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DRACO ITALIANA S.p.A. DRAP118 - PRIMER ES 40 COMP. A

Revision nr.3 Dated 08/06/2021 Printed on 08/06/2021 Page n. 1 / 21

Replaced revision:2 (Dated 06/10/2020)

Safety Data Sheet

According to Annex II to REACH - Regulation 2015/830

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

1.1. Product identifier

Code: DRAP118

Product name PRIMER ES 40 COMP. A

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

Intended use Epoxy primer two-component

1.3. Details of the supplier of the safety data sheet

Name DRACO ITALIANA S.p.A. Full address Via Monte Grappa, 11 D-E

District and Country 20067 Tribiano (MI)

Italia

Tel. +39 02.90632917 Fax +39 02.90631976

e-mail address of the competent person responsible for the Safety Data Sheet

info@draco-edilizia.it

1.4. Emergency telephone number

For urgent inquiries refer to Centro Antiveleni di Bergamo 800883300 (Azienda Ospedaliera Papa Giovanni XXII)

Centro Antiveleni di Firenze 0557947819 (Az. Osp. "Careggi" U.O. Tossicologia

Medica

Centro Antiveleni di Foggia 80018345 (Az. Osp. Univ. Foggia) Centro Antiveleni di Milano 0266101029 (Osp. Niguarda Ca' Granda) Centro Antiveleni di Napoli 0817472870 (Az. Osp. "A. Cardarelli")

Centro Antiveleni di Pavia 038224444 (CAV Centro Nazionale di Informazione

Tossicologica)

Centro Antiveleni di Roma 063054343 (CAV Policlinico "A. Gemelli") Centro Antiveleni di Roma 0649978000 (CAV Policlinico "Umberto I")

Centro Antiveleni di Roma 06 68593726 (CAV "Osp. Pediatrico Bambino Gesù" Dip.

Emergenza e Accettazione DEA)

SECTION 2. Hazards identification

2.1. Classification of the substance or mixture

The product is classified as hazardous pursuant to the provisions set forth in (EC) Regulation 1272/2008 (CLP) (and subsequent amendments and supplements). The product thus requires a safety datasheet that complies with the provisions of (EU) Regulation 2015/830. Any additional information concerning the risks for health and/or the environment are given in sections 11 and 12 of this sheet.

Hazard classification and indication:

Flammable liquid, category 2 H225 Highly flammable liquid and vapour. Eye irritation, category 2 H319 Causes serious eye irritation. Skin irritation, category 2 H315 Causes skin irritation. Skin sensitization, category 1 H317 May cause an allergic skin reaction. Specific target organ toxicity - single exposure, H336 May cause drowsiness or dizziness. category 3

Hazardous to the aquatic environment, chronic H411 Toxic to aquatic life with long lasting effects.

toxicity, category 2

2.2. Label elements

Hazard labelling pursuant to EC Regulation 1272/2008 (CLP) and subsequent amendments and supplements.

Hazard pictograms:



DRAP118 - PRIMER ES 40 COMP. A

Revision nr.3 Dated 08/06/2021 Printed on 08/06/2021 Page n. 2 / 21

Replaced revision:2 (Dated 06/10/2020)

SECTION 2. Hazards identification .../>>

Signal words: Danger

Hazard statements:

H225 Highly flammable liquid and vapour.
H319 Causes serious eye irritation.
H315 Causes skin irritation.
H317 May cause an allergic skin reaction.

H317 May cause an allergic skin reaction.
 H336 May cause drowsiness or dizziness.
 H411 Toxic to aquatic life with long lasting effects.

Precautionary statements:

P210 Keep away from heat, hot surfaces, sparks, open flames and other ignition sources. No smoking.

P280 Wear protective gloves/ protective clothing / eye protection / face protection.

P370+P378 In case of fire: use . . . to extinguish.
P273 Avoid release to the environment.

P391 Collect spillage.

P261 Avoid breathing dust / fume / gas / mist / vapours / spray.

Contains: Reaction product: bisphenol-A-epichlorohydrin and epoxy resins (average molecular weight <= 700)

Reaction product: bisphenol-F-epichlorohydrin and epoxy resins (average molecular weight <= 700)

Oxirane, mono [(C12-14- alkyloxy) methyl] derivatives

Acetone

2.3. Other hazards

On the basis of available data, the product does not contain any PBT or vPvB in percentage ≥ than 0,1%.

SECTION 3. Composition/information on ingredients

Xylene, mixture of isomers

XYLENE (MIXTURE OF ISOMERS)

3.2. Mixtures

Contains:

Identification x = Conc. % Classification 1272/2008 (CLP)

Acetone

CAS 67-64-1 30 ≤ x < 50 Flam. Liq. 2 H225, Eye Irrit. 2 H319, STOT SE 3 H336, EUH066

EC 200-662-2 INDEX 606-001-00-8

Reg. no. 01-2119471330-49-XXXX

Reaction product: bisphenol-A-epichlorohydrin and epoxy resins (average molecular weight <= 700)

CAS 25068-38-6 25 ≤ x < 30 Eye Irrit. 2 H319, Skin Irrit. 2 H315, Skin Sens. 1 H317, Aquatic Chronic 2 H411

EC 500-033-5 INDEX 603-074-00-8

Reg. no. 01-2119456619-26-XXXX

Reaction product: bisphenol-F-epichlorohydrin and epoxy resins (average molecular weight <= 700)

CAS 9003-36-5 10 ≤ x < 25 Skin Irrit. 2 H315, Skin Sens. 1 H317, Aquatic Chronic 2 H411

EC 500-006-8

INDEX

Reg. no. 01-2119454392-40-XXXX

Oxirane, mono [(C12-14- alkyloxy) methyl] derivatives

CAS 68609-97-2 $5 \le x < 9$ Skin Irrit. 2 H315, Skin Sens. 1 H317

EC 271-846-8 INDEX 603-103-00-4

Reg. no. 01 2119485289-22-XXXX

Xylene, mixture of isomers

CAS 1330-20-7 1 ≤ x < 5 Flam. Liq. 3 H226, Acute Tox. 4 H312, Acute Tox. 4 H332, Asp. Tox. 1 H304,

Eye Irrit. 2 H319, Skin Irrit. 2 H315, STOT SE 3 H335, Aquatic Chronic 3 H412

EC 215-535-7

INDEX 601-022-00-9

Reg. no. 01-2119488216-32-XXXX

DRAP118 - PRIMER ES 40 COMP. A

Revision nr.3 Dated 08/06/2021 Printed on 08/06/2021 Page n. 3 / 21 Replaced revision:2 (Dated 06/10/2020)

SECTION 3. Composition/information on ingredients .../>>

2-BUTOXYETHANOL

CAS 111-76-2 1≤x<5 Acute Tox. 4 H302, Acute Tox. 4 H312, Acute Tox. 4 H332, Eye Irrit. 2 H319,

Skin Irrit, 2 H315

EC 203-905-0 INDEX 603-014-00-0

ETHYLBENZENE

CAS 100-41-4 $1 \le x < 5$ Flam. Liq. 2 H225, Acute Tox. 4 H332, Asp. Tox. 1 H304, STOT RE 2 H373,

Aquatic Chronic 3 H412

EC 202-849-4 INDEX 601-023-00-4

Reg. no. 01-2119489370-35-XXXX

The full wording of hazard (H) phrases is given in section 16 of the sheet.

SECTION 4. First aid measures

4.1. Description of first aid measures

EYES: Remove contact lenses, if present. Wash immediately with plenty of water for at least 15 minutes, opening the eyelids fully. If problem persists, seek medical advice.

SKIN: Remove contaminated clothing. Rinse skin with a shower immediately. Get medical advice/attention immediately. Wash contaminated clothing before using it again.

INHALATION: Remove to open air. If the subject stops breathing, administer artificial respiration. Get medical advice/attention immediately. INGESTION: Get medical advice/attention immediately. Do not induce vomiting. Do not administer anything not explicitly authorised by a doctor

4.2. Most important symptoms and effects, both acute and delayed

Specific information on symptoms and effects caused by the product are unknown.

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

Information not available

SECTION 5. Firefighting measures

5.1. Extinguishing media

SUITABLE EXTINGUISHING EQUIPMENT

Extinguishing substances are: carbon dioxide, foam, chemical powder. For product loss or leakage that has not caught fire, water spray can be used to disperse flammable vapours and protect those trying to stem the leak.

UNSUITABLE EXTINGUISHING EQUIPMENT

Do not use jets of water. Water is not effective for putting out fires but can be used to cool containers exposed to flames to prevent explosions.

5.2. Special hazards arising from the substance or mixture

HAZARDS CAUSED BY EXPOSURE IN THE EVENT OF FIRE

Excess pressure may form in containers exposed to fire at a risk of explosion. Do not breathe combustion products.

5.3. Advice for firefighters

GENERAL INFORMATION

Use jets of water to cool the containers to prevent product decomposition and the development of substances potentially hazardous for health. Always wear full fire prevention gear. Collect extinguishing water to prevent it from draining into the sewer system. Dispose of contaminated water used for extinction and the remains of the fire according to applicable regulations.

SPECIAL PROTECTIVE EQUIPMENT FOR FIRE-FIGHTERS

Normal fire fighting clothing i.e. fire kit (BS EN 469), gloves (BS EN 659) and boots (HO specification A29 and A30) in combination with self-contained open circuit positive pressure compressed air breathing apparatus (BS EN 137).

SECTION 6. Accidental release measures

6.1. Personal precautions, protective equipment and emergency procedures

Block the leakage if there is no hazard.

Wear suitable protective equipment (including personal protective equipment referred to under Section 8 of the safety data sheet) to prevent any contamination of skin, eyes and personal clothing. These indications apply for both processing staff and those involved in emergency procedures.

Send away individuals who are not suitably equipped. Use explosion-proof equipment. Eliminate all sources of ignition (cigarettes, flames,

Revision nr.3 Dated 08/06/2021 Printed on 08/06/2021 Page n. 4 / 21

Replaced revision:2 (Dated 06/10/2020)

SECTION 6. Accidental release measures .../>>

sparks, etc.) from the leakage site.

6.2. Environmental precautions

The product must not penetrate into the sewer system or come into contact with surface water or ground water.

6.3. Methods and material for containment and cleaning up

Collect the leaked product into a suitable container. Evaluate the compatibility of the container to be used, by checking section 10. Absorb the remainder with inert absorbent material.

Make sure the leakage site is well aired. Contaminated material should be disposed of in compliance with the provisions set forth in point 13.

6.4. Reference to other sections

Any information on personal protection and disposal is given in sections 8 and 13.

SECTION 7. Handling and storage

7.1. Precautions for safe handling

Keep away from heat, sparks and naked flames; do not smoke or use matches or lighters. Without adequate ventilation, vapours may accumulate at ground level and, if ignited, catch fire even at a distance, with the danger of backfire. Avoid bunching of electrostatic charges. When performing transfer operations involving large containers, connect to an earthing system and wear antistatic footwear. Vigorous stirring and flow through the tubes and equipment may cause the formation and accumulation of electrostatic charges. In order to avoid the risk of fires and explosions, never use compressed air when handling. Open containers with caution as they may be pressurised. Do not eat, drink or smoke during use. Avoid leakage of the product into the environment.

7.2. Conditions for safe storage, including any incompatibilities

Store only in the original container. Store the containers sealed, in a well ventilated place, away from direct sunlight. Store in a cool and well ventilated place, keep far away from sources of heat, naked flames and sparks and other sources of ignition. Keep containers away from any incompatible materials, see section 10 for details.

7.3. Specific end use(s)

Information not available

SECTION 8. Exposure controls/personal protection

8.1. Control parameters

Regulatory References:

FRA France Valeurs limites d'exposition professionnelle aux agents chimiques en France. ED 984 - INRS

ITA Italia Decreto Legislativo 9 Aprile 2008, n.81

GBR United Kingdom EH40/2005 Workplace exposure limits (Fourth Edition 2020)

EU OEL EU Directive (EU) 2019/1831; Directive (EU) 2019/130; Directive (EU) 2019/983; Directive (EU)

2017/2398; Directive (EU) 2017/164; Directive 2009/161/EU; Directive 2006/15/EC; Directive

2004/37/EC; Directive 2000/39/EC; Directive 98/24/EC; Directive 91/322/EEC.

TLV-ACGIH ACGIH 2020

Revision nr.3 Dated 08/06/2021 Printed on 08/06/2021 Page n. 5 / 21 Replaced revision:2 (Dated 06/10/2020)

SECTION 8. Exposure controls/personal protection	/ >>
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				Ad	cetone				
Threshold Limit Va	lue								
Туре	Country	untry TWA/8h			STEL/15min Remarks / O				
,,		mg/m3	ppm	mg/m3	ppm				
VLEP	FRA	1210	500	2420	1000				
VLEP	ITA	1210	500						
WEL	GBR	1210	500	3620	1500				
OEL	EU	1210	500						
TLV-ACGIH		250	594	1187	500		irr oclr, TR	S, ssnc	
redicted no-effect	t concentra	ation - PNEC	;						
Normal value in f	resh water						10,6	mg/l	
Normal value in r	marine wate	er					1,06	mg/l	
Normal value for	fresh wate	r sediment					30,4	mg/kg/d	
Normal value for	marine wa	ter sediment					3,04	mg/kg/d	
Normal value for	water, inte	rmittent relea	ase				21	mg/l	
Normal value of	STP micro	organisms					100	mg/l	
Normal value for	the terresti	rial compartn	nent				29,5	mg/kg/d	
lealth - Derived no	effect lev	el - DNEL / I	DMEL						
	Effe	cts on consu	mers			Effects on worke	ers		
Route of exposur	re Acu	te Acu	ite	Chronic	Chronic	Acute local	Acute	Chronic	Chronic
	loca	ıl sys	temic	local	systemic		systemic	local	systemic
Oral							62		
							mg/kg		
							bw/d		
Inhalation					200	2420			1210
					mg/m3	mg/m3			mg/m3
Skin					62				186
					mg/kg bw/d				mg/kg
									bw/d

Reacti	on product	: bisphenol-A-epi	ichlorohydrin	and epoxy resi	ns (average mo	olecular weig	ht <= 700)	
redicted no-effect cor	ncentration	- PNEC	-		,	_		
Normal value in fresh	n water					3	μg/l	
Normal value in mari	ne water					0,3	μg/l	
Normal value for fres	h water sed	iment				0,5	mg/kg/d	
Normal value for mar	ine water se	ediment				0,5	mg/kg/d	
Normal value for wat	er, intermitte	ent release				0,013	mg/l	
Normal value of STP	microorgan	isms				10	mg/l	
Normal value for the	0,05	mg/kg/d						
lealth - Derived no-eff	ect level - D	ONEL / DMEL						
	Effects o	n consumers	kers					
Route of exposure	Acute	Acute	Chronic	Chronic	Acute local	Acute	Chronic	Chronic
	local	systemic	local	systemic		systemic	local	systemic
Oral		0,75		0,75				
		mg/kg bw/d		mg/kg bw/d				
Inhalation		0,75		0,75		12,3		12,3
		mg/m3		mg/m3		mg/m3		mg/m3
Skin		3,6		3,6	8,3	8,3		8,3
		mg/kg bw/d		mg/kg bw/d		mg/kg		mg/kg
		-		- -		bw/d		bw/d

Revision nr.3 Dated 08/06/2021 Printed on 08/06/2021 Page n. 6 / 21 Replaced revision:2 (Dated 06/10/2020)

bw/d

Reaction product: bisphenol-F-epichlorohydrin and epoxy resins (average molecular weight <= 700) Predicted no-effect concentration - PNEC Normal value in fresh water 0,003 mg/l Normal value in marine water 0,0003 mg/l Normal value for fresh water sediment 0,294 mg/kg/d Normal value for marine water sediment 0,0294 mg/kg/d Normal value for water, intermittent release 0,0254 mg/l Normal value of STP microorganisms 10 mg/l Normal value for the terrestrial compartment 0,237 mg/kg/d Health - Derived no-effect level - DNEL / DMEL	
Predicted no-effect concentration - PNEC Normal value in fresh water 0,003 mg/l Normal value in marine water 0,0003 mg/l Normal value for fresh water sediment 0,294 mg/kg/d Normal value for marine water sediment 0,0294 mg/kg/d Normal value for water, intermittent release 0,0254 mg/l Normal value of STP microorganisms 10 mg/l Normal value for the terrestrial compartment 0,237 mg/kg/d Health - Derived no-effect level - DNEL / DMEL	
Normal value in fresh water 0,003 mg/l Normal value in marine water 0,0003 mg/l Normal value for fresh water sediment 0,294 mg/kg/d Normal value for marine water sediment 0,0294 mg/kg/d Normal value for water, intermittent release 0,0254 mg/l Normal value of STP microorganisms 10 mg/l Normal value for the terrestrial compartment 0,237 mg/kg/d Health - Derived no-effect level - DNEL / DMEL	
Normal value in marine water Normal value for fresh water sediment Normal value for marine water sediment Normal value for marine water sediment Normal value for water, intermittent release Normal value of STP microorganisms Normal value for the terrestrial compartment Normal value for the terrestrial compartment O,237 mg/kg/d	
Normal value for fresh water sediment Normal value for marine water sediment Normal value for marine water sediment Normal value for water, intermittent release Normal value of STP microorganisms Normal value for the terrestrial compartment Normal value for the terrestrial compartment O,237 mg/kg/d Health - Derived no-effect level - DNEL / DMEL	
Normal value for marine water sediment 0,0294 mg/kg/d Normal value for water, intermittent release 0,0254 mg/l Normal value of STP microorganisms 10 mg/l Normal value for the terrestrial compartment 0,237 mg/kg/d Health - Derived no-effect level - DNEL / DMEL	
Normal value for water, intermittent release 0,0254 mg/l Normal value of STP microorganisms 10 mg/l Normal value for the terrestrial compartment 0,237 mg/kg/d Health - Derived no-effect level - DNEL / DMEL	
Normal value of STP microorganisms Normal value for the terrestrial compartment 0,237 mg/kg/d Health - Derived no-effect level - DNEL / DMEL	
Normal value for the terrestrial compartment 0,237 mg/kg/d Health - Derived no-effect level - DNEL / DMEL	
Health - Derived no-effect level - DNEL / DMEL	
Effects on consumers Effects on workers	
Route of exposure Acute Acute Chronic Chronic Acute local Acute Chronic Chronic	iic
local systemic local systemic systemic local systemi	nic
Oral 6,25	
mg/kg bw/d	
Inhalation 8,7 29,39	
mg/m3 mg/m3	3
Skin 62,5 8,3 104,15	5
mg/kg bw/d μg/cm2 mg/kg	

				Xylene, mix	ture of isome	ers			
hreshold Limit V	/alue								
Туре	Country TWA/8h			STEL/15	min	Remarks / O	bservations		
		mg/m3	ppm	mg/m3	ppm				
VLEP	FRA	221	50	442	100	SKIN			
VLEP	ITA	221	50	442	100				
WEL	GBR	220	50	441	100	SKIN			
OEL	EU	221	50	442	100	SKIN			
TLV-ACGIH		434	100	651	150				
redicted no-effe	ct concentra	ation - PNE	3						
Normal value in	r fresh water						0,327	mg/l	
Normal value in	n marine wate	er					0,327	mg/l	
Normal value for	or fresh wate	r sediment					12,46	mg/kg	
Normal value for	or marine wa	ter sediment					12,46	mg/kg	
Normal value for	or water, inte	rmittent relea	ase				0,327	mg/l	
Normal value of	f STP microo	organisms					6,58	mg/l	
Normal value for	or the terresti	rial compartr	nent				2,31	mg/kg	
lealth - Derived r	no-effect lev	el - DNEL /	DMEL						
	Effe	cts on consu	ımers			Effects on workers			
Route of expos	ure Acu	te Acı	ıte	Chronic	Chronic	Acute local	Acute	Chronic	Chronic
	loca	ıl sys	temic	local	systemic		systemic	local	systemic
Inhalation		•			•	442 mg/m3	442 mg/m3	221 mg/m3	221 mg/m3

Revision nr.3 Dated 08/06/2021 Printed on 08/06/2021 Page n. 7 / 21 Replaced revision:2 (Dated 06/10/2020)

SECTION 8. Exposure controls/personal protection .../>>

•				 1100	DE1175115				
				ETHYL	BENZENE				
hreshold Limit V									
Туре	Country	TWA/8h		STEL/15r	nin	Remarks / Ol	oservations		
		mg/m3	ppm	mg/m3	ppm				
VLEP	FRA	88,4	20	442	100	SKIN			
VLEP	ITA	442	100	884	200	SKIN			
WEL	GBR	441	100	552	125	SKIN			
OEL	EU	442	100	884	200	SKIN			
TLV-ACGIH		87	20						
redicted no-effe	ct concentra	ation - PNE	C						
Normal value in	fresh water						0,1	mg/l	
Normal value in	marine water	er					0,01	mg/l	
Normal value for	r fresh wate	r sediment					13,7	mg/kg	
Normal value for	r marine wa	ter sediment					1,37	mg/kg	
Normal value for	r water, inte	rmittent rele	ase				0,1	mg/l	
Normal value of	STP micro	organisms					9,6	mg/l	
Normal value for	r the terresti	rial compartr	nent				2,68	mg/kg	
lealth - Derived n	o-effect lev	el - DNEL /	DMEL						
	Effe	cts on consu	ımers			Effects on workers			
Route of exposu	ure Acu	te Acı	ute	Chronic	Chronic	Acute local	Acute	Chronic	Chronic
	loca	ıl sys	temic	local	systemic		systemic	local	systemic
Oral		_			1,6		•		•
					mg/kg bw/d				
Inhalation					15	293			77
					mg/m3	mg/m3			mg/m3
Skin									180
									mg/kg/d

				2-BUTO	XYETHANOL				
hreshold Limit Val	lue								
Type Coun		TWA/8h		STEL/15	min	Remarks / Ob	servations		
		mg/m3	ppm	mg/m3	ppm				
VLEP	FRA	49	10	246	50	SKIN			
VLEP	ITA	98	20	246	50	SKIN			
WEL	GBR	123	25	246	50	SKIN			
OEL	EU	98	20	246	50	SKIN			
TLV-ACGIH		97	20						
Predicted no-effect	concentra	ation - PNE	С						
Normal value in fr						8,8	mg/l		
Normal value in n	narine wate	er					0,88	mg/l	
Normal value for	fresh wate	sediment					34,6	mg/kg	
Normal value for	marine wa	ter sedimen	t				3,46	mg/kg	
Normal value for	water, inte	mittent rele	ase				9,1	mg/l	
Normal value of S	STP microc	rganisms					463	mg/l	
Normal value for	the terresti	ial compartı	ment				2,33	mg/kg/d	
lealth - Derived no	-effect lev	el - DNEL /	DMEL						
	Effe	cts on consu	umers			Effects on workers			
Route of exposure	e Acu	te Acı	ute	Chronic	Chronic	Acute local	Acute	Chronic	Chronic
	loca	l sys	stemic	local	systemic		systemic	local	systemic
Oral		26,	7		6,3				
		mg	/kg bw/d		mg/kg bw/d				
Inhalation	147	426	3	147	59	246	1091		98
	mg/	m3 mg	/m3	mg/m3	mg/m3	mg/m3	mg/m3		mg/m3
Skin		89			75		89		125
		mg	/kg bw/d		mg/kg bw/d		mg/kg		mg/kg
							bw/d		bw/d

Legend

(C) = CEILING ; INHAL = Inhalable Fraction ; RESP = Respirable Fraction ; THORA = Thoracic Fraction.

VND = hazard identified but no DNEL/PNEC available ; NEA = no exposure expected ; NPI = no hazard identified.

Acetone

Biological index of exposure:

Components with biological limit values: CAS: 67-64-1 acetone

IBE (ACGIH 2019) 25 mg / I

Samples: urine

Time of withdrawal: at the end of the shift

Biological indicator: acetone

Notes: the biological indicator is not specific, since it is also possible to detect its presence after exposure to other chemicals.

DRAP118 - PRIMER ES 40 COMP. A

Printed on 08/06/2021
Printed on 08/06/2021
Page n. 8 / 21
Replaced revision:2 (Dated 06/10/2020)

Revision nr.3

SECTION 8. Exposure controls/personal protection .../>>

8.2. Exposure controls

As the use of adequate technical equipment must always take priority over personal protective equipment, make sure that the workplace is well aired through effective local aspiration.

When choosing personal protective equipment, ask your chemical substance supplier for advice.

Personal protective equipment must be CE marked, showing that it complies with applicable standards.

Provide an emergency shower with face and eye wash station.

HAND PROTECTION

Protect hands with category III work gloves (see standard EN 374).

The following should be considered when choosing work glove material: compatibility, degradation, failure time and permeability.

The work gloves' resistance to chemical agents should be checked before use, as it can be unpredictable. The gloves' wear time depends on the duration and type of use.

SKIN PROTECTION

Wear category II professional long-sleeved overalls and safety footwear (see Regulation 2016/425 and standard EN ISO 20344). Wash body with soap and water after removing protective clothing.

Consider the appropriateness of providing antistatic clothing in the case of working environments in which there is a risk of explosion.

EYE PROTECTION

Wear airtight protective goggles (see standard EN 166).

RESPIRATORY PROTECTION

If the threshold value (e.g. TLV-TWA) is exceeded for the substance or one of the substances present in the product, wear a mask with a type AX filter, whose limit of use will be defined by the manufacturer (see standard EN 14387). In the presence of gases or vapours of various kinds and/or gases or vapours containing particulate (aerosol sprays, fumes, mists, etc.) combined filters are required.

Respiratory protection devices must be used if the technical measures adopted are not suitable for restricting the worker's exposure to the threshold values considered. The protection provided by masks is in any case limited.

If the substance considered is odourless or its olfactory threshold is higher than the corresponding TLV-TWA and in the case of an emergency, wear open-circuit compressed air breathing apparatus (in compliance with standard EN 137) or external air-intake breathing apparatus (in compliance with standard EN 138). For a correct choice of respiratory protection device, see standard EN 529.

ENVIRONMENTAL EXPOSURE CONTROLS

The emissions generated by manufacturing processes, including those generated by ventilation equipment, should be checked to ensure compliance with environmental standards.

Product residues must not be indiscriminately disposed of with waste water or by dumping in waterways.

Acetone

Respiratory protection:

for short exposures or in the event of an accident: filter devices, type AX (EN 371). Having a breathing apparatus that does not depend on circulating air ready for emergencies.

Hand protection:

protective gloves compliant with EN 374.

Glove material: butyl rubber (butyl rubber) - layer thickness> = 0.5 mm.

Breakthrough time:> 480 min.

Observe the glove manufacturer's instructions regarding penetrability and breakthrough time.

Eye protection:

hermetically sealed safety goggles according to EN 166.

Body protection:

use solvent resistant protective clothing.

Recommendation:

flame retardant, antistatic protective clothing. safety shoes according to EN 345-347.

General protection and hygiene measures

Wash hands before breaks and after work. Avoid contact with skin and eyes. Do not eat, drink or smoke during use. Have an eye wash bottle or eye rinse ready at work.

Alternatives to the following personal protective measures can only be determined in consultation with a responsible safety expert.

Xylene, mixture of isomers

XYLENE (MIXTURE OF ISOMERS)

Protect your hands with nitrile latex gloves compliant with EN 374-1: 2016.

SECTION 9. Physical and chemical properties

9.1. Information on basic physical and chemical properties

PropertiesValueInformationAppearanceliquidColouryellow

DRAP118 - PRIMER ES 40 COMP. A

Revision nr.3 Dated 08/06/2021 Printed on 08/06/2021 Page n. 9 / 21

Replaced revision:2 (Dated 06/10/2020)

SECTION 9. Physical and chemical properties .../>>

solvent Not available Odour threshold Not available pН Melting point / freezing point Not available Initial boiling point °C 56 Not available Boiling range °C Flash point 23 Evaporation rate Not available Flammability (solid, gas) Not available Lower inflammability limit Not available Upper inflammability limit Not available Not available Lower explosive limit Upper explosive limit Not available Vapour pressure Not available Vapour density Not available Relative density 1,05 g/cc Not available Solubility Partition coefficient: n-octanol/water Not available Not available Auto-ignition temperature Decomposition temperature Not available Not available Viscosity Not available Explosive properties Oxidising properties Not available

9.2. Other information

VOC (Directive 2010/75/EC) : 50,00 % - 525,00 g/litre VOC (volatile carbon) : 31,97 % - 335,63 g/litre

SECTION 10. Stability and reactivity

10.1. Reactivity

There are no particular risks of reaction with other substances in normal conditions of use.

2-BUTOXYETHANOL

Decomposes under the effect of heat.

10.2. Chemical stability

The product is stable in normal conditions of use and storage.

10.3. Possibility of hazardous reactions

The vapours may also form explosive mixtures with the air.

Acetone

Risk of explosion on contact with: bromine trifluoride, fluorine dioxide, hydrogen peroxide, nitrosyl chloride, 2-methyl-1,3-butadiene, nitromethane, nitrosyl perchlorate. May react dangerously with: potassium tert-butoxide, alkaline hydroxides, bromine, bromoform, isoprene, sodium, sulfur dioxide, chromium trioxide, cromyl chloride, nitric acid, chloroform, peroxymonosulfuric acid, phosphorus oxychloride, chromosulfuric acid, fluorine, strong oxidizing agents, strong reducing agents. Develop flammable gases in contact with: nitrosyl perchlorate.

Xylene, mixture of isomers

XYLENE (MIXTURE OF ISOMERS)

Stable under normal conditions of use and storage Reacts violently with: strong oxidants, strong acids, acid nitric, perchlorates.May form explosive mixtures with: air.

ETHYLBENZENE

Reacts violently with: strong oxidants. Attacks various types of plastic materials. May form explosive mixtures with: air. Reacts violently with: strong oxidants Attacks various types of plastics May form explosive mixtures with: air.

2-BUTOXYETHANOL

May react dangerously with: aluminium,oxidising agents.Forms peroxides with: air. May react dangerously with: aluminum, oxidizing agents: Forms peroxides with: air.

10.4. Conditions to avoid

Avoid overheating. Avoid bunching of electrostatic charges. Avoid all sources of ignition.

Acetone

Dated 08/06/2021 Printed on 08/06/2021 Page n. 10 / 21

Revision nr.3

Replaced revision:2 (Dated 06/10/2020)

SECTION 10. Stability and reactivity .../>>

Avoid exposure to: heat sources, open flames.

2-BUTOXYETHANOL

Avoid exposure to: sources of heat,naked flames. Avoid exposure to: heat sources, open flames.

10.5. Incompatible materials

Acetone

Incompatible with: acids, oxidizing substances.

10.6. Hazardous decomposition products

In the event of thermal decomposition or fire, gases and vapours that are potentially dangerous to health may be released.

Acetone

It can develop: ketene, irritants.

ETHYLBENZENE

May develop: methane, styrene, hydrogen, ethane. It can develop: methane, styrene, hydrogen, ethane.

2-BUTOXYETHANOL

May develop: hydrogen. It can develop hydrogen.

SECTION 11. Toxicological information

In the absence of experimental data for the product itself, health hazards are evaluated according to the properties of the substances it contains, using the criteria specified in the applicable regulation for classification.

It is therefore necessary to take into account the concentration of the individual hazardous substances indicated in section 3, to evaluate the toxicological effects of exposure to the product.

11.1. Information on toxicological effects

Oxirane, mono [(C12-14- alkyloxy) methyl] derivatives

Inhalation: May cause respiratory tract irritation.

Ingestion: Gastrointestinal symptoms, including stomach pain.

Skin contact: Irritating to skin. May cause sensitization by skin contact.

Contact with eyes: Irritating to eyes.

Metabolism, toxicokinetics, mechanism of action and other information

Acetone

Acetone appears in the human and mammalian organisms as an endogenous product of normal metabolism with considerably increased levels during altered physiological states.

Acetone from dermal, inhaled and oral exposure is rapidly absorbed. Relative airway absorption was approximately 50% in humans. It passes into the blood within a few minutes. Acetone is not selectively absorbed into any tissue but is more evenly distributed in body water

The metabolic fate of exogenous acetone is independent of the pathway of absorption and involves three separate low-dose gluconeogenic pathways with acetol (1-hydroxyacetone), methylglyoxal and 1,2-propanediol as intermediates. Both methylglyoxal and propanediol are oxidized to pyruvate, which is the basic component for the biosynthesis of many endogenous biochemicals. At high doses, an alternating metabolic pathway appears with cleavage of 1,2-propanediol to acetate and formate. The elimination of acetone is effective even at high internal doses and occurs through metabolic transformation to endogenous biochemical substances, such as acetone vapor through the airways and skin surface, through the exhalation of CO2 and into the urine as acetone or acetol, methylglyoxal or as D-lactoyl-GSH. The acetone turnover rates were linear up to a plasma concentration of 5 mM (260 mg / L) with a turnover rate of ca. 9 µmol / kg bw / min = approx. 0.52 mg / kg of body weight / minute corresponding to a daily turnover of 750 mg / kg of body weight / day. Studies with repeated daily exposures of 6 or 8 hours have confirmed that bioaccumulation is not expected to occur until approx. 1,000 ppm (approximately 2,400 mg / m3 for 8 h / day 5 d / w) in humans and during 14 days of daily exposure in rats up to 11,000 ppm (26,550 mg / m3). For oral application to rats as a single bolus by gavage, the elimination of acetone appears to be saturated when blood levels rise above 300-400 mg / L corresponding to a dose of approximately 200 mg / kg body weight. Source ECHA

Information on likely routes of exposure

Xylene, mixture of isomers

XYLENE (MIXTURE OF ISOMERS)

WORKERS: inhalation; contact with the skin.

POPULATION: ingestion of contaminated food or water; inhalation of ambient air.

DRAP118 - PRIMER ES 40 COMP. A

Revision nr.3 Dated 08/06/2021 Printed on 08/06/2021 Page n. 11 / 21

Replaced revision:2 (Dated 06/10/2020)

SECTION 11. Toxicological information .../>>

ETHYLBENZENE

WORKERS: inhalation: contact with the skin.

POPULATION: ingestion of contaminated food or water; contact with the skin of products containing the substance.

Delayed and immediate effects as well as chronic effects from short and long-term exposure

Xylene, mixture of isomers

XYLENE (MIXTURE OF ISOMERS)

Toxic action on the central nervous system (encephalopathies); irritant action on the skin, conjunctiva, cornea and respiratory system.

FTHYI BENZENE

As the counterparts of benzene, may have an acute effect on the central nervous system, with depression, narcosis, often preceded by dizziness and associated with headache (IspesI). Is irritating for skin, conjunctiva and respiratory tract.

Interactive effects

Xylene, mixture of isomers

XYLENE (MIXTURE OF ISOMERS)

Alcohol intake interferes with the metabolism of the substance, inhibiting it. Consumption of ethanol (0.8 g / kg) before
4-hour exposure to xylenes vapors (145 and 280 ppm) causes a 50% decrease in metilippuric acid excretion,
while the blood concentration of xylenes rises about 1.5-2 times. At the same time there is an increase in side effects
secondary to ethanol. The metabolism of xylenes is enhanced by phenobarbital and 3-methyl-colanthrene-type enzyme inducers.
Aspirin and xylenes mutually inhibit their conjugation with glycine, which results in a decrease
urinary excretion of metilippuric acid. Other industrial products can interfere with the metabolism of xylenes.

ACUTE TOXICITY

ATE (Inhalation) of the mixture: > 20 mg/l
ATE (Oral) of the mixture: >2000 mg/kg
ATE (Dermal) of the mixture: >2000 mg/kg

Xylene, mixture of isomers

 LD50 (Oral)
 3523 mg/kg Ratto

 LD50 (Dermal)
 2000 mg/kg Coniglio

 LC50 (Inhalation)
 27,541 mg/l/4h Ratto

Reaction product: bisphenol-A-epichlorohydrin and epoxy resins (average molecular weight <= 700)

LD50 (Oral) > 11400 mg/kg Rat

Reaction product: bisphenol-F-epichlorohydrin and epoxy resins (average molecular weight <= 700)

 LD50 (Oral)
 > 2000 mg/kg Rat, OECD 420

 LD50 (Dermal)
 > 5000 mg/kg Rat, OECD 401

Oxirane, mono [(C12-14- alkyloxy) methyl] derivatives

LD50 (Oral) 17100 mg/kg Rat

ETHYLBENZENE

 LD50 (Oral)
 3500 mg/kg Rat

 LD50 (Dermal)
 15354 mg/kg Rabbit

 LC50 (Inhalation)
 17,2 mg/l/4h Rat

2-BUTOXYETHANOL

 LD50 (Oral)
 1300 mg/kg Rat

 LD50 (Dermal)
 > 2000 mg/kg Rabbit

 LC50 (Inhalation)
 450 ppm/4h Rat

Acetone

 LD50 (Oral)
 5800 mg/kg Rat

 LD50 (Dermal)
 7426 mg/kg Rat

 LC50 (Inhalation)
 76 mg/l/4h Rabbit

Reaction product: bisphenol-A-epichlorohydrin and epoxy resins (average molecular weight <= 700)

Acute toxicity - inhalation: Due to the very low vapor pressure (saturated atmosphere = 0.008 ppb), significant studies on the effects of acute inhalation could not be performed.

Acute toxicity - dermal: In a rat study according to the OECD standard n. 402 the dermal LD50 was> 2000 mg / kg. In several acute dermal toxicity studies in rabbits, LD50 was> 2000 mg / kg. In a rabbit study a LD50 value of 23 grams / kg was reported.

DRAP118 - PRIMER ES 40 COMP. A

Revision nr.3 Dated 08/06/2021 Printed on 08/06/2021 Page n. 12 / 21

Replaced revision:2 (Dated 06/10/2020)

SECTION 11. Toxicological information .../>>

Reaction product: bisphenol-F-epichlorohydrin and epoxy resins (average molecular weight <= 700)

Acute toxicity - inhalation: in accordance with Annex VII of the REACH Regulation, it is not necessary to conduct the acute toxicity study by inhalation absorption, since oral and dermal absorption studies are available for this substance.

In a rat study according to the OECD standard n. 402 the dermal LD50 was> 2000 mg / kg. In several acute dermal toxicity studies in rabbits, LD50 was> 2000 mg / kg. In a rabbit study a LD50 value of 23 grams / kg was reported.

Oxirane, mono [(C12-14- alkyloxy) methyl] derivatives

Acute toxicity - inhalation: No mortality was observed in rats exposed for 7 hours to saturated vapor (150 mg / m3).

LC50 (4h) 0.206 mg / I, Inhalation, Dusts / mists, Rat (0 Death.)

SKIN CORROSION / IRRITATION

Causes skin irritation

Reaction product: bisphenol-A-epichlorohydrin and epoxy resins (average molecular weight <= 700)

Result: skin - erythema / eschar 404 Acute Dermal Irritation / Corrosion

Species: rabbit Score 1.5 -2

Result: Skin - Edema 404 Acute Dermal Irritation / Corrosion

Species: Rabbit Score: 1.0 - 1.5

Result: eyes - 405 Acute Eye Irritation / Corrosion

Species: Rabbit Score: 0

Result: Conjunctive redness

Species: Rabbit Score: 0.7

Result: Skin - Moderately irritating

Species: Rabbit Exposure: 24 h

Result: Skin - Strongly irritating

Species: rabbit Exposure: 24 h

Result: eyes - mild irritant

Species: rabbit

Reaction product: bisphenol-F-epichlorohydrin and epoxy resins (average molecular weight <= 700)

Result: Skin - erythema / eschar 404 Acute Dermal Irritation / Corrosion

Species: rabbit Score: 0.7 Exposure: 4 h Observation: 72 h

Result: Skin - edema 404 Acete Dermal Irritation / Corrosion

Species: rabbit Score: 0 Exposure: 4 h Observation: 4-504 h

Result: eyes - corneal opacity 405 Acute Eye Irritation / corrosion

Species: Rabbit

Observation: 1 - 168 h

Result: eyes - Injury of the iris 405 Acute Eye Irritation / Corrosion

Species: Rabbit Score: 0

Observation: 1 - 168 h

Result: eyes - Redness of the conjunctiva 405 Acute Eye Irritation / Corrosion

Species: rabbit Score: 0

Observation: 1 - 168 h

DRAP118 - PRIMER ES 40 COMP. A

Revision nr.3 Dated 08/06/2021 Printed on 08/06/2021 Page n. 13 / 21

Replaced revision:2 (Dated 06/10/2020)

SECTION 11. Toxicological information .../>>

Result: eyes - conjunctiva edema 405 Acute Eye Irritation / Corrosion

Species: rabbit Score: 0

Observation: 1 - 168 h

Result: Skin - Slight irritation

Species: rabbit Exposure: 24 h

Oxirane, mono [(C12-14- alkyloxy) methyl] derivatives

Result: Skin - Primary index of skin irritation (PDII) OTS 798.4450 Acute Dermal Irritation

Species: Rabbit Score: 4.1 Exposure: 24 h Observation: 72 h

Result: Skin - Primary Index of Skin Irritation (PDII) 404 Acute Dermal Irritation / Corrosion

Species: Rabbit Score: 5.75 Exposure: 24 h Observation: 72 h

Result: eyes - corneal opacity 405 Acute Eye Irritation / Corrosion

Species: Rabbit Score: 2

Observation: 1 - 24 h

Species: Skin - Moderately irritating

Species: Rabbit Exposure: 24 h

SERIOUS EYE DAMAGE / IRRITATION

Causes serious eye irritation

RESPIRATORY OR SKIN SENSITISATION

Sensitising for the skin

Reaction product: bisphenol-F-epichlorohydrin and epoxy resins (average molecular weight <= 700)

Species: rabbit Test: OECD 405 Result: no eye irritation

Skin sensitization

Reaction product: bisphenol-A-epichlorohydrin and epoxy resins (average molecular weight <= 700)

In a study with an LLNA assay on mice conducted according to the OECD standard n. 429, the estimated EC3 corresponded to a concentration of 5.7%; this result suggests that BADGE is a moderate skin sensitizer in this test system. In a guinea pig maximization study according to the OECD standard n. 406, BADGE induced a positive skin reaction in 100% of the experimental animals at a stimulus dose with a concentration of 50%. Therefore, BADGE is an "extreme" skin sensitizer in the conditions of this study. BADGE was positive for skin sensitization also in a study with the Buehler method on guinea pig conducted according to OCSE standard n. 406

Reaction product: bisphenol-F-epichlorohydrin and epoxy resins (average molecular weight <= 700)

The Buehler method was used to assess the skin sensitization potential of the liquid epoxy BPFDGE. Ten male guinea pigs were given 0.4 ml of the test substance topically once a week for three weeks. A positive control of BPFDGE liquid epoxy resin was used on ten additional animals. The stimulation phase began two weeks later with the addition of 5 animals exposed to 0.4ml of liquid BPFDGE liquid epoxy resin. The negative control had 0 positive reactions; BPFDGE liquid epoxy resin produced positive reactions in 4 out of 10 guinea pigs and the positive control had 8 out of ten positive reactions. Under the conditions of this study, the test material resulted in delayed hypersensitivity in guinea pigs.

Route of exposure: skin Species: mouse Method: OECD 429

Result: can cause sensitization in contact with the skin

DRAP118 - PRIMER ES 40 COMP. A

Dated 08/06/2021 Printed on 08/06/2021 Page n. 14 / 21 Replaced revision:2 (Dated 06/10/2020)

Revision nr.3

SECTION 11. Toxicological information .../>>

Oxirane, mono [(C12-14- alkyloxy) methyl] derivatives

In a sensitization study with the Buehler method carried out according to the test specification OTS 870.2600 of the US EPA, positive skin reactions were observed in 20/20 guinea pigs. An extreme sensitizer in a study with maximization test on guinea pig conducted according to the OECD test specification No. 406.

GERM CELL MUTAGENICITY

Does not meet the classification criteria for this hazard class

Reaction product: bisphenol-A-epichlorohydrin and epoxy resins (average molecular weight <= 700)

In several studies it was found that BADGE induces gene mutation in experimental strains Ames / Salmonella TA1535 and TA100. In general, mutagenic activity was greater without S9 metabolic activation of the liver. Induced gene mutation in L5178Y mouse lymphoma cells. Induced gene mutation and chromosomal damage in V79 Chinese hamster cells. Induced cell transformation in Syrian hamster BHK cells based on clonal growth in soft agar. It did not induce evidence of chromosomal damage in a study with an oral probe in a test of the dominant lethal on mice conducted up to a high dosage level of 10 grams / kg and in a micronuclear test on mice conducted up to a high dose of 5000 mg / kg. Negative in a spermatocytic cytogenetic assay on male mice with treatment for 5 days by oral probe up to a high dose of 3000 mg / kg. It did not induce an increase in the frequency of chromosomal damage in a cytogenetic assay on bone marrow cells on a Chinese hamster using an oral probe up to a high dose of 3300 mg / kg. It did not induce an increase in DNA strand breaks in rat liver cells after treatment with oral gavage with 500 mg / kg, measured by alkaline elution

Reaction product: bisphenol-F-epichlorohydrin and epoxy resins (average molecular weight <= 700)

Bisphenol F diglycidyl ether induced a gene mutation in the Ames / Salmonella mutation test and chromosomal aberrations in human lymphocytes in multiple independent GLP studies conducted according to test guidelines. Furthermore, the structural analog, bisphenol A diglycidyl ether (BPADGE), induced a significant increase in the frequency of mutations in cultured L5178Y mouse lymphoma cells, supporting the other conclusions. Therefore, BPFDGE is genotoxic in vitro. When the genotoxic potential of bisphenol F diglycidyl ether was evaluated in multiple GLP compliant in vivo tests, including mouse micronucleus tests, UDS in vivo / in vitro tests, and MutaMouse on rat, no evidence of genotoxicity was observed. Results from other in vivo genotoxicity tests also support these negative results for BPFDGE. It is concluded that bisphenol F diglycidyl ether is not genotoxic in vivo.

In vitro genotoxicity:

Metabolic activation: with or without metabolic activation

Method: OECD 471 Result: positive

Metabolic activation: with or without metabolic activation

Method: OECD 473 Result: positive

Metabolic activation: with or without metabolic activation

Method: OECD 476 Result: positive

Genotoxicity in vivo: Type of gellule: germs Method of application: oral

Result: negative

Type of gellula: somatic Method of application: oral Dose: 0 - 5000 mg / kg Result: negative

Oxirane, mono [(C12-14- alkyloxy) methyl] derivatives

Positive in a bacterial mutation test conducted according to the OECD test specification No. 471 in an experimental Salmonella TA1535 strain with and without metabolic activation with S9. Negative in a gene mutation test on Chinese hamster ovary cells (CHO) HGPRT conducted according to the OECD test specification No. 476 up to cytotoxic levels with and without metabolic activation with S9. Negative in a gene mutation assay on L5178Y / TK mouse lymphoma cells tested up to cytotoxic dose levels. Negative by micronucleus induction (chromosomal damage) in a mouse study conducted according to OECD specification No. 474 up to a high dose of intraperitoneal injection of 4.0 grams / kg. Negative in a study of chromosomal aberrations on rat bone marrow conducted similarly to the OECD test specification No. 475 by intraperitoneal injection, up to a high dose of about 700 mg / kg.

CARCINOGENICITY

Does not meet the classification criteria for this hazard class

Acetone

The ACGIH classifies acetone as A4, that is, not classifiable as a human carcinogen: An agent that suggests that it may be carcinogenic to humans but which cannot be definitively assessed due to insufficient data. In vitro or animal studies do not provide sufficient carcinogenicity indications to classify the agent in one of the other categories.

DRAP118 - PRIMER ES 40 COMP. A

Revision nr.3 Dated 08/06/2021 Printed on 08/06/2021 Page n. 15 / 21

Replaced revision:2 (Dated 06/10/2020)

SECTION 11. Toxicological information .../>>

Reaction product: bisphenol-A-epichlorohydrin and epoxy resins (average molecular weight <= 700)

In a study with an oral rat probe according to the OECD standard n. 453 there was no evidence of carcinogenicity up to the high dose level of 100 mg / kg / day. Dermal exposure studies were performed in male mice and female rats according to OECD regulation n. 453. No evidence of carcinogenicity was observed in male mice treated up to the high dose of 100 mg / kg / day and female rats exposed up to the high dose of 1000 mg / kg / day.

Reaction product: bisphenol-F-epichlorohydrin and epoxy resins (average molecular weight <= 700)

È stata valutata la capacità del Bisfenolo F diglicidiletere (BPFDGE) di indurre tumori locali e sistemici in uno studio di 24 mesi con test cutaneo ("skin painting") sul topo. Il trattamento cutaneo di topi per due volte alla settimana con una soluzione fino al 10% di diglicidiletere bisfenolo F (BPFDGE) non ha indotto alcun risultato negativo di incidenza di tumori o effetti cutanei locali. Pertanto, il BPFDGE non è da considerare cancerogeno per il topo nelle condizioni di questo studio. Il NOAEL è stato stimato pari a circa 800 mg/kg/die.

Specie: ratto, maschio e femmina Modalità di applicazione: orale Tempo diesposizione: 24 mesi

Dosi: 15 mg/kg

Frequenza del trattamento: 7 al giorno

Metodo: OECD 453 Risultato: negativo

Specie: topo, maschio

Modalità di applicazione: dermico Tempo di esposizione: 24 mesi

Dosi: 1 mg/kg

Frequenza del trattamento: 3 al giorno

Metodo: OECD 453 Risultato: negativo

Specie: ratto, femmina

Modalità di applicazione: dermico Tempo di esposizione: 24 mesi

Dosi: 1 mg/kg

Frequenza del trattamento: 5 al giorno

Metodo: OECD 453 Risultato: negativo

Xylene, mixture of isomers

XYLENE (MIXTURE OF ISOMERS)

Classified in group 3 (not classifiable as a human carcinogen) by the International Agency for Research on Cancer (IARC).

The US Environmental Protection Agency (EPA) argues that "the data was found to be inadequate for a potential assessment carcinogenic".

ETHYLBENZENE

Classified in Group 2B (possible human carcinogen) by the International Agency for Research on Cancer (IARC) - (IARC, 2000). Classified in Group D (not classifiable as a human carcinogen) by the US Environmental Protection Agency (EPA) - (US EPA file on-line 2014).

REPRODUCTIVE TOXICITY

Does not meet the classification criteria for this hazard class

Reaction product: bisphenol-A-epichlorohydrin and epoxy resins (average molecular weight <= 700)

BADGE did not induce any evidence of developmental toxicity in rats and rabbits exposed by oral probe, or in cutaneously treated rabbits, in BPL studies according to the OECD standard n. 414. Studies with an oral probe were conducted up to a high dose level of 180 mg / kg / day which produced maternal toxicity based on the reduction in body weight gain. The rabbit skin toxicity study was conducted up to a high dose of 300 mg / kg / day which induced maternal toxicity based on the reduction in body weight gain.

Reaction product: bisphenol-F-epichlorohydrin and epoxy resins (average molecular weight <= 700)

The bisphenol A diglycidylether (DGEBPA) has been tested for its embryo / fetal toxicity and teratogenicity in pregnant rabbits. The DGEBPA was applied daily to the back (depilated) of white New Zealand rabbits at doses of 0 (polyethylene glycol, vehicle control), 30, 100 or 300 mg / kg of body weight / day at a volumetric dose of 1 ml / kg of body weight / day on days 6 to 18 of gestation. Twenty-six inseminated rabbits were used per dosage group, obtaining a minimum of 20 pregnant rabbits per exposure level. An occlusive bandage of absorbent gauze and non-absorbent cotton was placed on the dosing area on the back of each rabbit. The bandage was held in place for a minimum of 6 hours / day with a lycra / spandex protective cover. Following the period of occlusion the bandage and the protective wrapper were removed. Maternal toxicity effects were observed among pregnant rabbits in the 300 mg / kg dose group, as evidenced by moderate to severe erythema, fissures, haemorrhages and mild edema at the site of exposure. Similar, but less severe skin lesions were observed in pregnant rabbits in the 100 mg / kg / day exposure group. Effects on the skin (mild erythema) observed in pregnant rabbits in the 30 mg / kg / day group were not considered toxicologically significant. No

DRAP118 - PRIMER ES 40 COMP. A

Revision nr.3 Printed on 08/06/2021

Replaced revision:2 (Dated 06/10/2020)

SECTION 11. Toxicological information .../>>

evidence of embryo / fetotoxicity or teratogenicity was observed at any dose, which results in a level at which no effect (NOEL) is observed at an embryonic / fetal level of 300 mg / kg body weight / day.

Oxirane, mono [(C12-14- alkyloxy) methyl] derivatives

In a dermal toxicological study conducted in the rat according to the US EPA OTS 798.4420 method and according to the OECD test specification No. 414, the NOAEL for adverse effects on both mother and development was above the high dose level of 200 mg / kg /

Adverse effects on sexual function and fertility

Reaction product: bisphenol-F-epichlorohydrin and epoxy resins (average molecular weight <= 700)

Species: rat, male and female Method of application: oral Method: OECD 416

Result: There was no effect on fertility and early embryonic development.

Adverse effects on development of the offspring

Reaction product: bisphenol-F-epichlorohydrin and epoxy resins (average molecular weight <= 700)

Species: rabbit, female Method of application: dermal

General toxicity in mothers: no level of harmfulness

observed: 30 mg / kg body weight Result: no teratogenic effect

STOT - SINGLE EXPOSURE

May cause drowsiness or dizziness

STOT - REPEATED EXPOSURE

Does not meet the classification criteria for this hazard class

Reaction product: bisphenol-F-epichlorohydrin and epoxy resins (average molecular weight <= 700)

NOAEL: 250 mg / kg

Species: rat, male and female Method of application: ingestion Exposure time: 13 weeks Number of exposures: 7 d Mode: subchronic toxicity

ASPIRATION HAZARD

Does not meet the classification criteria for this hazard class

SECTION 12. Ecological information

This product is dangerous for the environment and is toxic for aquatic organisms. In the long term, it have negative effects on acquatic environment.

12.1. Toxicity

Reaction product: bisphenol-F-epichlorohydrin and epoxy resins (average molecular weight <= 700)

EC50 (Selenastrum capricornutum): 1.8 mg / I

Exposure time: 72 h

Static test

Method: OECD 201

CI50 (activated sludge)> 100 mg / I

Exposure time: 3 h Static test

Oxirane, mono [(C12-14- alkyloxy) methyl] derivatives

LC50, 96 hour:> 5000 mg / I, Oncorhynchus mykiss (rainbow trout)

LC50, 96 hour: 1800 mg / I, Lepomis macrochirus (Perch) EC50, 72 hours: 843 mg / I, Pseudokirchneriella subcapitata NOEC, 72 hours: 500 mg / I, Pseudokirchneriella subcapitata

EC50, 3 hours:> 100 mg / I, Activated sludge

DRAP118 - PRIMER ES 40 COMP. A

Revision nr.3 Dated 08/06/2021 Printed on 08/06/2021 Page n. 17 / 21

Replaced revision:2 (Dated 06/10/2020)

SECTION 12. Ecological information .../>>

Reaction product: bisphenol-A-epichlorohydrin and epoxy resins (average molecular weight <= 700) LC50 - for Fish 1,3 mg/l/96h 203 Fish, Acute Toxicity Test

EC50 - for Crustacea 2,1 mg/l/48h 202 Daphnia sp. Acute Immobilization Test and Reproduction Test

EC50 - for Algae / Aquatic Plants > 11 mg/l/72h Algae

Chronic NOEC for Crustacea 0,3 mg/l 11 Daphnia Magna Reproduction Test (21 d)

Reaction product: bisphenol-F-epichlorohydrin and epoxy resins (average molecular weight <= 700)

LC50 - for Fish 2,54 mg/l/96h Fish

EC50 - for Crustacea 2,55 mg/l/48h 202 Daphnia sp. Acute Immobilization Test and Reproduction Tes

EC50 - for Algae / Aquatic Plants > 1000 mg/l/72h 201 Alga, Growth Inhibition Test
Chronic NOEC for Crustacea 0,3 mg/l Daphnia magna, 21 d, OECD 211 semistatic

Oxirane, mono [(C12-14- alkyloxy) methyl] derivatives

LC50 - for Fish > 5 g/l 203 Fish, Acute Toxicity Test. Bluegill

EC50 - for Crustacea 7,2 mg/l/48h 202 Daphnia sp. Acute Immobilization Test and Reproduction Test

EC50 - for Algae / Aquatic Plants 844 mg/l/72h 201 Alga, Growth Inhibition Test

2-BUTOXYETHANOL

LC50 - for Fish 1474 mg/l/96h Oncorhynchus mykiss EC50 - for Crustacea 1550 mg/l/48h Daphnia magna

Acetone

LC50 - for Fish 5540 mg/l/96h Lepomis macrochirus EC50 - for Crustacea 8800 mg/l/48h Daphnia pulex Chronic NOEC for Crustacea 2212 mg/l Daphnia magna , 28 d

12.2. Persistence and degradability

Reaction product: bisphenol-A-epichlorohydrin and epoxy resins (average molecular weight <= 700)

The level of biodegradation in an "improved" OECD 301F study was 5% within the 28-day contact period. Biodegradation has reached 6 - 12% after 28 days of contact in a study conducted according to the OECD standard n. 301B. Therefore BADGE is not readily biodegradable under the conditions of the studies.

Reaction product: bisphenol-F-epichlorohydrin and epoxy resins (average molecular weight <= 700)

Bisphenol F diglycidyl ether was not readily biodegradable under the conditions of the screening studies according to OECD test specifications Nos. 301 B and 301 D. The maximum rate of biodegradation observed in one of the OECD 301 B studies was 16% per 10 mg / the 28 days of contact.

Inoculum: activated sludge Concentration: 3 mg / I Result: not biodegradable Biodegradation: approx. 0% Exposure time: 28 d

Method: Directive 67/548 / EEC Annex V C.4.E

Oxirane, mono [(C12-14- alkyloxy) methyl] derivatives

In a study conducted according to OECD Test Specification No. 301 F, biodegradation was 57-655 after 7 days. However, in a study conducted according to OECD Test Specification No. 301 D (unopened bottle) biodegradation was only 34.7% after 28 days.

87% degradation: 28 days $\,$ OECD 301F $\,$

Xylene, mixture of isomers

Solubility in water 60 mg/l ASTM E1148

Degradability: information not available

ETHYLBENZENE

Solubility in water 1000 - 10000 mg/l

Rapidly degradable

2-BUTOXYETHANOL

Solubility in water 900 g/l 20°C

Rapidly degradable

Acetone

Rapidly degradable

12.3. Bioaccumulative potential

DRAP118 - PRIMER ES 40 COMP. A

Revision nr.3 Dated 08/06/2021 Printed on 08/06/2021 Page n. 18 / 21 Replaced revision:2 (Dated 06/10/2020)

SECTION 12. Ecological information .../>>

Reaction product: bisphenol-A-epichlorohydrin and epoxy resins (average molecular weight <= 700)

LogPOW: 2.64 - 3.78 BCF: 3 - 31 31.00 Potential: low

Reaction product: bisphenol-F-epichlorohydrin and epoxy resins (average molecular weight <= 700)

BCF: 150 150.00 Potential: low

Oxirane, mono [(C12-14- alkyloxy) methyl] derivatives

BCF: 160 - 263 160.00

Potential: low

Xylene, mixture of isomers

Partition coefficient: n-octanol/water 3,16 BCF 25,9

Reaction product: bisphenol-F-epichlorohydrin and epoxy resins (average molecular weight <= 700)

Partition coefficient: n-octanol/water 2,7 Log Kow

BCF 150

Oxirane, mono [(C12-14- alkyloxy) methyl] derivatives

Partition coefficient: n-octanol/water 3,77 Log Kow

ETHYLBENZENE

Partition coefficient: n-octanol/water 3,6

2-BUTOXYETHANOL

Partition coefficient: n-octanol/water 0,81

Acetone

Partition coefficient: n-octanol/water -0,24 BCF 3

12.4. Mobility in soil

Reaction product: bisphenol-F-epichlorohydrin and epoxy resins (average molecular weight <= 700)
Partition coefficient: soil/water 4460 OECD 121

12.5. Results of PBT and vPvB assessment

On the basis of available data, the product does not contain any PBT or vPvB in percentage ≥ than 0,1%.

12.6. Other adverse effects

Information not available

SECTION 13. Disposal considerations

13.1. Waste treatment methods

Reuse, when possible. Product residues should be considered special hazardous waste. The hazard level of waste containing this product should be evaluated according to applicable regulations.

Disposal must be performed through an authorised waste management firm, in compliance with national and local regulations.

Waste transportation may be subject to ADR restrictions.

CONTAMINATED PACKAGING

Contaminated packaging must be recovered or disposed of in compliance with national waste management regulations.

SECTION 14. Transport information

14.1. UN number

ADR / RID, IMDG, IATA: 1263

DRAP118 - PRIMER ES 40 COMP. A

Revision nr.3 Dated 08/06/2021 Printed on 08/06/2021 Page n. 19 / 21

Replaced revision:2 (Dated 06/10/2020)

SECTION 14. Transport information .../>>

14.2. UN proper shipping name

ADR / RID: PAINT OF PAINT RELATED MATERIAL IMDG: PAINT OF PAINT RELATED MATERIAL IATA: PAINT OF PAINT RELATED MATERIAL

14.3. Transport hazard class(es)

ADR / RID: Class: 3 Label: 3

IMDG: Class: 3 Label: 3

IATA: Class: 3 Label: 3



14.4. Packing group

ADR / RID, IMDG, IATA: III

14.5. Environmental hazards

ADR / RID: NO IMDG: NO IATA: NO

14.6. Special precautions for user

ADR / RID: HIN - Kemler: 30 Limited Quantities: 5 L Tunnel restriction code: (D/E)

Special provision: IMDG: EMS: F-E, S-E Limited Quantities: 5 L

IATA: Cargo: Maximum quantity: 220 L Packaging instructions: 366

Pass.: Maximum quantity: 60 L Packaging instructions: 355

Special provision: A3, A72, A192

14.7. Transport in bulk according to Annex II of Marpol and the IBC Code

Information not relevant

SECTION 15. Regulatory information

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

Seveso Category - Directive 2012/18/EC: P5c-E2

Restrictions relating to the product or contained substances pursuant to Annex XVII to EC Regulation 1907/2006

<u>Product</u>

Point

Point 3 - 40

Contained substance

Point 75 Reaction product: bisphenol-A-epichlorohydrin and epoxy resins (average molecular weight <=

700)

Reg. no.: 01-2119456619-26-XXXX

Point 75 Oxirane, mono [(C12-14- alkyloxy) methyl] derivatives

Reg. no.: 01 2119485289-22-XXXX

Xylene, mixture of isomers

Reg. no.: 01-2119488216-32-XXXX

Point 75 2-BUTOXYETHANOL

75

Regulation (EC) No. 2019/1148 - on the marketing and use of explosives precursors

Regulated explosives precursor

The acquisition, introduction, possession or use of that regulated explosives precursor by members of the general public is subject to reporting obligations as set out in Article 9.

All suspicious transactions and significant disappearances and thefts must be reported to the relevant national contact point.

DRAP118 - PRIMER ES 40 COMP. A

Revision nr.3 Dated 08/06/2021 Printed on 08/06/2021 Page n. 20 / 21 Replaced revision:2 (Dated 06/10/2020)

SECTION 15. Regulatory information ... / >>

Substances in Candidate List (Art. 59 REACH)

On the basis of available data, the product does not contain any SVHC in percentage ≥ than 0,1%.

Substances subject to authorisation (Annex XIV REACH)

None

Substances subject to exportation reporting pursuant to (EC) Reg. 649/2012:

None

Substances subject to the Rotterdam Convention:

None

Substances subject to the Stockholm Convention:

None

Healthcare controls

Workers exposed to this chemical agent must not undergo health checks, provided that available risk-assessment data prove that the risks related to the workers' health and safety are modest and that the 98/24/EC directive is respected.

15.2. Chemical safety assessment

A chemical safety assessment has been performed for the following contained substances Xylene, mixture of isomers

SECTION 16. Other information

Text of hazard (H) indications mentioned in section 2-3 of the sheet:

Flam. Liq. 2 Flammable liquid, category 2
Flam. Liq. 3 Flammable liquid, category 3
Acute Tox. 4 Acute toxicity, category 4
Asp. Tox. 1 Aspiration hazard, category 1

STOT RE 2 Specific target organ toxicity - repeated exposure, category 2

Eye Irrit. 2 Eye irritation, category 2 Skin Irrit. 2 Skin irritation, category 2

STOT SE 3 Specific target organ toxicity - single exposure, category 3

Skin Sens. 1 Skin sensitization, category 1

Aquatic Chronic 2 Hazardous to the aquatic environment, chronic toxicity, category 2 **Aquatic Chronic 3** Hazardous to the aquatic environment, chronic toxicity, category 3

H225Highly flammable liquid and vapour.H226Flammable liquid and vapour.H302Harmful if swallowed.H312Harmful in contact with skin.

H332 Harmful if inhaled.

H304 May be fatal if swallowed and enters airways.

H373 May cause damage to organs through prolonged or repeated exposure.

H319 Causes serious eye irritation.H315 Causes skin irritation.

H335 May cause respiratory irritation.
 H317 May cause an allergic skin reaction.
 H336 May cause drowsiness or dizziness.
 H411 Toxic to aquatic life with long lasting effects.
 H412 Harmful to aquatic life with long lasting effects.

EUH066 Repeated exposure may cause skin dryness or cracking.

LEGEND:

- ADR: European Agreement concerning the carriage of Dangerous goods by Road
- CAS NUMBER: Chemical Abstract Service Number
- CE50: Effective concentration (required to induce a 50% effect)
- CE NUMBER: Identifier in ESIS (European archive of existing substances)
- CLP: EC Regulation 1272/2008
- DNEL: Derived No Effect Level
- EmS: Emergency Schedule
- GHS: Globally Harmonized System of classification and labeling of chemicals
- IATA DGR: International Air Transport Association Dangerous Goods Regulation
- IC50: Immobilization Concentration 50%
- IMDG: International Maritime Code for dangerous goods

DRAP118 - PRIMER ES 40 COMP. A

Revision nr.3 Dated 08/06/2021 Printed on 08/06/2021 Page n. 21 / 21 Replaced revision:2 (Dated 06/10/2020)

SECTION 16. Other information .../>>

- IMO: International Maritime Organization- INDEX NUMBER: Identifier in Annex VI of CLP
- LC50: Lethal Concentration 50%
- LD50: Lethal dose 50%
- OEL: Occupational Exposure Level
- PBT: Persistent bioaccumulative and toxic as REACH Regulation
- PEC: Predicted environmental Concentration
- PEL: Predicted exposure level
- PNEC: Predicted no effect concentration
- REACH: EC Regulation 1907/2006
- RID: Regulation concerning the international transport of dangerous goods by train
- TLV: Threshold Limit Value
- TLV CEILING: Concentration that should not be exceeded during any time of occupational exposure.
- TWA STEL: Short-term exposure limit
- TWA: Time-weighted average exposure limit
- VOC: Volatile organic Compounds
- vPvB: Very Persistent and very Bioaccumulative as for REACH Regulation
- WGK: Water hazard classes (German).

GENERAL BIBLIOGRAPHY

- 1. Regulation (EC) 1907/2006 (REACH) of the European Parliament
- 2. Regulation (EC) 1272/2008 (CLP) of the European Parliament
- 3. Regulation (EU) 790/2009 (I Atp. CLP) of the European Parliament
- 4. Regulation (EU) 2015/830 of the European Parliament
- 5. Regulation (EU) 286/2011 (II Atp. CLP) of the European Parliament
- 6. Regulation (EU) 618/2012 (III Atp. CLP) of the European Parliament
- 7. Regulation (EU) 487/2013 (IV Atp. CLP) of the European Parliament
- 8. Regulation (EU) 944/2013 (V Atp. CLP) of the European Parliament
- 9. Regulation (EU) 605/2014 (VI Atp. CLP) of the European Parliament
- 10. Regulation (EU) 2015/1221 (VII Atp. CLP) of the European Parliament
- 11. Regulation (EU) 2016/918 (VIII Atp. CLP) of the European Parliament
- 12. Regulation (EU) 2016/1179 (IX Atp. CLP)
- 13. Regulation (EU) 2017/776 (X Atp. CLP)
- 14. Regulation (EU) 2018/669 (XI Atp. CLP)
- 15. Regulation (EU) 2018/1480 (XIII Atp. CLP)
- 16. Regulation (EU) 2019/521 (XII Atp. CLP)
- 17. Regulation (EU) 2019/1148
- 18. Regulation (EU) 2020/217 (XIV Atp. CLP)
- The Merck Index. 10th Edition
- Handling Chemical Safety
- INRS Fiche Toxicologique (toxicological sheet)
- Patty Industrial Hygiene and Toxicology
- N.I. Sax Dangerous properties of Industrial Materials-7, 1989 Edition
- IFA GESTIS website
- ECHA website
- Database of SDS models for chemicals Ministry of Health and ISS (Istituto Superiore di Sanità) Italy

Note for users:

The information contained in the present sheet are based on our own knowledge on the date of the last version. Users must verify the suitability and thoroughness of provided information according to each specific use of the product.

This document must not be regarded as a guarantee on any specific product property.

The use of this product is not subject to our direct control; therefore, users must, under their own responsibility, comply with the current health and safety laws and regulations. The producer is relieved from any liability arising from improper uses.

Provide appointed staff with adequate training on how to use chemical products.

CALCULATION METHODS FOR CLASSIFICATION

Chemical and physical hazards: Product classification derives from criteria established by the CLP Regulation, Annex I, Part 2. The data for evaluation of chemical-physical properties are reported in section 9.

Health hazards: Product classification is based on calculation methods as per Annex I of CLP, Part 3, unless determined otherwise in Section

Environmental hazards: Product classification is based on calculation methods as per Annex I of CLP, Part 4, unless determined otherwise in Section 12.

Changes to previous review:

The following sections were modified:

02/03/08/09/10/11/12/15/16.