exhibit estrogenic activity in human breast cancer cell line MCF-7, but there were remarkable differences in activity. Several derivatives of BPA exhibited significant thyroid hormonal activity towards rat pitutary cell line GH3, which releases growth hormone in a thyroid hormone-dependent manner. However, BPA and several other derivatives did not show such activity. Results suggest that the 4-hydroxyl group of the A-phenyl ring and the B-phenyl ring of BPA derivatives are required for these hormonal activities, and substituents at the 3,5-positions of the phenyl rings and the bridging alkly molety markedly influence the activities. Bi		
CARBON BLACK	occurrence of these is greatly reduced in solid grades of the resin. No significant acute toxicological data identified in literature search. WARNING: This substance has been classified by the IARC as Group 2B: Possibly Carcinogenic to Humans.		
NEOPENTYL GLYCOL DIGLYCIDYL ETHER & (C12-14)ALKYLGLYCIDYL ETHER	Inhalation (rat) TCLo: 50 mg/m3/6h/90D-I Nil reported The following information refers to contact allergens as a group and may not be specific to this product. Contact allergies quickly manifest themselves as contact eczema, more rarely as urticaria or Quincke's oedema. The pathogenesis of contact eczema involves a cell-mediated (T lymphocytes) immune reaction of the delayed type. Other allergic skin reactions, e.g. contact urticaria, involve antibody-mediated immune reactions. The significance of the contact allergen is not simply determined by its sensitisation potential: the distribution of the substance and the opportunities for contact with it are equally important. A weakly sensitising substance which is widely distributed can be a more important allergen than one with stronger sensitising potential with which few individuals come into contact. From a clinical point of view, substances are noteworthy if they produce an allergic test reaction in more than 1% of the persons tested.		
Acute Toxicity	⊗ Car	rcinogenicity 🛇	
Skin Irritation/Corrosion		productivity 🛇	
Serious Eye Damage/Irritation		le Exposure	
Respiratory or Skin sensitisation	✓ STOT - Repeate	ed Exposure	
Mutagenicity	⊗ Aspira	ation Hazard	
		Legend: X − Data available but does not fill the criteria for classification ↓ − Data required to make classification available ○ − Data Not Available to make classification	

12.1. Toxicity

Ingredient	Endpoint	Test Duration (hr)	Species	Value	Source
bisphenol A/ diglycidyl ether resin, liquid	LC50	96	Fish	1.2mg/L	2
bisphenol A/ diglycidyl ether resin, liquid	EC50	48	Crustacea	1.1mg/L	2
bisphenol A/ diglycidyl ether resin, liquid	EC50	48	Crustacea	1.7mg/L	2
bisphenol A/ diglycidyl ether resin, liquid	NOEC	504	Crustacea	0.3mg/L	2
bisphenol A/ diglycidyl ether resin, liquid	EC50	72	Algae or other aquatic plants	9.4mg/L	2
neopentyl glycol diglycidyl ether	LC50	96	Fish	12.318mg/L	3
naphtha petroleum, heavy alkylate	EC50	72	Algae or other aquatic plants	=13mg/L	1
naphtha petroleum, heavy alkylate	EC50	72	Algae or other aquatic plants	=30000mg/L	1
naphtha petroleum, heavy alkylate	NOEC	72	Algae or other aquatic plants	=0.1mg/L	1
bisphenol A diglycidyl ether resin, solid	LC50	96	Fish	1.2mg/L	2
bisphenol A diglycidyl ether resin, solid	EC50	48	Crustacea	1.1mg/L	2
bisphenol A diglycidyl ether resin, solid	EC50	48	Crustacea	1.7mg/L	2
bisphenol A diglycidyl ether resin, solid	NOEC	504	Crustacea	0.3mg/L	2
bisphenol A diglycidyl ether resin, solid	EC50	72	Algae or other aquatic plants	9.4mg/L	2
carbon black	LC50	96	Fish	>100mg/L	2
carbon black	NOEC	720	Fish	17mg/L	2
carbon black	EC50	48	Crustacea	>100mg/L	2
carbon black	EC50	384	Crustacea	4.9mg/L	2
carbon black	EC50	96	Algae or other aquatic plants	95mg/L	2
(C12-14)alkylglycidyl ether	LC50	96	Fish	>5000mg/L	2
(C12-14)alkylglycidyl ether	EC50	48	Crustacea	6.07mg/L	2
(C12-14)alkylglycidyl ether	NOEC	48	Crustacea	<10mg/L	2

Legend:

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. For bisphenol A and related bisphenols:

In general, studies have shown that bisphenol A can affect growth, reproduction and development in aquatic organisms. Among freshwater organisms, fish appear to be the most sensitive species. Evidence of endocrine-related effects in fish, aquatic invertebrates, amphibians and reptiles has been reported at environmentally relevant exposure levels lower than those required for acute toxicity. There is a widespread variation in reported values for endocrine-related effects, but many fall in the range of 1 ug/L to 1 mg/L

Bisphenol A, its derivatives and analogues, can be released from polymers, resins and certain substances by metabolic products

As an environmental contaminant, bisphenol A interferes with nitrogen fixation at the roots of leguminous plants associated with the bacterial symbiont Sinorhizobium meliloti. Despite a half-life in the soil of only 1-10 days, its ubiquity makes it an important pollutant. According to Environment Canada, 'initial assessment shows that at low levels, bisphenol A can harm fish and organisms over time. Studies also indicate that it can currently be found in municipal wastewater.' However, a study conducted in the United States found that 91-98% of bisphenol A may be removed from water during treatment at municipal water treatment plants.

A 2009 review of the biological impacts of plasticisers on wildlife published by the Royal Society with a focus on annelids (both aquatic and terrestrial), molluscs, crustaceans, insects, fish and amphibians concluded that bisphenol A has been shown to affect reproduction in all studied animal groups, to impair development in crustaceans and amphibians and to induce genetic aberrations. A large 2010 study of two rivers in Canada found that areas contaminated with hormone-like chemicals including bisphenol A showed females made up 85 per cent of the population of a certain fish, while females made up only 55 per cent in uncontaminated areas.

Although abundant data are available on the toxicity of bisphenol-A (2,2-bis (4-hydroxydiphenyl)propane;(BPA) A variety of BPs were examined for their acute toxicity against Daphnia magna, mutagenicity, and oestrogenic activity using the Daphtoxkit (Creasel Ltd.), the umu test system, and the yeast two-hybrid system, respectively, in comparison with BPA. BPA was moderately toxic to D. magna (48-h EC50 was 10 mg/l) according to the current U.S. EPA acute toxicity evaluation standard, and it was weakly oestrogenic with 5 orders of magnitude lower activity than that of the natural estrogen 17 beta-oestradiol in the yeast screen, while no mutagenicity was observed. All seven BPs tested here showed moderate to slight acute toxicity, no mutagenicity, and weak oestrogenic activity as well as BPA. Some of the BPs showed considerably higher oestrogenic activity than BPA, and others exhibited much lower activity. Bisphenol S (bis(4-hydroxydiphenyl)sulfide) showed oestrogenic activity.

Biodegradation is a major mechanism for eliminating various environmental pollutants. Studies on the biodegradation of bisphenols have mainly focused on bisphenol A. A number of BPA-degrading bacteria have been isolated from enrichments of sludge from wastewater treatment plants. The first step in the biodegradation of BPA is the hydroxylation of the carbon atom of a methyl group or the quaternary carbon in the BPA molecule. Judging from these features of the biodegradation mechanisms, it is possible that the same mechanism used for BPA is used to biodegrade all bisphenols that have at least one methyl or methylene group bonded at the carbon atom between the two phenol groups. However, bisphenol F ((bis(4-hydroxyphenyl)methane; BPF), which has no substituent at the bridging carbon, is unlikely to be metabolised by such a mechanism. Nevertheless BPF is readily degraded by river water microorganisms under aerobic conditions. From this evidence, it was clear that a specific mechanism for biodegradation of BPF does exist in the natural ecosystem,

Algae can enhance the photodegradation of bisphenols. The photodegradation rate of BPF increased with increasing algae concentration. Humic acid and Fe3+ ions also enhanced the photodegradation of BPF. The effect of pH value on the BPF photodegradation was also important.

Significant environmental findings are limited. Oxiranes (including glycidyl ethers and alkyl oxides, and epoxides) exhibit common characteristics with respect to environmental fate and

ecotoxicology. One such oxirane is ethyloxirane and data presented here may be taken as representative.

for 1,2-butylene oxide (ethyloxirane):

Environmental fate: Ethyloxirane is highly soluble in water and has a very low soil-adsorption coefficient, which suggests that if released to water, adsorption of ethyloxirane to sediment and suspended solids is not expected. Volatilisation of ethyloxirane from water surfaces would be expected based on the moderate estimated Henry's Law constant. If ethyloxirane is released to soil, it is expected to have low adsorption and thus very high mobility. Volatilisation from moist soil and dry soil surfaces is expected, based on its vapour pressure. It is expected that ethyloxirane exists solely as a vapour in ambient atmosphere, based on its very high vapour pressure. Ethyloxirane may also be removed from the atmosphere by wet deposition processes, considering its relatively high water solubility.

Persistence: The half-life in air is about 5.6 days from the reaction of ethyloxirane with photochemically produced hydroxyl radicals which indicates that this chemical meets the persistence criterion in air (half-life of = 2 days)*

Ethyloxirane is hydrolysable, with a half-life of 6.5 days, and biodegradable up to 100% degradation and is not expected to persist in water. A further model-predicted biodegradation half-life of 15 days in water was obtained and used to predict the half-life of this chemical in soil and sediment by applying Boethling's extrapolation factors (t1/2water: t1/2 soil: t1/2 sediment = 1: 1: 4) (Boethling 1995). According to these values, it can be concluded that ethyloxirane does not meet the persistence criteria in water and soil (half-lives = 182 days) and sediments (half-life = 365 days).

Experimental and modelled log Kow values of 0.68 and 0.86, respectively, indicate that the potential for bioaccumulation of ethyloxirane in organisms is likely to be low. Modelled bioaccumulation -factor (BAF) and bioconcentration -factor (BCF) values of 1 to 17 L/kg indicate that ethyloxirane does not meet the bioaccumulation criteria (BCF/BAF = 5000)* Ecotoxicity:

Experimental ecotoxicological data for ethyloxirane (OECD 2001) indicate low to moderate toxicity to aquatic organisms. For fish and water flea, acute LC50/EC50 values vary within a narrow range of 70-215 mg/L; for algae, toxicity values exceed 500 mg/L, while for bacteria they are close to 5000 mg/L

* Persistence and Bioaccumulation Regulations (Canada 2000).

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

Environmentaltoxicity is a function of the n-octanol/water partition coefficient (log Pow,log Kow). Compounds with log Pow >5 act as neutral organics, but at a lowerlog Pow, the toxicity of epoxidecontaining polymers is greater than that predicted for simple narcotics.

DO NOT discharge into sewer or waterways

12.2. Persistence and degradability

Ingredient	Persistence: Water/Soil	Persistence: Air
bisphenol A/ diglycidyl ether resin, liquid	HIGH	HIGH
neopentyl glycol diglycidyl ether	HIGH	HIGH
bisphenol A diglycidyl ether resin, solid	HIGH	HIGH

12.3. Bioaccumulative potential

Ingredient	Bioaccumulation
bisphenol A/ diglycidyl ether resin, liquid	LOW (LogKOW = 2.6835)
neopentyl glycol diglycidyl ether	LOW (LogKOW = 0.2342)
bisphenol A diglycidyl ether resin, solid	LOW (LogKOW = 2.6835)

12.4. Mobility in soil

Ingredient	Mobility
bisphenol A/ diglycidyl ether resin, liquid	LOW (KOC = 51.43)
neopentyl glycol diglycidyl ether	LOW (KOC = 10)
bisphenol A diglycidyl ether resin, solid	LOW (KOC = 51.43)

12.5.Results of PBT and vPvB assessment

	Р	В	т
Relevant available data	Not Available	Not Available	Not Available
PBT Criteria fulfilled?	Not Available	Not Available	Not 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 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
	 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 process equipment to enter drains.
	It may be necessary to collect all wash water for treatment before disposal.
	In all cases disposal to sewer may be subject to local laws and regulations and these should be considered first.
	Where in doubt contact the responsible authority.
	 Recycle wherever possible or consult manufacturer for recycling options.
	Consult State Land Waste Authority for disposal.
	Bury or incinerate residue at an approved site.
	 Recycle containers if possible, or dispose of in an authorised landfill.
Waste treatment options	Not Available
Sewage disposal options	Not Available

SECTION 14 TRANSPORT INFORMATION

Labels Required



Land transport (ADR)

14.1.UN number	3082
14.2.Packing group	II
14.3.UN proper shipping name	ENVIRONMENTALLY HAZARDOUS SUBSTANCE, LIQUID, N.O.S. (contains bisphenol a diglycidyl ether resin, solid)
14.4.Environmental hazard	Not Applicable
14.5. Transport hazard class(es)	Class9SubriskNot Applicable
14.6. Special precautions for user	Hazard identification (Kemler)90Classification codeM6Hazard Label9Special provisions274 335 375 601Limited quantity5 L

Air transport (ICAO-IATA / DGR)

14.4. Environmental hazard Not	I	dous substance, liquid, n.o.s. * (contains	s bisphenol a digly	sidyl ether resin, solid)
14.3. UN proper shipping name Environmental hazard 14.4. Environmental hazard Not 14.5. Transport hazard IC/	nvironmentally hazaro	dous substance, liquid, n.o.s. * (contains	s bisphenol a digly	xidyl ether resin, solid)
14.4. Environmental hazard Not 14.5. Transport hazard IC/	lot Applicable	dous substance, liquid, n.o.s. * (contains	s bisphenol a digly	sidyl ether resin, solid)
14.5. Transport hazard				
14.5. Transport hazard	ICAO/IATA Class			
	ICAO / IATA Subrisk	9 Not Applicable 9L		
14.6. Special precautions for user Pa	Passenger and Cargo		A97 A158 A197 964 450 L 964 450 L Y964	

Passenger and Cargo Limited Maximum Qty / Pack 3

30 kg G

Sea transport (IMDG-Code / GGVSee)

	,	
14.1. UN number	3082	
14.2. Packing group	Ш	
14.3. UN proper shipping name	ENVIRONMENTALL	LY HAZARDOUS SUBSTANCE, LIQUID, N.O.S. (contains bisphenol a diglycidyl ether resin, solid)
14.4. Environmental hazard	Marine Pollutant	
14.5. Transport hazard class(es)	IMDG Class 9 IMDG Subrisk No	ot Applicable
14.6. Special precautions for user	EMS Number Special provisions Limited Quantities	F-A, S-F 274 335 969 5 L

Inland waterways transport (ADN)

14.1. UN number	3082				
14.2. Packing group	Ш				
14.3. UN proper shipping name	ENVIRONMENTALLY HAZARDOUS SUBSTANCE, LIQUID, N.O.S. (contains bisphenol a diglycidyl ether resin, solid)				
14.4. Environmental hazard	Not Applicable				
14.5. Transport hazard class(es)	9 Not Applicable				
14.6. Special precautions for user	Classification codeM6Special provisions274; 335; 375; 601Limited quantity5 LEquipment requiredPPFire cones number0				

Transport in bulk according to Annex II of MARPOL and the IBC code Not Applicable

SECTION 15 REGULATORY INFORMATION

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

BISPHENOL A/ DIGLYCIDYL ETHER RESIN, LIQUID(25085-99-8) IS FOUND ON THE FOLLOWING REGULATORY LISTS

EU European Chemicals Agency (ECHA) Community Rolling Action Plan (CoRAP) List of	European Union (EU) No-Longer Polymers List (NLP) (67/548/EEC)				
Substances	European Union (EU) Regulation (EC) No 1272/2008 on Classification, Labelling and				
EU REACH Regulation (EC) No 1907/2006 - Annex XVII - Restrictions on the manufacture, placing on the market and use of certain dangerous substances, mixtures and articles	Packaging of Substances and Mixtures - Annex VI				
European Union (EU) Annex I to Directive 67/548/EEC on Classification and Labelling of					
Dangerous Substances - updated by ATP: 31					

NEOPENTYL GLYCOL DIGLYCIDYL ETHER(17557-23-2) IS FOUND ON THE FOLLOWING REGULATORY LISTS

EU REACH Regulation (EC) No 1907/2006 - Annex XVII - Restrictions on the manufacture,	European Union (EU) Annex I to Directive 67/548/EEC on Classification and Labelling of
placing on the market and use of certain dangerous substances, mixtures and articles	Dangerous Substances - updated by ATP: 31
European Union - European Inventory of Existing Commercial Chemical Substances (EINECS) (English)	European Union (EU) Regulation (EC) No 1272/2008 on Classification, Labelling and Packaging of Substances and Mixtures - Annex VI

NAPHTHA PETROLEUM, HEAVY ALKYLATE(64741-65-7.) IS FOUND ON THE FOLLOWING REGULATORY LISTS

EU REACH Regulation (EC) No 1907/2006 - Annex XVII (Appendix 2) Carcinogens: category 1B (Table 3.1)/category 2 (Table 3.2)

EU REACH Regulation (EC) No 1907/2006 - Annex XVII (Appendix 4) Mutagens: category 1B (Table 3.1)/category 2 (Table 3.2)

European Union - European Inventory of Existing Commercial Chemical Substances (EINECS) (English)

European Union (EU) Annex I to Directive 67/548/EEC on Classification and Labelling of Dangerous Substances - updated by ATP: 31 European Union (EU) Annex I to Directive 67/548/EEC on Classification and Labelling of Dangerous Substances (updated by ATP: 31) - Carcinogenic Substances European Union (EU) Annex I to Directive 67/548/EEC on Classification and Labelling of Dangerous Substances (updated by ATP: 31) - Mutagenic Substances

European Union (EU) Regulation (EC) No 1272/2008 on Classification, Labelling and Packaging of Substances and Mixtures - Annex VI

BISPHENOL A DIGLYCIDYL ETHER RESIN, SOLID(25068-38-6) IS FOUND ON THE FOLLOWING REGULATORY LISTS

EU European Chemicals Agency (ECHA) Community Rolling Action Plan (CoRAP) List of Substances	European Union (EU) No-Longer Polymers List (NLP) (67/548/EEC)
EU REACH Regulation (EC) No 1907/2006 - Annex XVII - Restrictions on the manufacture, placing on the market and use of certain dangerous substances, mixtures and articles	European Union (EU) Regulation (EC) No 1272/2008 on Classification, Labelling and Packaging of Substances and Mixtures - Annex VI
European Union (EU) Annex I to Directive 67/548/EEC on Classification and Labelling of Dangerous Substances - updated by ATP: 31	
CARBON BLACK(1333-86-4) IS FOUND ON THE FOLLOWING REGULATORY LISTS	
EU REACH Regulation (EC) No 1907/2006 - Annex XVII - Restrictions on the manufacture, placing on the market and use of certain dangerous substances, mixtures and articles	European Union - European Inventory of Existing Commercial Chemical Substances (EINECS) (English)
European Customs Inventory of Chemical Substances ECICS (English) European List of Notified Chemical Substances (ELINCS)	International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Monographs
European Trade Union Confederation (ETUC) Priority List for REACH Authorisation	UK Workplace Exposure Limits (WELs)
(C12-14)ALKYLGLYCIDYL ETHER(68609-97-2) IS FOUND ON THE FOLLOWING REGULA	TORY LISTS
EU European Chemicals Agency (ECHA) Community Rolling Action Plan (CoRAP) List of	European Union - European Inventory of Existing Commercial Chemical Substances (EINECS)

Substances	(English)
EU REACH Regulation (EC) No 1907/2006 - Annex XVII - Restrictions on the manufacture, placing on the market and use of certain dangerous substances, mixtures and articles	European Union (EU) Annex I to Directive 67/548/EEC on Classification and Labelling of Dangerous Substances - updated by ATP: 31
European Customs Inventory of Chemical Substances ECICS (English)	European Union (EU) Regulation (EC) No 1272/2008 on Classification, Labelling and
European Trade Union Confederation (ETUC) Priority List for REACH Authorisation	Packaging of Substances and Mixtures - Annex VI

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/EC, 94/33/EC, 91/689/EEC, 1999/13/EC, Commission Regulation (EU) 2015/830, Regulation (EC) No 1272/2008 and their amendments as well as the following British legislation: - The Control of Substances Hazardous to Health Regulations (COSHH) 2002 - COSHH Essentials - The Management of Health and Safety at Work Regulations 1999

15.2. Chemical safety assessment

For further information please look at the Chemical Safety Assessment and Exposure Scenarios prepared by your Supply Chain if available.

ECHA SUMMARY

Ingredient	CAS number	Index No	ECHA Dossier				
bisphenol A/ diglycidyl ether resin, liquid	25085-99-8	603-074-00-8	01-2119456619-26-XXXX				
Harmonisation (C&L Inventory) Pictograms Signal Hazard Statem Word Code(s) Vorde(s) Code(s)							
1	Skin Irrit. 2, Skin Sens. 1, Eye Irrit. 2, Aquat	GHS07, GHS09, Wng	H315, H317, H319				
2	Skin Irrit. 2, Skin Sens. 1, Eye Irrit. 2, Aqua Chronic 3, Aquatic Chronic 4, Skin Corr. 1/	GHS09, Wng, GHS08, Dgr, GHS07	H315, H317, H319, H372				
1	Skin Irrit. 2, Skin Sens. 1, Eye Irrit. 2, Aquat		GHS07, GHS09, Wng	H315, H317, H319			
2	Skin Irrit. 2, Skin Sens. 1, Eye Irrit. 2, Aquat	Not Classified	GHS07, GHS09, Wng	H315, H317, H319			
1	Skin Irrit. 2, Skin Sens. 1A, Aquatic Chronic		GHS07, GHS09, Wng	H315, H317			
2	Skin Irrit. 2, Skin Sens. 1A, Aquatic Chronic		GHS07, GHS09, Wng	H315, H317			
Harmonisation Code 1 = The most prevalent classification. Harmonisation Code 2 = The most severe classification.							

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Ingredient	CAS number	Ind	Index No		ECHA Dossier	
neopentyl glycol diglycidyl ether	17557-23-2	603	603-094-00-7		lable	
Harmonisation (C&L Inventory)	Hazard Class and Category Code(s)		Pictograms Signal Word Code(s)		Hazard Statement Code(s)	
1	Skin Irrit. 2, Skin Sens. 1		GHS07, Wng		H315, H317	
2	Skin Irrit. 2, Skin Sens. 1		GHS07, Wng		H315, H317	

Harmonisation Code 1 = The most prevalent classification. Harmonisation Code 2 = The most severe classification.

Ingredient	CAS number	Index No	EC	HA Dossier		
naphtha petroleum, heavy alkylate	64741-65-7.	649-275-00-4	01-2	119850115-46-XXXX		
Harmonisation (C&L Inventory)	Hazard Class and Category Code(s)		Pictograms Signal Word Code(s)	Hazard Statement Code(s)		
1	Asp. Tox. 1, Muta. 1B, Carc. 1B			GHS08, Dgr	H304, H340, H350	
2	Asp. Tox. 1, Muta. 1B, Carc. 1B, Flam. Liq. 3, Aquatic Chronic 4, Aquatic Chronic 2, Not Classified, Acute Tox. 3, STOT SE 3, Flam. Liq. 1, Skin Irrit. 2, Flam. Liq. 2, Repr. 2, Aquatic Chronic 3		GHS08, Dgr, GHS02, GHS09, GHS06, Wng	H304, H340, H350, H331, H336, H224, H315, H361		

Harmonisation Code 1 = The most prevalent classification. Harmonisation Code 2 = The most severe classification.

Ingredient	CAS number	Index No	dex No ECHA Dossier		
bisphenol A diglycidyl ether resin, solid	25068-38-6	603-074-00-8	01-2119456619-26-XXXX		
Harmonisation (C&L Inventory)	&L Hazard Class and Category Code(s)		Pictograms Signal Word Code(s)	Hazard Statement Code(s)	
1	Skin Irrit. 2, Skin Sens. 1, Eye Irrit. 2, Aquatic Chronic 2		GHS07, GHS09, Wng	H315, H317, H319	

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832HD Black 1:1 Epoxy Potting and Encapsulating Compound (Part A)

2	Skin Irrit. 2, Skin Sens. 1, Eye Irrit. 2, Aquatic Chronic 2, Not Classified, Acute Tox. 4, Aquatic Chronic 3, Aquatic Chronic 4, Skin Corr. 1A, Aquatic Acute 1, Aquatic Chronic 1	GHS09, Wng, GHS08, Dgr, GHS07	H315, H317, H319, H372
1	Skin Irrit. 2, Skin Sens. 1, Eye Irrit. 2, Aquatic Chronic 2	GHS07, GHS09, Wng	H315, H317, H319
2	Skin Irrit. 2, Skin Sens. 1, Eye Irrit. 2, Aquatic Chronic 2, Skin Sens. 1B, Skin Sens. 1A, Not Classified	GHS07, GHS09, Wng	H315, H317, H319
1	Skin Irrit. 2, Skin Sens. 1A, Aquatic Chronic 2	GHS07, GHS09, Wng	H315, H317
2	Skin Irrit. 2, Skin Sens. 1A, Aquatic Chronic 2	GHS07, GHS09, Wng	H315, H317
Harmonisation Co	de 1 = The most prevalent classification. Harmonisation Code 2 = The most severe classification.	_	1

Ingredient	CAS number	Index No	ECHA Dossier		
carbon black	1333-86-4	Not Available	01-2119384822-32-XXXX, 01-2119489801-30-XXXX, 01-2119475601-40-XXXX		
Harmonisation (C&L Inventory)	Hazard Class and Ca	tegory Code(s)		Pictograms Signal Word Code(s)	Hazard Statement Code(s)
1	Not Classified	Not Classified		GHS08, Wng, Dgr, GHS06, GHS02, GHS09	H351, H335, H319, H372, H251, H315, H228, H370, H332
2	,	Not Classified, Carc. 2, STOT SE 3, Eye Irrit. 2, STOT RE 2, STOT RE 1, Aquatic Chronic 4, Self-heat. 1, Self-heat. 2, Skin Irrit. 2, STOT SE 1, Aquatic Chronic 1, Flam. Sol. 2, Acute Tox. 4		GHS08, Wng, Dgr, GHS06, GHS02, GHS09	H351, H335, H319, H372, H251, H315, H228, H370, H332
2	,	ssified, Carc. 2, STOT SE 3, Eye Irrit. 2, STOT RE 2, STOT RE 1, Aquatic : 4, Self-heat. 1, Self-heat. 2, Skin Irrit. 2, STOT SE 1, Aquatic Chronic 1, Flam. Acute Tox. 4		GHS08, Wng, Dgr, GHS06, GHS02, GHS09	H351, H335, H319, H372, H251, H315, H228, H370, H332

Harmonisation Code 1 = The most prevalent classification. Harmonisation Code 2 = The most severe classification.

Ingredient	CAS number	Index No	ECHA Dossier	
(C12-14)alkylglycidyl ether	68609-97-2 603-103-00-4		01-2119485289-22-XXXX	
Harmonisation (C&L Inventory)	Hazard Class and Category Code(s)		Pictograms Signal Word Code(s)	Hazard Statement Code(s)
1	Skin Irrit. 2, Skin Sens. 1		GHS07, Wng	H315, H317
2	Skin Irrit. 2, Skin Sens. 1, Aquatic Chronic 2, Not Classified, Acute Tox. 4, Eye Irrit. 2		GHS07, Wng, GHS09	H315, H317

Harmonisation Code 1 = The most prevalent classification. Harmonisation Code 2 = The most severe classification.

National Inventory	Status	
Australia - AICS	Y	
Canada - DSL	Y	
Canada - NDSL	N (bisphenol A diglycidyl ether resin, solid; (C12-14)alkylglycidyl ether; bisphenol A/ diglycidyl ether resin, liquid; neopentyl glycol diglycidyl ether; naphtha petroleum, heavy alkylate; carbon black)	
China - IECSC	Υ	
Europe - EINEC / ELINCS / NLP	Y	
Japan - ENCS	N (bisphenol A diglycidyl ether resin, solid; (C12-14)alkylglycidyl ether; bisphenol A/ diglycidyl ether resin, liquid; neopentyl glycol diglycidyl ether; naphtha petroleum, heavy alkylate)	
Korea - KECI	Y	
New Zealand - NZIoC	Υ	
Philippines - PICCS	Y	
USA - TSCA	Y	
Legend:	Y = All ingredients are on the inventory $N = Not$ determined or one or more ingredients are not on the inventory and are not exempt from listing(see specific ingredients in brackets)	

SECTION 16 OTHER INFORMATION

Full text Risk and Hazard codes

H224	Extremely flammable liquid and vapour.	
H226	Flammable liquid and vapour.	
H228	Flammable solid.	
H251	Self-heating: may catch fire.	
H304	May be fatal if swallowed and enters airways.	
H331	Toxic if inhaled.	
H332	Harmful if inhaled.	
H335	May cause respiratory irritation.	
H336	May cause drowsiness or dizziness.	
H340	May cause genetic defects.	
H350	May cause cancer.	
H351	Suspected of causing cancer.	
H361	Suspected of damaging fertility or the unborn child.	

H370	Causes damage to organs.
H372	Causes damage to organs.

Other information

Ingredients with multiple cas numbers

Name	CAS No
bisphenol A/ diglycidyl ether resin, liquid	25068-38-6, 25085-99-8
bisphenol A diglycidyl ether resin, solid	25068-38-6, 25085-99-8

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.

A list of reference resources used to assist the committee may be found at: www.chemwatch.net

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.

For detailed advice on Personal Protective Equipment, refer to the following EU CEN Standards:

- EN 166 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

Definitions and abbreviations

PC-TWA: Permissible Concentration-Time Weighted Average

 $\label{eq:product} \mathsf{PC-STEL:}\ \mathsf{Permissible}\ \mathsf{Concentration}\text{-}\mathsf{Short}\ \mathsf{Term}\ \mathsf{Exposure}\ \mathsf{Limit}$

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