

Product brands by Wilhelmsen



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L.REACH.NOR.EN

CargoClean HD

Wilhelmsen Ships Service AS

Part Number: 779104 (25 liter) Version No: 5.23 Safety Data Sheet (Conforms to Annex II of REACH (1907/2006) - Regulation 2020/878)

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

1.1. Product Identifier

Product name	CargoClean HD
Chemical Name	Not Applicable
Synonyms	Not Available
Proper shipping name	CAUSTIC ALKALI LIQUID, N.O.S. (contains potassium hydroxide)
Chemical formula	Not Applicable
Other means of identification	779104 (25 liter), 779104

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

Procedural Category	PROC7 Industrial spraying		
Chemical Product Category	PC35 Washing and cleaning products		
Sectors of Use SU22 Professional uses: Public domain (administration, education, entertainment, services, craftsmen) SU3 Industrial uses: Uses of substances as such or in preparations* at industrial sites			
Relevant identified uses	Heavy duty alkaline cleaner		
Uses advised against	No specific uses advised against are identified.		

1.3. Details of the manufacturer or supplier of the safety data sheet

Registered company name	Wilhelmsen Ships Service AS	Outback (M)SDS portal: http://jr.chemwatch.net/outb/account /autologin?login=wilhelmsen	Wilhelmsen Ships Service AS* Central Warehouse
Address	Strandveien 20 Lysaker 1366 Norway	Use our Outback portal to obtain our (M)SDSs in other languages and/or formatFor questions relating to our SDSs please use Email: WSS.GLOBAL.SDSINFO@wilhelmsen.com Norway	Willem Barentszstraat 50 Rotterdam Netherlands
Telephone	+47 67 58 40 00	Not Available	+31 10 4877 777
Fax	Not Available	Not Available Not Available	
Website	http://www.wilhelmsen.com/	ww.wilhelmsen.com/ http://www.wilhelmsen.com	
Email	wss.norway.cs@wilhelmsen.com	wss.global.sdsinfo@wilhelmsen.com	wss.rotterdam@wilhelmsen.com
	1		
Registered company name	Wilhelmsen Ships Service AS* Central Warehouse		
Address	Willem Barentszstraat 50 Rotterdam Netherlands		

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Telephone	+31 10 4877 777
Fax	Not Available
Website	http://www.wilhelmsen.com
Email	wss.rotterdam@wilhelmsen.com

1.4. Emergency telephone number

Association / Organisation	Giftinformasjonssentralen - 24 timer 24hrs - Chemwar		vatch	Dutch nat. poison centre	
Emergency telephone numbers	+47 22591300	+31-10-4877700		+ 31 88 7558561	
Other emergency telephone numbers	+31-10-4877700	+31-10-4877700		+ 31 10 4877700	
Association / Organisation	Dutch nat. poison centre		CHEMWATCH EMERGENCY RESPONSE (24/7)		
Emergency telephone numbers	+ 31 30 274 88 88		+47 23 25 25 84		
Other emergency telephone numbers	+ 31-10-4877700		+61 3 9573 3188		

Once connected and if the message is not in your preferred language then please dial 01

SECTION 2 Hazards identification

2.1. Classification of the substance or mixture

Classification according to regulation (EC) No 1272/2008 [CLP] and amendments ^[1]	H290 - Corrosive to Metals Category 1, H314 - Skin Corrosion/Irritation Category 1A	
Legend:	1. Classified by Chemwatch; 2. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI	

2.2. Label elements



Hazard statement(s)

H290	May be corrosive to metals.
H314	Causes severe skin burns and eye damage.

Supplementary statement(s)

EUH208	Contains N,N-dimethyldecanamide. May produce an allergic reaction.
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Precautionary statement(s) General

P101	If medical advice is needed, have product container or label at hand.		
P102	Keep out of reach of children.		
P103	Read carefully and follow all instructions.		

Precautionary statement(s) Prevention

P260	Do not breathe mist/vapours/spray.
P264	Wash all exposed external body areas thoroughly after handling.
P280	Wear protective gloves, protective clothing, eye protection and face protection.
P234	Keep only in original packaging.

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Precautionary statement(s) Response

P301+P330+P331	IF SWALLOWED: Rinse mouth. Do NOT induce vomiting.		
P303+P361+P353	IF ON SKIN (or hair): Take off immediately all contaminated clothing. Rinse skin with water [or shower].		
P305+P351+P338	IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.		
P310	Immediately call a POISON CENTER/doctor/physician/first aider.		
P363	Wash contaminated clothing before reuse.		
P390	Absorb spillage to prevent material damage.		
P304+P340	IF INHALED: Remove person to fresh air and keep comfortable for breathing.		

Precautionary statement(s) Storage

P405 Store locked up.

Precautionary statement(s) Disposal

P501 Dispose of contents/container to authorised hazardous or special waste collection point in accordance with any local regulation.

2.3. Other hazards

hydrocarbons, C10-13-	Determined to have and earling discupting properties according to Europe Pegulation (EU) 528/2012, Europe Pegulation (EU)
n-alkanes, isoalkanes,	2017/2100 and Europe Regulation (EU) 2018/605
cyclics, < 2% aromatics	

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] and amendments	SCL / M-Factor	Nanoform Particle Characteristics
1. 1310-58-3 2.215-181-3 3.019-002-00-8 4.Not Available	1-5	potassium hydroxide	Acute Toxicity (Oral) Category 4, Skin Corrosion/Irritation Category 1A; H302, H314 ^[2]	Skin Corr. 1A; H314: $C \ge 5 \%$ Skin Corr. 1B; H314: $2 \% \le C <$ 5 % Skin Irrit. 2; H315: $0,5 \% \le C$ < 2 % Eye Irrit. 2; H319: $0,5 \% \le$ C < 2 %	Not Available
1. 10213-79-3 2.Not Available 3.014-010-00-8 4.Not Available	1-5	sodium metasilicate, pentahydrate	Skin Corrosion/Irritation Category 1B, Specific Target Organ Toxicity - Single Exposure (Respiratory Tract Irritation) Category 3; H314, H335 ^[2]	Not Available	Not Available
1. 1554325-20-0 2.Not Available 3.Not Available 4.Not Available	1-5	<u>C12-14-</u> alkyl(hydroxyethyl)dimethyl chloride, ethoxylated	Acute Toxicity (Oral) Category 4, Skin Corrosion/Irritation Category 2, Serious Eye Damage/Eye Irritation Category 1, Hazardous to the Aquatic Environment Acute Hazard Category 1; H302, H315, H318, H400 ^[1]	Not Available	Not Available
1. 160875-66-1* 2.605-233-7 3.Not Available 4.Not Available	1-5	Fatty alcohol ethoxylates*	Serious Eye Damage/Eye Irritation Category 1, Acute Toxicity (Oral) Category 3; H318, H301 ^[1]	0	Not Available
1. 68891-38-3 2.500-234-8 3.Not Available 4.Not Available	1-5	sodium linear-(C12-14)alkyl ether sulfate	Acute Toxicity (Oral) Category 4, Skin Corrosion/Irritation Category 2, Serious Eye Damage/Eye Irritation Category 2; H302, H315, H319 ^[1]	Not Available	Not Available
1. 68155-07-7* 2.268-935-9	1-5	cocamide diethanolamide.	Skin Corrosion/Irritation Category 2, Hazardous to the Aquatic	Not Available	Not Available

1. CAS No 2.EC No 3.Index No 4.REACH No	%[weight]	Name	Classification according to regulation (EC) No 1272/2008 [CLP] and amendments	SCL / M-Factor	Nanoform Particle Characteristics
3.Not Available 4.Not Available			Environment Long-Term Hazard Category 2, Serious Eye Damage/Eye Irritation Category 1; H315, H411, H318 ^[1]		
1. 14433-76-2 2.238-405-1 3.Not Available 4.Not Available	1-5	N.N-dimethyldecanamide	Skin Corrosion/Irritation Category 2, Serious Eye Damage/Eye Irritation Category 1, Sensitisation (Skin) Category 1, Specific Target Organ Toxicity - Repeated Exposure Category 2, Hazardous to the Aquatic Environment Long-Term Hazard Category 3; H315, H318, H317, H373, H412 ^[1]	Not Available	Not Available
1. Not Available 2.Not Available 3.Not Available 4.Not Available	1-5	hydrocarbons, C10-13- n-alkanes, isoalkanes, cyclics, < 2% aromatics ^[e]	Flammable Liquids Category 3, Specific Target Organ Toxicity - Single Exposure (Narcotic Effects) Category 3, Aspiration Hazard Category 1; H226, H336, H304 ^[1]	Not Available	Not Available
1. 111-76-2 2.203-905-0 3.603-014-00-0 4.Not Available	1-5	ethylene glycol monobutyl ether *	Acute Toxicity (Oral) Category 4, Acute Toxicity (Inhalation) Category 4, Skin Corrosion/Irritation Category 2, Serious Eye Damage/Eye Irritation Category 2; H302, H332, H315, H319 ^[2]	oral: ATE = 1200 mg/kg bw	Not Available
Legen	d: 1. Classified	by Chemwatch; 2. Classification	drawn from Regulation (EU) No 1272/200 dentified as baving endocrine disrupting pi	08 - Annex VI; 3. Class	sification drawn from

SECTION 4 First aid measures

4.1. Description of first aid measures If this product comes in contact with the eyes: Immediately hold eyelids apart and flush the eye continuously with running water. • Ensure complete irrigation of the eye by keeping eyelids apart and away from eye and moving the eyelids by occasionally Eve Contact lifting the upper and lower lids. Continue flushing until advised to stop by the Poisons Information Centre or a doctor, or for at least 15 minutes. Transport to hospital or doctor without delay. ▶ Removal of contact lenses after an eye injury should only be undertaken by skilled personnel. If skin or hair contact occurs: Immediately flush body and clothes with large amounts of water, using safety shower if available. Skin Contact Quickly remove all contaminated clothing, including footwear. • Wash skin and hair with running water. Continue flushing with water until advised to stop by the Poisons Information Centre. Transport to hospital, or doctor. ▶ If fumes or combustion products are inhaled remove from contaminated area. Lay patient down. Keep warm and rested. Prostheses such as false teeth, which may block airway, should be removed, where possible, prior to initiating first aid procedures. Apply artificial respiration if not breathing, preferably with a demand valve resuscitator, bag-valve mask device, or pocket mask as trained. Perform CPR if necessary. Transport to hospital, or doctor, without delay. Inhalation Inhalation of vapours or aerosols (mists, fumes) may cause lung oedema. Corrosive substances may cause lung damage (e.g. lung oedema, fluid in the lungs). + As this reaction may be delayed up to 24 hours after exposure, affected individuals need complete rest (preferably in semi-recumbent posture) and must be kept under medical observation even if no symptoms are (vet) manifested. Before any such manifestation, the administration of a spray containing a dexamethasone derivative or beclomethasone derivative may be considered. This must definitely be left to a doctor or person authorised by him/her. (ICSC13719) ▶ For advice, contact a Poisons Information Centre or a doctor at once. Urgent hospital treatment is likely to be needed. If swallowed do NOT induce vomiting. Ingestion • If vomiting occurs, lean patient forward or place on left side (head-down position, if possible) to maintain open airway and prevent aspiration. Observe the patient carefully.

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

Any material aspirated during vomiting may produce lung injury. Therefore emesis should not be induced mechanically or pharmacologically. Mechanical means should be used if it is considered necessary to evacuate the stomach contents; these include gastric lavage after endotracheal intubation. If spontaneous vomiting has occurred after ingestion, the patient should be monitored for difficult breathing, as adverse effects of aspiration into the lungs may be delayed up to 48 hours. For acute or short term repeated exposures to ethylene glycol:

- Early treatment of ingestion is important. Ensure emesis is satisfactory.
- Test and correct for metabolic acidosis and hypocalcaemia.
- Apply sustained diuresis when possible with hypertonic mannitol.
- Evaluate renal status and begin haemodialysis if indicated. [I.L.O]
- Rapid absorption is an indication that emesis or lavage is effective only in the first few hours. Cathartics and charcoal are generally not effective.
- Correct acidosis, fluid/electrolyte balance and respiratory depression in the usual manner. Systemic acidosis (below 7.2) can be treated with intravenous sodium bicarbonate solution.
- Ethanol therapy prolongs the half-life of ethylene glycol and reduces the formation of toxic metabolites.
- Pyridoxine and thiamine are cofactors for ethylene glycol metabolism and should be given (50 to 100 mg respectively) intramuscularly, four times per day for 2 days
- * Magnesium is also a cofactor and should be replenished. The status of 4-methylpyrazole, in the treatment regime, is still uncertain. For clearance of the material and its metabolites, haemodialysis is much superior to peritoneal dialysis.

[Ellenhorn and Barceloux: Medical Toxicology]

It has been suggested that there is a need for establishing a new biological exposure limit before a workshift that is clearly below 100 mmol ethoxy-acetic acids per mole creatinine in morning urine of people occupationally exposed to ethylene glycol ethers. This arises from the finding that an increase in urinary stones may be associated with such exposures.

Laitinen J., et al: Occupational & Environmental Medicine 1996; 53, 595-600

For acute or short-term repeated exposures to highly alkaline materials:

- Respiratory stress is uncommon but present occasionally because of soft tissue edema.
- Unless endotracheal intubation can be accomplished under direct vision, cricothyroidotomy or tracheotomy may be necessary.
- Oxygen is given as indicated
- The presence of shock suggests perforation and mandates an intravenous line and fluid administration.
- Damage due to alkaline corrosives occurs by liquefaction necrosis whereby the saponification of fats and solubilisation of proteins allow deep penetration into the tissue

Alkalis continue to cause damage after exposure.

INGESTION:

Milk and water are the preferred diluents

No more than 2 glasses of water should be given to an adult.

- Neutralising agents should never be given since exothermic heat reaction may compound injury.
- * Catharsis and emesis are absolutely contra-indicated.
- * Activated charcoal does not absorb alkali.
- * Gastric lavage should not be used.

Supportive care involves the following:

- Withhold oral feedings initially.
- If endoscopy confirms transmucosal injury start steroids only within the first 48 hours.
- Carefully evaluate the amount of tissue necrosis before assessing the need for surgical intervention.
- Patients should be instructed to seek medical attention whenever they develop difficulty in swallowing (dysphagia).

SKIN AND EYE:

Injury should be irrigated for 20-30 minutes.

Eye injuries require saline. [Ellenhorn & Barceloux: Medical Toxicology]

SECTION 5 Firefighting measures

5.1. Extinguishing media

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

5.2. Special hazards arising from the substrate or mixture

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Continued...

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 firefighter	S
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 fire fighting procedures suitable for surrounding area. Do not approach containers suspected to be hot. Cool fire exposed containers with water spray from a protected location. If safe to do so, remove containers from path of fire.
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. May emit corrosive fumes.

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	 Drains for storage or use areas should have retention basins for pH adjustments and dilution of spills before discharge or disposal of material. Check regularly for spills and leaks. 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	Chemical Class: bases For release onto land: recommended SORBENT RANK APPLICA TYPE RANK APPLICA LAND SPILL - SMALL cross-linked polymer - particulate cross-linked polymer - pillow sorbent clay - particulate foamed glass - pillow expanded minerals - particulate foamed glass - particulate foamed glass - particulate cross-linked polymer - particulate sorbent clay - particulate cross-linked polymer - particulate expanded minerals - particulate cross-linked polymer - particulate sorbent clay - particulate sorbent clay - particulate expanded mineral - particulate sorbent clay - particulate expanded mineral - particulate foamed glass - particulate cross-linked polymer - pillow	1 sorb TION 1 2 2 3 4 1 2 3 3 4	ents listed COLLI shovel throw shovel throw shovel shovel blower blower blower throw	in order of ECTION shovel pitchfork shovel pitchfork shovel skipload skipload skipload	i priority. LIMITATIONS R,W,SS R, DGC, RT R, I, P K, R, P, DGC, RT R, I, W, P, DGC R, W, SS er R, I, P er R, W, SS er R, I, W, P, DGC er R, I, W, P, DGC er R, I, W, P, DGC er R, W, SS er R, I, W, P, DGC er R, U, P, DGC	
	foamed glass - pillow Legend	4	throw	skipload	er R, P, DGC., RT	

DO	GC: Not effective where ground cover is dense
R;	Not reusable
	Not incinerable
P:	Effectiveness reduced when rainy
R	CNot effective where terrain is runged
S	Not for use within environmentally sensitive sites
W	- Ffectiveness reduced when windy
R	- Encourding of the second which which will be and the second of the second s
	W Maked at al. Bollution Tabralagy Barjaw No. 150-100 and Comparision 1098
N	w weivold et al. Politicion Technology Review No. 150. Noves Data Colporation 1988
•	Clear area of personnel and move upwind.
•	· 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.
	Stop leak if safe to do so.
	Contain spill with sand, earth or vermiculite.
	Collect recoverable product into labelled containers for recycling.

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. WARNING: To avoid violent reaction, ALWAYS add material to water and NEVER water to material. Avoid smoking, naked lights or ignition sources. Avoid contact with incompatible materials. When handling, DO NOT eat, drink or smoke.
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. DO NOT store near acids, or oxidising agents No smoking, naked lights, heat or ignition sources.

7.2. Conditions for safe storage, including any incompatibilities

Suitable container	 Lined metal can, lined metal pail/ can. Plastic pail. Polyliner drum. Packing as recommended by manufacturer. Check all containers are clearly labelled and free from leaks. For low viscosity materials Drums and jerricans must be of the non-removable head type. Where a can is to be used as an inner package, the can must have a screwed enclosure. For materials with a viscosity of at least 2680 cSt. (23 deg. C) and solids (between 15 C deg. and 40 deg C.): Removable head packaging; Cans with friction closures and low pressure tubes and cartridges may be used. Where combination packages are used, and the inner packages are of glass, porcelain or stoneware, there must be sufficient inert cushioning material in contact with inner and outer packages unless the outer packaging is a close fitting moulded plastic box and the substances are not incompatible with the plastic.
Storage incompatibility	 Avoid strong acids, acid chlorides, acid anhydrides and chloroformates. Avoid contact with copper, aluminium and their alloys. Avoid reaction with oxidising agents
Hazard categories in accordance with Regulation (EC) No 1272/2008	Not Available



X — Must not be stored together

0 — May be stored together with specific preventions

+ — May be stored together

Note: Depending on other risk factors, compatibility assessment based on the table above may not be relevant to storage situations, particularly where large volumes of dangerous goods are stored and handled. Reference should be made to the Safety Data Sheets for each substance or article and risks assessed accordingly.

7.3. Specific end use(s)

See section 1.2

SECTION 8 Exposure controls / personal protection

8.1. Control parameters

Ingredient	DNELs Exposure Pattern Worker	PNECs Compartment
potassium hydroxide	Inhalation 1 mg/m ³ (Local, Chronic) Inhalation 1 mg/m ³ (Local, Chronic) *	Not Available
sodium metasilicate, pentahydrate	Dermal 1.49 mg/kg bw/day (Systemic, Chronic) Inhalation 6.22 mg/m ³ (Systemic, Chronic) Inhalation 2 mg/m ³ (Local, Chronic) Inhalation 2 mg/m ³ (Local, Acute) Dermal 0.74 mg/kg bw/day (Systemic, Chronic) * Inhalation 1.55 mg/m ³ (Systemic, Chronic) * Oral 0.74 mg/kg bw/day (Systemic, Chronic) *	7.5 mg/L (Water (Fresh)) 1 mg/L (Water - Intermittent release) 7.5 mg/L (Water (Marine)) 1000 mg/L (STP)
sodium linear-(C12-14)alkyl ether sulfate	Dermal 2 750 mg/kg bw/day (Systemic, Chronic) Inhalation 7.9 mg/m ³ (Systemic, Chronic) Dermal 132 μg/cm ² (Local, Chronic) Dermal 1 650 mg/kg bw/day (Systemic, Chronic) * Inhalation 1.4 mg/m ³ (Systemic, Chronic) * Oral 1.125 mg/kg bw/day (Systemic, Chronic) * Dermal 79 μg/cm ² (Local, Chronic) *	0.052 mg/L (Water (Fresh)) 0.024 mg/L (Water - Intermittent release) 0.071 mg/L (Water (Marine)) 0.2 mg/kg sediment dw (Sediment (Fresh Water)) 0.02 mg/kg sediment dw (Sediment (Marine)) 7.5 mg/kg soil dw (Soil) 1 g/L (STP)
cocamide diethanolamide.	Dermal 0.75 mg/kg bw/day (Systemic, Chronic) Inhalation 11.5 mg/m ³ (Systemic, Chronic) Dermal 89.3 µg/kg bw/day (Systemic, Chronic) * Inhalation 2.03 mg/m ³ (Systemic, Chronic) * Oral 1.17 mg/kg bw/day (Systemic, Chronic) *	 7 μg/L (Water (Fresh)) 0.7 μg/L (Water - Intermittent release) 24 μg/L (Water (Marine)) 0.23 mg/kg sediment dw (Sediment (Fresh Water)) 23 μg/kg sediment dw (Sediment (Marine)) 32 mg/kg soil dw (Soil) 830 mg/L (STP)
N,N-dimethyldecanamide	Dermal 23.81 mg/kg bw/day (Systemic, Chronic) Inhalation 166.67 mg/m ³ (Systemic, Chronic) Dermal 14.29 mg/kg bw/day (Systemic, Chronic) * Inhalation 50 mg/m ³ (Systemic, Chronic) * Oral 14.29 mg/kg bw/day (Systemic, Chronic) *	28 μg/L (Water (Fresh)) 2.8 μg/L (Water - Intermittent release) 77 μg/L (Water (Marine)) 1.58 mg/kg sediment dw (Sediment (Fresh Water)) 0.158 mg/kg sediment dw (Sediment (Marine)) 10.6 mg/kg soil dw (Soil) 2.12 mg/L (STP) 12.71 mg/kg food (Oral)
ethylene glycol monobutyl ether	Inhalation 98 mg/m ³ (Systemic, Chronic) Inhalation 1 091 mg/m ³ (Systemic, Acute) Inhalation 246 mg/m ³ (Local, Acute) Inhalation 59 mg/m ³ (Systemic, Chronic) * Oral 6.3 mg/kg bw/day (Systemic, Acute) * Inhalation 426 mg/m ³ (Systemic, Acute) * Oral 26.7 mg/kg bw/day (Systemic, Acute) * Inhalation 147 mg/m ³ (Local, Acute) *	 8.8 mg/L (Water (Fresh)) 0.88 mg/L (Water - Intermittent release) 26.4 mg/L (Water (Marine)) 34.6 mg/kg sediment dw (Sediment (Fresh Water)) 3.46 mg/kg sediment dw (Sediment (Marine)) 2.33 mg/kg soil dw (Soil) 463 mg/L (STP) 0.02 g/kg food (Oral)

* Values for General Population

Occupational Exposure Limits (OEL)

INGREDIENT DATA

Source	Ingredient	Material name	TWA	STEL	Peak	Notes
Norway regulations on action rvalues cand limif values physical and chemical factors in the work environment and infection risk groups for biological factors (Norwegian)	potassium hydroxide	Kaliumhydroksid	Not Available	Not Available	2 mg/m3	Not Available
EU Consolidated List of Indicative Occupational Exposure Limit Values (IOELVs)	ethylene glycol monobutyl ether	2-Butoxyethanol	20 ppm / 98 mg/m3	246 mg/m3 / 50 ppm	Not Available	Skin
Norway regulations on action rvalues cand limit values physical and chemical factors in the work environment and infection risk groups for biological factors (Norwegian)	ethylene glycol monobutyl ether	2-butoksyetanol	10 ppm / 50 mg/m3	Not Available	Not Available	HE

Emergency Limits

Ingredient	TEEL-1	TEEL-2	TEEL-3
potassium hydroxide	0.18 mg/m3	2 mg/m3	54 mg/m3
sodium metasilicate, pentahydrate	6.6 mg/m3	73 mg/m3	440 mg/m3
sodium metasilicate, pentahydrate	3.8 mg/m3	42 mg/m3	250 mg/m3
ethylene glycol monobutyl ether	60 ppm	120 ppm	700 ppm

Ingredient	Original IDLH	Revised IDLH
potassium hydroxide	Not Available	Not Available
sodium metasilicate, pentahydrate	Not Available	Not Available
C12-14- alkyl(hydroxyethyl)dimethyl chloride, ethoxylated	Not Available	Not Available
Fatty alcohol ethoxylates*	Not Available	Not Available
sodium linear-(C12-14)alkyl ether sulfate	Not Available	Not Available
cocamide diethanolamide.	Not Available	Not Available
N,N-dimethyldecanamide	Not Available	Not Available
hydrocarbons, C10-13- n-alkanes, isoalkanes, cyclics, < 2% aromatics	Not Available	Not Available
ethylene glycol monobutyl ether	700 ppm	Not Available

Occupational Exposure Banding

Ingredient	Occupational Exposure Band Rating	Occupational Exposure Band Limit
sodium metasilicate, pentahydrate	E	≤ 0.01 mg/m³
C12-14- alkyl(hydroxyethyl)dimethyl chloride, ethoxylated	E	≤ 0.01 mg/m³
Fatty alcohol ethoxylates*	E	≤ 0.1 ppm
Notes:	Occupational exposure banding is a process of assigning chemicals into specific categories or bands based on a chemical's potency and the adverse health outcomes associated with exposure. The output of this process is an occupational exposure band (OEB), which corresponds to a range of exposure concentrations that are expected to protect worker health.	

Continued...

Ingredient	Occupational Exposure Band Rating	Occupational Exposure Band Limit
sodium linear-(C12-14)alkyl ether sulfate	E	≤ 0.01 mg/m³
cocamide diethanolamide.	E	≤ 0.1 ppm
N,N-dimethyldecanamide	E	≤ 0.01 mg/m³
Notes:	Occupational exposure banding is a process of assigning chemicals into specific categories or bands based on a chemical's potency and the adverse health outcomes associated with exposure. The output of this process is an occupational exposure band (OEB), which corresponds to a range of exposure concentrations that are expected to protect worker health.	

MATERIAL DATA

for potassium hydroxide:

The TLV-TWA is protective against respiratory tract irritation produced at higher concentrations

These exposure guidelines have been derived from a screening level of risk assessment and should not be construed as unequivocally safe limits. ORGS represent an 8-hour time-weighted average unless specified otherwise.

CR = Cancer Risk/10000; UF = Uncertainty factor:

TLV believed to be adequate to protect reproductive health:

LOD: Limit of detection

Toxic endpoints have also been identified as:

D = Developmental; R = Reproductive; TC = Transplacental carcinogen

Jankovic J., Drake F.: A Screening Method for Occupational Reproductive

American Industrial Hygiene Association Journal 57: 641-649 (1996)

For ethylene glycol monobutyl ether (2-butoxyethanol)

Odour Threshold Value: 0.10 ppm (detection), 0.35 ppm (recognition)

Although rats appear to be more susceptible than other animals anaemia is not uncommon amongst humans following exposure. The TLV reflects the need to maintain exposures below levels found to cause blood changes in experimental animals. It is concluded that this limit will reduce the significant risk of irritation, haematologic effects and other systemic effects observed in humans and animals exposed to higher vapour concentrations. The toxic effects typical of some other glycol ethers (pancytopenia, testis atrophy and teratogenic effects) are not found with this substance.

Odour Safety Factor (OSF)

OSF=2E2 (2-BUTOXYETHANOL)

8.2. Exposure controls

8.2.1. Appropriate engineering controls	Engineering controls are used to remove a hazard or place a barrier between the worker and the hazard. Well-designed engineering controls can be highly effective in protecting workers and will typically be independent of worker interactions to provide this high level of protection. The basic types of engineering controls are: Process controls which involve changing the way a job activity or process is done to reduce the risk. Enclosure and/or isolation of emission source which keeps a selected hazard "physically" away from the worker and ventilation that strategically "adds" and "removes" air in the work environment. Ventilation can remove or dilute an air contaminant if designed properly. The design of a ventilation system must match the particular process and chemical or contaminant in use. Employers may need to use multiple types of controls to prevent employee overexposure.
8.2.2. Individual protection measures, such as personal protective equipment	
Eye and face protection	 Safety glasses with unperforated side shields may be used where continuous eye protection is desirable, as in laboratories; spectacles are not sufficient where complete eye protection is needed such as when handling bulk-quantities, where there is a danger of splashing, or if the material may be under pressure. Chemical goggles. Whenever there is a danger of the material coming in contact with the eyes; goggles must be properly fitted. [AS/NZS 1337.1, EN166 or national equivalent] Full face shield (20 cm, 8 in minimum) may be required for supplementary but never for primary protection of eyes; these afford face protection. Alternatively a gas mask may replace splash goggles and face shields. Contact lenses may pose a special hazard; soft contact lenses may absorb and concentrate irritants. A written policy document, describing the wearing of lenses or restrictions on use, should be created for each workplace or task.
Skin protection	See Hand protection below
Hands/feet protection	 Elbow length PVC gloves When handling corrosive liquids, wear trousers or overalls outside of boots, to avoid spills entering boots.
Body protection	See Other protection below
Other protection	 Overalls. PVC Apron. PVC protective suit may be required if exposure severe. Eyewash unit. Ensure there is ready access to a safety shower.

GLOVE SELECTION INDEX

Glove selection is based on a modified presentation of the:

"Forsberg Clothing Performance Index".

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

CargoClean HD

Material	СРІ
BUTYL	А
NEOPRENE	В
NITRILE	В
PVC	В
NAT+NEOPR+NITRILE	С
NATURAL RUBBER	С
NATURAL+NEOPRENE	С
NITRILE+PVC	С
PE/EVAL/PE	С
PVA	С
SARANEX-23	С

* CPI - Chemwatch Performance Index

A: Best Selection

B: Satisfactory; may degrade after 4 hours continuous immersion

C: Poor to Dangerous Choice for other than short term immersion

NOTE: As a series of factors will influence the actual performance of the glove, a final selection must be based on detailed observation. -

* Where the glove is to be used on a short term, casual or infrequent basis,

factors such as "feel" or convenience (e.g. disposability), may dictate a choice of gloves which might otherwise be unsuitable following long-term or frequent use. A qualified practitioner should be consulted.

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

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

Required Minimum Protection Factor	Half-Face Respirator	Full-Face Respirator	Powered Air Respirator
up to 10 x ES	AK-AUS P2	-	AK-PAPR-AUS / Class 1 P2
up to 50 x ES	-	AK-AUS / Class 1 P2	-
up to 100 x ES	-	AK-2 P2	AK-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 point organic compounds(below 65 degC)

- Cartridge respirators should never be used for emergency ingress or in areas of unknown vapour concentrations or oxygen content.
- The wearer must be warned to leave the contaminated area immediately on detecting any odours through the respirator. The odour may indicate that the mask is not functioning properly, that the vapour concentration is too high, or that the mask is not properly fitted. Because of these limitations, only restricted use of cartridge respirators is considered appropriate.
- Cartridge performance is affected by humidity. Cartridges should be changed after 2 hr of continuous use unless it is determined that the humidity is less than 75%, in which case, cartridges can be used for 4 hr. Used cartridges should be discarded daily, regardless of the length of time used

76ak-p()

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	Yellow		
Physical state	Liquid	Relative density (Water = 1)	1.030-1.055
Odour	Not Available	Partition coefficient n-octanol / water	Not Available
Odour threshold	Not Available	Auto-ignition temperature (°C)	Not Available
pH (as supplied)	>13	Decomposition temperature (°C)	Not Available
Melting point / freezing point (°C)	Not Available	Viscosity (cSt)	Not Available
Initial boiling point and boiling range (°C)	100	Molecular weight (g/mol)	Not Available
Flash point (°C)	Not Available	Taste	Not Available
Evaporation rate	Not Available	Explosive properties	Not Available
Flammability	Not Available	Oxidising properties	Not Available
Upper Explosive Limit (%)	Not Available	Surface Tension (dyn/cm or mN/m)	Not Available
Lower Explosive Limit (%)	Not Available	Volatile Component (%vol)	Not Available
Vapour pressure (kPa)	Not Available	Gas group	Not Available

Solubility in water	Miscible	pH as a solution (1%)	Not Available
Vapour density (Air = 1)	Not Available	VOC g/L	Not Available
Nanoform Solubility	Not Available	Nanoform Particle Characteristics	Not Available
Particle Size	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 hazard classes as defined in Regulation (EC) No 1272/2008

Inhaled	Inhalation of vapours or aerosols (mists, fumes), generated by the material during the course of normal handling, may produce toxic effects. Evidence shows, or practical experience predicts, that the material produces irritation of the respiratory system, in a substantial number of individuals, following inhalation. In contrast to most organs, the lung is able to respond to a chemical insult by first removing or neutralising the irritant and then repairing the damage. The repair process, which initially evolved to protect mammalian lungs from foreign matter and antigens, may however, produce further lung damage resulting in the impairment of gas exchange, the primary function of the lungs. Respiratory tract irritation often results in an inflammatory response involving the recruitment and activation of many cell types, mainly derived from the vascular system. Inhalation of alkaline corrosives may produce irritation of the respiratory tract with coughing, choking, pain and mucous membrane damage. Pulmonary oedema may develop in more severe cases; this may be immediate or in most cases following a latent period of 5-72 hours. Symptoms may include a tightness in the chest, dyspnoea, frothy sputum, cyanosis and dizziness. Findings may include hypotension, a weak and rapid pulse and moist rales.
Ingestion	Ingestion of alkaline corrosives may produce immediate pain, and circumoral burns. Mucous membrane corrosive damage is characterised by a white appearance and soapy feel; this may then become brown, oedematous and ulcerated. Profuse salivation with an inability to swallow or speak may also result. Even where there is limited or no evidence of chemical burns, both the oesophagus and stomach may experience a burning pain; vomiting and diarrhoea may follow. The vomitus may be thick and may be slimy (mucous) and may eventually contain blood and shreds of mucosa. Epiglottal oedema may result in respiratory distress and asphyxia. Marked hypotension is symptomatic of shock; a weak and rapid pulse, shallow respiration and clammy skin may also be evident. Swallowing of the liquid may cause aspiration of vomit into the lungs with the risk of haemorrhaging, pulmonary oedema, progressing to chemical pneumonitis; serious consequences may result. Signs and symptoms of chemical (aspiration) pneumonitis may include coughing, gasping, choking, burning of the mouth, difficult breathing, and bluish coloured skin (cyanosis). The material is not thought to produce adverse health effects following ingestion (as classified by EC Directives using animal models). Nevertheless, adverse systemic effects have been produced following exposure of animals by at least one other route and good hygiene practice requires that exposure be kept to a minimum.
Skin Contact	The material can produce severe chemical burns following direct contact with the skin. Skin contact is not thought to produce harmful health effects (as classified under EC Directives using animal models). Systemic harm, however, has been identified following exposure of animals by at least one other route and the material may still produce health damage following entry through wounds, lesions or abrasions. Good hygiene practice requires that exposure be kept to a minimum and that suitable gloves be used in an occupational setting. Skin contact with alkaline corrosives may produce severe pain and burns; brownish stains may develop. The corroded area may be soft, gelatinous and necrotic; tissue destruction may be deep. Anionic surfactants/ hydrotropes generally produce skin reactions following the removal of natural oils. The skin may appear red and may become sore. Papular dermatitis may also develop. Sensitive individuals may exhibit cracking, scaling and blistering. 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	Direct contact with alkaline corrosives may produce pain and burns. Oedema, destruction of the epithelium, corneal opacification and iritis may occur. In less severe cases these symptoms tend to resolve. In severe injuries the full extent of the damage may not be immediately apparent with late complications comprising a persistent oedema, vascularisation and corneal scarring, permanent opacity, staphyloma, cataract, symblepharon and loss of sight. Direct eye contact with some concentrated anionic surfactants/ hydrotropes produces corneal damage, in some cases severe. Low concentrations may produce immediate discomfort, conjunctival hyperaemia, and oedema of the corneal epithelium. Healing may take several days. Temporary clouding of the cornea may occur.
Chronic	Repeated or prolonged exposure to corrosives may result in the erosion of teeth, inflammatory and ulcerative changes in the mouth and necrosis (rarely) of the jaw. Bronchial irritation, with cough, and frequent attacks of bronchial pneumonia may ensue. Gastrointestinal disturbances may also occur. Chronic exposures may result in dermatitis and/or conjunctivitis. Repeated or long-term occupational exposure is likely to produce cumulative health effects involving organs or biochemical systems. Long-term exposure to respiratory irritants may result in disease of the airways involving difficult breathing and related systemic problems. Exposure to the material may cause concerns for human fertility, generally on the basis that results in animal studies provide sufficient evidence to cause a strong suspicion of impaired fertility in the absence of toxic effects, or evidence of impaired fertility occurring at around the same dose levels as other toxic effects, but which are not a secondary non-specific consequence of other toxic effects. On the basis, primarily, of animal experiments, concern has been expressed that the material may produce carcinogenic or mutagenic effects; in respect of the available information, however, there presently exists inadequate data for making a satisfactory assessment.

	тохісіту	IRRITATION
CargoClean HD	Not Available	Not Available
	ΤΟΧΙΟΙΤΥ	IRRITATION
neteopium hudrovide	Oral (Rat) LD50: 273 mg/kg ^[2]	Eye (rabbit):1mg/24h rinse-moderate
potassium hydroxide		Skin (human): 50 mg/24h SEVERE
		Skin (rabbit): 50 mg/24h SEVERE
	ΤΟΧΙΟΙΤΥ	IRRITATION
sodium metasilicate, pentahydrate	Oral (Rat) LD50: 1153 mg/kg ^[2]	Skin (human): 250 mg/24h SEVERE
ponian, a are		Skin (rabbit): 250 mg/24h SEVERE
C12-14-	ΤΟΧΙΟΙΤΥ	IRRITATION
chloride, ethoxylated	Oral (Rat) LD50: >300 mg/kg ^[2]	Not Available
	ΤΟΧΙΟΙΤΥ	IRRITATION
Fatty alcohol ethoxylates*	Dermal (Other) LD50: >2000 mg/kg ^[2]	Not Available
	Oral (Rat) LD50: >300-2000 mg/kg ^[2]	
	ΤΟΧΙΟΙΤΥ	IRRITATION
sodium linear-(C12-14)alkyl ether sulfate	dermal (rat) LD50: >=2000 mg/kg ^[1]	Eye: adverse effect observed (irritating) ^[1]
	Oral (Rat) LD50: >540 mg/kg ^[1]	Skin: adverse effect observed (irritating) ^[1]
	ΤΟΧΙΟΙΤΥ	IRRITATION
cocamide diethanolamide.	dermal (rat) LD50: >2000 mg/kg ^[2]	Not Available
	Oral (Rat) LD50: >2000 mg/kg ^[1]	
	ΤΟΧΙΟΙΤΥ	IRRITATION
N N dimetholdssemaids	dermal (rat) LD50: >2000 mg/kg ^[2]	Not Available
N,N-dimethyidecanamide	Inhalation(Rat) LC50: >3.551 mg/L4h ^[1]	
	Oral (Rat) LD50: >2000 mg/kg ^[1]	
hydrocarbons, C10-13-	TOXICITY	IRRITATION
n-aikanes, isoaikanes, cyclics, < 2% aromatics	Not Available	Not Available
ethylene glycol monobutyl ether	ΤΟΧΙΟΙΤΥ	IRRITATION

Continued...

dermal (guinea pig) LD50: 210 mg/kg ^[2]	Eye (rabbit): 100 mg SEVERE * [Union Carbide]
Inhalation(Rat) LC50: 450 ppm4h ^[2]	Eye (rabbit): 100 mg/24h-moderate
Oral (Rat) LD50: 250 mg/kg ^[2]	Eye: adverse effect observed (irritating) ^[1]
	Skin (rabbit): 500 mg, open; mild
	Skin: adverse effect observed (irritating) ^[1]
	Skin: no adverse effect observed (not irritating) ^[1]

POTASSIUM HYDROXIDE	The material may produce moderate eye irritation leading to inflammation. Repeated or prolonged exposure to irritants may produce conjunctivitis. The material may produce severe skin irritation after prolonged or repeated exposure, and may produce a contact dermatitis (nonallergic). This form of dermatitis is often characterised by skin redness (erythema) thickening of the epidermis. Histologically there may be intercellular oedema of the spongy layer (spongiosis) and intracellular oedema of the epidermis. Prolonged contact is unlikely, given the severity of response, but repeated exposures may produce severe ulceration.			
SODIUM METASILICATE, PENTAHYDRATE	sodium metasilicate anhydrous: The material may be irritating to the eye, with prolonged contact causing inflammation. Repeated or prolonged exposure to irritants may produce conjunctivitis. The material may produce respiratory tract irritation. Symptoms of pulmonary irritation may include coughing, wheezing, laryngitis, shortness of breath, headache, nausea, and a burning sensation. Unlike most organs, the lung can respond to a chemical insult or a chemical agent, by first removing or neutralising the irritant and then repairing the damage (inflammation of the lungs may be a consequence). The repair process (which initially developed to protect mammalian lungs from foreign matter and antigens) may, however, cause further damage to the lungs (fibrosis for example) when activated by hazardous chemicals. Often, this results in an impairment of gas exchange, the primary function of the lungs. Therefore prolonged exposure to respiratory irritants may cause sustained breathing difficulties.			
C12-14- ALKYL(HYDROXYETHYL)DIMETHYL CHLORIDE, ETHOXYLATED	 * SDS for Berol R648 NG (70% aqueous solution) No specific data describing the health effects of cationic dialkyldimethylammonium (DADMA - dimonium) salts are readily available. However, many of the properties described for alkyltrimethylammonium (ATMA)) salts also apply to DADMA salts, although these are generally less irritating than the corresponding ATMA salts For alkyltrimethylammonium chloride (ATMAC) Most undiluted cationic surfactants satisfy the criteria for classification as Harmful (Xn) with R22 and as Irritant (Xi) for skin and eyes with R38 and R41. In addition, certain surfactants will satisfy the criteria for classification as Corrosive with R34 in addition to the acute toxicity. According to Centre Europeen des Agents de Surface et de leurs Intermediaires Organiques (CESIO), C8-18 alkyltrimethylammonium chloride (ATMAC) (i.e., lauryl, coco, soya, and tallow) are classified as Corrosive (C) with the risk phrases R22 (Harmful if swallowed) and R34 (Causes burns). C16 ATMAC is classified as Harmful (Xn) with the risk phrases R22 (Harmful if swallowed), R38 (Irritating to skin), and R41 (Risk of serious damage to eyes). C20-22 ATMAC are classified as Irritant (Xi) with R36/38 (Irritating to eyes and skin). Toxokinetics and Acute Toxicity: The few available absorption studies conducted with cationic surfactants indicate that absorption occurs in small amounts through the skin. Percutaneous absorption of radiolabelled C12 alkyltrimethylammonium bromide (ATMAB) in 3% aqueous solution (applied to an 8 cm2 area with occlusion) in the rat was low and corresponded to 0.6% of the applied 14C activity within the first 24 hours, whereas 13.2% remained on the skin after rinsing. Cutaneous application of the surfactant without rinsing resulted in a greater degree of percutaneous absorption (3.15%) in 48 hours. 			
Fatty alcohol ethoxylates*	Alcohol ethoxylates are according to CESIO (2000) classified as Irritant or Harmful depending on the number of EO-units: EO < 5 gives Irritant (Xi) with R38 (Irritating to skin) and R41 (Risk of serious damage to eyes) EO > 5-15 gives Harmful (Xn) with R22 (Harmful if swallowed) - R38/41 EO > 15-20 gives Harmful (Xn) with R22-41 >20 EO is not classified (CESIO 2000) Oxo-AE, C13 EO10 and C13 EO15, are Irritating (Xi) with R36/38 (Irritating to eyes and skin) . AE are not included in Annex 1 of the list of dangerous substances of the Council Directive 67/548/EEC In general, alcohol ethoxylates (AE) are readily absorbed through the skin of guinea pigs and rats and through the gastrointestinal mucosa of rats. AE are quickly eliminated from the body through the urine, faeces, and expired air (CO2).Orally dosed AE was absorbed rapidly and extensively in rats, and more than 75% of the dose was absorbed. When applied to the skin of humans, the doses were absorbed slowly and incompletely (50% absorbed in 72 hours). Half of the absorbed surfactant was excreted promptly in the urine and smaller amounts of AE appeared in the faeces and expired air (CO2)). The metabolism of C12 AE yields PEG, carboxylic acids, and CO2 as metabolites. The LD50 values after oral administration to rats range from about 1-15 g/kg body weight indicating a low to moderate acute			

	toxicity.
	The ability of nonionic surfactants to cause a swelling of the stratum corneum of guinea pig skin has been studied. The swelling mechanism of the skin involves a combination of ionic binding of the hydrophilic group as well as hydrophobic interactions of the alkyl chain with the substrate.
SODIUM LINEAR-(C12-14)ALKYL ETHER SULFATE	Alkyl ether sulfates (alcohol or alkyl ethoxysulfates) (AES) (syn: AAASD ,alkyl alcohol alkoxylate sulfates, SLES) are generally classified according to Comité Européen des Agents de Surface et leurs Intermédiaires Organiques (CESIO) as Irritant (Xi) with the risk phrases R38 (Irritating to skin) and R36 (Irritating to eyes). An exception has been made for AES (2-3E0) in a concentration of 70-75% where R36 is substituted with R41 (Risk of serious damage to eyes). AES are not included in Annex 1 of the list of dangerous substances of Council Directive 67/548/EEC. In assessing this family the Cosmetic Ingredient Review (CIR) Expert Panel recognized that most of the acute oral toxicity, dermal irritation and sensitization, subchronic and chronic oral toxicity, reproductive and developmental toxicity, carcinogenicity, and photosensitization studies have been conducted on ammonium laureth sulfate and sodium laureth sulfate. Sodium and ammonium laureth sulfate have not evoked adverse responses in any toxicological testing, including acute oral toxicity, sub-chronic and chronic oral toxicity, reproductive and develop-mental toxicity, carcinogenicity, and photosensitization studies. These data, however, are considered a sufficient basis for concluding that the other ingredients are safe in the practices of use and concentration described in the safety assessment because of the fundamental chemical similarities between them and because they all are chemically similar salts(salts are expected to be dissociated in any product formulation independent of where the salt is sodium, ammonium, magnesium, or zinc) of sulfated ethoxylated alcohols, and they all function as surfactants in cosmetic formulations. Based on these considerations, safety test data on one ingredient may be extrapolated to all of them. The panel noted that sodium laureth sulfate and ammonium laureth sulfate can produce eye and/or skin irritation in experimental animals and in some human test subjects; irritation may occur in some users of cosmetic formula
cocamide diethanolamide.	Coconut oil diethanolamine condensate is possibly carcinogenic to humans (IARC Group 2B) In a study of the dermal application in mice, coconut oil diethanolamine condensate increased the incidence of hepatocellular carcinoma and hepatocellular adenoma in males and females, and of hepatoblastoma in males. The incidence of renal tubule adenoma and carcinoma combined was also increased in males. In a study of dermal application in rats, no increase in tumour incidence was observed. Tumours of the kidney and hepatoblastoma are rare spontaneous neoplasms in experimental animals. The amide linkage between diethanolamine and of the fatty acid moiety is resistant to metabolic hydrolysis. The carcinogenic effects of the coconut diethanolamine condensate used in the cancer bioassay may be due to the levels of diethanolamine (18.2%) in the solutions tested. Mechanistic data are very weak to evaluate the carcinogenic potential of coconut oil diethanolamine condensate per se. A test material composed primarily of diethanolamides of coconut oil acids, with unreacted diethanolamine, alkanolamides of unsaturated acids, and amine salts of the acids, was evaluated. The polar nitrosamine, N-nitrosodiethanolamine, was detected at a concentration of 219 ppb Under test conditions, there was no evidence of carcinogenic activity of the test material in male rats administered 50 or 100 mg/kg bw. There was an equivocal evidence of carcinogenic activity in female rats based on a marginal increase in the incidences of renal tubule neoplasms. for diethanolamine (DEA): In animal studies, DEA has low acute toxicity via the oral and dermal routes with moderate skin irritation and severe eye irritation. In subchronic toxicity testing conducted via the oral route in rats and mice, the main effects observed were increased organ weights and histopathology of the kidney and/or liver, with the majority of other tissue effects noted only at relatively high dosages. In subchronic studies conducted via the dermal route, skin irrita
N,N-DIMETHYLDECANAMIDE	Toxicity test were performed with a mixture of N,N-dimethyldecanamide and N,N-dimethyloctanamide (with traces of N,N-dimethyl-dodecanamide and N,N-dimethyl-hexanamide). Due to the fact that a high amount in the mixture was N,N-dimethyloctanamide and the rest of the mixture are homologues with a lower and higher molecular weight which can be assumed to have a similar toxicological behaviour it is concluded that the mixture has nearly an similar toxicological behaviour like pure N,N-dimethyloctanamide. A 90 days repeated dose studies with a mixture of a mixture of N,N-dimethyldecanamide and N,N-dimethyloctanamide in beagle dogs via gavage (40, 200 and 1000 mg/kg bw/d) reported no relevant findings regarding the male or female fertility/developmental toxicity . It is assumed that a reproductive screening study or two generation study does not need to be conducted as results from a developmental toxicity study and a subchronic toxicity study did not reveal any reason of concern for offspring and for parent animals with respect to developmental toxicity or fertility. There were no hints for gene mutation or cytogenicity from in vitro genotoxicity test performed with the pure N,N-Dimethyloctanamide or from a mixture of N,N-Dimethyldecanamide

	and/or N,N-Dimethyloctanamide (with traces of N,N-dimethyldodecanamide and N,N-dimethylhexanamide). * REACh Dossier
HYDROCARBONS, C10-13- N-ALKANES, ISOALKANES, CYCLICS, < 2% AROMATICS	Studies indicate that normal, branched and cyclic paraffins are absorbed from the mammalian gastrointestinal tract and that the absorption of n-paraffins is inversely proportional to the carbon chain length, with little absorption above C30. With respect to the carbon chain lengths likely to be present in mineral oil, n-paraffins may be absorbed to a greater extent that iso- or cyclo-paraffins. The major classes of hydrocarbons have been shown to be well absorbed by the gastrointestinal tract in various species. In many cases, the hydrophobic hydrocarbons are ingested in association with dietary lipids. The dependence of hydrocarbon absorption on concomitant triglyceride digestion and absorption, is known as the "hydrocarbon continuum hypothesis", and asserts that a series of solubilising phases in the intestinal lumen, created by dietary triglycerides and their digestion products, afford hydrocarbons a route to the lipid phase of the intestinal absorptive cell (enterocyte) membrane. While some hydrocarbons may traverse the mucosal epithelium unmetabolised and appear as solutes in lipoprotein particles in intestinal lymph, there is evidence that most hydrocarbons partially separate from nutrient lipids and undergo metabolic transformation in the enterocyte. The enterocyte may play a major role in determining the proportion of an absorbed hydrocarbon that, by escaping initial biotransformation, becomes available for deposition in its unchanged form in peripheral tissues such as adipose tissue, or in the liver. For alkanes: Exposure to the commercial hexane (a representative of the ECHA group of hydrocarbons, C5-C7, n-alkanes, isoalkanes, n-hexane rich) had no effect on the behavior of rats. Rats were tested monthly throughout the exposure for hindlimb splay and grip strength. The NOAEC for sub-chronic neurological effects is 9000 ppm in rats. In a 13 week subchronic inhalation study, the neurotoxicity of light alkylate naphtha distillate (LAND-2; carbon range C5-C8) was examined in male and female rats and aside
ETHYLENE GLYCOL MONOBUTYL ETHER	NOTE: Changes in kidney, liver, spleen and lungs are observed in animals exposed to high concentrations of this substance by all routes. ** ASCC (NZ) SDS For ethylene glycol monoalkyl ethers and their acetates (EGMAEs): Typical members of this category are ethylene glycol propylene ether (EGPE), ethylene glycol butyl ether (EGBE) and ethylene glycol hexyl ether (EGHE) and their acetates. EGMAEs are substrates for alcohol dehydrogenase isozyme ADH-3, which catalyzes the conversion of their terminal alcohols to aldehydes (which are transient metabolites). Further, rapid conversion of the aldehydes by aldehyde dehydrogenase produces alkoxyacetic acids, which are the predominant urinary metabolites of mono substituted glycol ethers. Acute Toxicity: Oral LD50 values in rats for all category members range from 739 (EGHE) to 3089 mg/kg bw (EGPE), with values increasing with decreasing molecular weight. Four to six hour acute inhalation toxicity studies were conducted for these chemicals in rats at the highest vapour concentrations practically achievable. Values range from LC0 > 85 ppm (508 mg/m3) for EGHE, LC50 > 400ppm (2620 mg/m3) for EGBEA to LC50 > 2132 ppm (9061 mg/m3) for FGBE. No lethality was observed for any of these materials under these conditions. Dermal LD50 values in rabbits range from 435 mg/kg bw (EGBE) to 1500 mg/kg bw (EGBEA). Overall these category members can be considered to be of low to moderate acute toxicity. Exposure of pregnant rats to ethylene glycol monobutyl ether (2-butoxyethanol) at 100 ppm or rabbits at 200 ppm during organogenesis resulted in maternal toxicity and embryotoxicity including a decreased number of viable implantations per litter. Slight foetoxicity in the form of poorly ossified or unossified skeletal elements was also apparent in rats. Teratogenic effects were not observed in other species. At least one researcher has stated that the reproductive effects were less than that of other monoalkyl ethers of ethylene glycol. Chronic exposure may cause anaemia, macrocytosis, a
CargoClean HD & POTASSIUM HYDROXIDE & SODIUM METASILICATE, PENTAHYDRATE & C12-14- ALKYL(HYDROXYETHYL)DIMETHYL CHLORIDE, ETHOXYLATED & cocamide diethanolamide. & N,N-DIMETHYLDECANAMIDE	Asthma-like symptoms may continue for months or even years after exposure to the material ends. This may be due to a non-allergic condition known as reactive airways dysfunction syndrome (RADS) which can occur after exposure to high levels of highly irritating compound. Main criteria for diagnosing RADS include the absence of previous airways disease in a non-atopic individual, with sudden onset of persistent asthma-like symptoms within minutes to hours of a documented exposure to the irritant. Other criteria for diagnosis of RADS include a reversible airflow pattern on lung function tests, moderate to severe bronchial hyperreactivity on methacholine challenge testing, and the lack of minimal lymphocytic inflammation, without eosinophilia. RADS (or asthma) following an irritating inhalation is an infrequent disorder with rates related to the concentration of and duration of exposure to the irritating substance. On the other hand, industrial bronchitis is a disorder that occurs as a result of exposure due to high concentrations of irritating substance (often particles) and is completely reversible after exposure ceases. The disorder is characterized by difficulty breathing, cough and mucus production.

SODIUM METASILICATE, PENTAHYDRATE & cocamide diethanolamide. & ETHYLENE GLYCOL MONOBUTYL ETHER	The material may cause skin irritation after prolonged or repeated exposure and may produce a contact dermatitis (nonallergic). This form of dermatitis is often characterised by skin redness (erythema) and swelling epidermis. Histologically there may be intercellular oedema of the spongy layer (spongiosis) and intracellular oedema of the epidermis.
C12-14- ALKYL(HYDROXYETHYL)DIMETHYL CHLORIDE, ETHOXYLATED & SODIUM LINEAR-(C12-14)ALKYL ETHER SULFATE	Polyethers, for example, ethoxylated surfactants and polyethylene glycols, are highly susceptible towards air oxidation as the ether oxygens will stabilize intermediary radicals involved. Investigations of a chemically well-defined alcohol (pentaethylene glycol mono-n-dodecyl ether) ethoxylate, showed that polyethers form complex mixtures of oxidation products when exposed to air. Sensitization studies in guinea pigs revealed that the pure nonoxidized surfactant itself is nonsensitizing but that many of the investigated oxidation products are sensitizers. Two hydroperoxides were identified in the oxidation mixture, but only one (16-hydroperoxy-3,6,9,12,15-pentaoxaheptacosan-1-ol) was stable enough to be isolated. It was found to be a strong sensitizer in LLNA (local lymph node assay for detection of sensitization capacity). The formation of other hydroperoxides was indicated by the detection of their corresponding aldehydes in the oxidation mixture . On the basis of the lower irritancy, nonionic surfactants are often preferred to ionic surfactants in topical products. However, their susceptibility towards autoxidation also increases the irritation. Because of their irritating effect, it is difficult to diagnose ACD to these compounds by patch testing. Allergic Contact Dermatitis—Formation, Structural Requirements,and Reactivity of Skin Sensitizers. Ann-Therese Karlberg et al; Chem.
SODIUM LINEAR-(C12-14)ALKYL ETHER SULFATE & cocamide diethanolamide. & HYDROCARBONS, C10-13- N-ALKANES, ISOALKANES, CYCLICS, < 2% AROMATICS	No significant acute toxicological data identified in literature search.
cocamide diethanolamide. & N,N-DIMETHYLDECANAMIDE	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 uricaria or Quincke's oedema. The pathogenesis of contact eczema involves a cell-mediated (T pymptocytes) immune reaction of the delayed type. Other allergic skin reactions, e.g. contact uricaria, 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. Fatty acid amides (FAA) are ubiquitous in household and commercial environments. The most common of these are based on coconut oil fatty acids alkanolamides. These are the most widely studied in terms of human exposure. Fatty acid diethanolamides (CB-C18) are classified by Comite Europeen des Agents de Surface et de leurs Intermediaires Organiques (CESIO) as Irritating (X) with the risk phrases R38 (firitating to skin) and R41 (Risk of serious damage to eyes). Fatty acid monoethanolamides are classified as Irritant (Xi) with the risk phrases R41 Several studies of the sensitization potential of cocoamide diethanolamide (DEA) indicate that this FAA induces occupational allergic contact dematitis and a number of reports on skin allergy patch testing of cocaamide DEA have been published. These tests indicate that allergy to cocoamide DEA is becoming more common. Alkanolamides (especially secondary alkanolamides) are susceptible to nitrosamine formation which constitutes a potential health problem. Nitrosamine contamination is possible either from pre-existing contaminat

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CargoClean HD

Imidazole derivatives. For Subcategory IV, two subchronic toxicity studies for one of the chemicals indicated a low order of repeat-dose toxicity for the FND amphoteric salts similar to that seen in the other categories. Genetic Toxicity in vitro: Based on the lack of effect of one or more chemicals in each subcategory, adequate data for mutagenic activity as measured by the Salmonella reverse mutation assay exist for all of the subcategories. Developmental Toxicity: A developmental toxicity study in Subcategory I and in Subcategory IV and a third study for a chemical in Subcategory III are available. The studies indicate these chemicals are not developmental toxicants, as expected based on their structures, molecular weights, physical properties and knowledge of similar chemicals. cocamide diethanolamide. & ETHYLENE GLYCOL MONOBUTYL The material may produce severe irritation to the eye causing pronounced inflammation. Repeated or prolonged ETHER exposure to irritants may produce conjunctivitis. Acute Toxicity × × Carcinogenicity Skin Irritation/Corrosion × ~ Reproductivity Serious Eye STOT - Single Exposure × X Damage/Irritation

 Respiratory or Skin sensitisation
 ×
 STOT - Repeated Exposure

 Mutagenicity
 ×
 Aspiration Hazard

> Legend: X − Data either not available or does not fill the criteria for classification ✓ − Data available to make classification

X

×

11.2 Information on other hazards

11.2.1. Endocrine disrupting properties

Many chemicals may mimic or interfere with the body's hormones, known as the endocrine system. Endocrine disruptors are chemicals that can interfere with endocrine (or hormonal) systems. Endocrine disruptors interfere with the synthesis, secretion, transport, binding, action, or elimination of natural hormones in the body. Any system in the body controlled by hormones can be derailed by hormone disruptors. Specifically, endocrine disruptors may be associated with the development of learning disabilities, deformations of the body various cancers and sexual development problems. Endocrine disrupting chemicals cause adverse effects in animals. But limited scientific information exists on potential health problems in humans. Because people are typically exposed to multiple endocrine disruptors at the same time, assessing public health effects is difficult.

11.2.2. Other information

See Section 11.1

SECTION 12 Ecological information

12.1. Toxicity

	Endpoint	Test Duration (hr)	S	pecies		Value	Source
CargoClean HD	Not Available	Not Available	N	ot Available		Not Available	Not Available
	Endpoint	Test Duration (hr)		Species		Value	Source
potassium hydroxide	LC50	96h		Fish		80mg/l	2
	NOEC(ECx)	24h		Fish		28mg/l	2
sodium metasilicate, pentahydrate	Endpoint	Test Duration (hr)	Spe	cies	Valu	e	Source
	EC50	72h	Algae or other aquatic plants		207mg/l		2
	EC50	48h	Crustacea		22.94-49.01mg/l		4
	LC50	96h	Fish	Fish 1		ng/l	1
	EC50(ECx)	48h	Crus	stacea	22.9	4-49.01mg/l	4
	Endpoint	Test Duration (hr)	Sp	ecies	Va	lue	Source
C12-14- alkyl(hydroxyethyl)dimethyl chloride, ethoxylated	EC50	48h	Cru	ustacea	>1	-10mg/l	Not Available
	LC50	96h	Fis	h	>1	0-100mg/l	Not Available
	EC50(ECx)	48h	Cru	ustacea	>1	-10mg/l	Not Available
Fatty alcohol ethoxylates*	Endpoint	Test Duration (hr)	Spec	ies	V	alue	Source

	EC50	48	Crustacea Daphnia magna	>10-100mg/L	8
	EC50	72	Algae/Plant Pseudokirchneriella subcapitata(Algae)	>10-100mg/L	8
	LC50	96	Fish Other	>10-100mg/L	8
	Endpoint	Test Duration (hr)	Species	Value	Source
	EC50	72h	Algae or other aquatic plants	1.8mg/l	2
sodium linear-(C12-14)alkyl	EC50	48h	Crustacea	7.4mg/l	2
ether sulfate	EC50	96h	Algae or other aquatic plants	7.5mg/l	2
	LC50	96h	Fish	>1<10mg/l	2
	NOEC(ECx)	672h	Fish	0.14mg/l	2
	Endpoint	Test Duration (hr)	Species	Value	Source
	EC50	48h	Crustacea	~3.2mg/l	2
cocamide diethanolamide.	LC50	96h	Fish	~2.4mg/l	2
	NOEC(ECx)	504h	Crustacea	~0.1mg/l	2
	EC50	72h	Algae or other aquatic plants	~2.1mg/l	2
	Endpoint	Test Duration (hr)	Species	Value	Source
	EC50	72h	Algae or other aquatic plants	0.805mg/l	2
N,N-dimethyldecanamide	EC50	48h	Crustacea	0.29mg/l	2
	LC50	96h	Fish	>0.88mg/l	2
	NOEC(ECx)	504h	Crustacea	0.079mg/l	2
hydrocarbons, C10-13-	Endpoint	Test Duration (hr)	Species	Value	Source
n-alkanes, isoalkanes, cyclics, < 2% aromatics	Not Available	Not Available	Not Available	Not Available	Not Available
	Endpoint	Test Duration (hr)	Species	Value	Source
	EC50	72h	Algae or other aquatic plants	623mg/l	2
othulana aluan menakutut	EC50	48h	Crustacea	164mg/l	2
emylene glycol monobutyl ether	EC50	96h	Algae or other aquatic plants	720mg/l	2
	LC50	96h	Fish	1700mg/l	Not Available
	EC10(ECx)	48h	Crustacea	7.2mg/l	2
Legend:	Extracted from	1. IUCLID Toxicity Data 2. Euro	ope ECHA Registered Substances - Ecotoxicolog	ical Information - Aqua	atic Toxicity

Bioconcentration Data 7. METI (Japan) - Bioconcentration Data 8. Vendor Data

Toxic to aquatic organisms.

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 Surfactants: Kow cannot be easily determined due to hydrophilic/hydrophobic properties of the molecules in surfactants. BCF value: 1-350.

Aquatic Fate: Surfactants tend to accumulate at the interface of the air with water and are not extracted into one or the other liquid phases.

Terrestrial Fate: Anionic surfactants are not appreciably sorbed by inorganic solids. Cationic surfactants are strongly sorbed by solids, particularly clays. Significant sorption of anionic and non-ionic surfactants has been observed in activated sludge and organic river sediments. Surfactants have been shown to improve water infiltration into soils with moderate to severe hydrophobic or water-repellent properties.

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

Prevent, by any means available, spillage from entering drains or water courses.

DO NOT discharge into sewer or waterways.

12.2. Persistence and degradability

Ingredient	Persistence: Water/Soil	Persistence: Air
N,N-dimethyldecanamide	LOW	LOW
ethylene glycol monobutyl ether	LOW (Half-life = 56 days)	LOW (Half-life = 1.37 days)

12.3. Bioaccumulative potential

Ingredient	Bioaccumulation
N,N-dimethyldecanamide	LOW (LogKOW = 3.4438)
ethylene glycol monobutyl ether	LOW (BCF = 2.51)

12.4. Mobility in soil

Ingredient	Mobility
N,N-dimethyldecanamide	LOW (KOC = 1307)
ethylene glycol monobutyl ether	HIGH (KOC = 1)

12.5. Results of PBT and vPvB assessment

	Ρ	В	т	
Relevant available data	Not Available	Not Available	Not Ava	ailable
PBT	×	×	×	
vPvB	×	×	×	
PBT Criteria fulfilled?				No
vPvB			No	

12.6. Endocrine disrupting properties

The evidence linking adverse effects to endocrine disruptors is more compelling in the environment than it is in humans. Endocrine disruptors profoundly alter reproductive physiology of ecosystems and ultimately impact entire populations. Some endocrine-disrupting chemicals are slow to break down in the environment. That characteristic makes them potentially hazardous over long periods of time. Some well established adverse effects of endocrine disruptors in various wildlife species include eggshell-thinning, displayed of characteristics of the opposite sex and impaired reproductive development. Other adverse changes in wildlife species that have been suggested, but not proven include reproductive abnormalities, immune dysfunction and skeletal deformaties.

12.7. Other adverse effects

No evidence of ozone depleting properties were found in the current literature.

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. 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. Consult manufacturer for recycling options or consult local or regional waste management authority for disposal if no suitable treatment or disposal facility can be identified. Treat and neutralise at an approved treatment plant. Treatment should involve: Neutralisation with suitable dilute acid followed by: burial in a land-fill specifically licensed to accept chemical and / or pharmaceutical wastes or Incineration in a licensed apparatus (after admixture with suitable combustible material).
	 material). Decontaminate empty containers. Observe all label safeguards until containers are cleaned and destroyed.
Waste treatment options	Not Available
Sewage disposal options	Not Available

SECTION 14 Transport information

Marine Pollutant	NO

Land transport (ADR-RID)

14.1. UN number or ID number	1719				
14.2. UN proper shipping name	CAUSTIC ALKALI	CAUSTIC ALKALI LIQUID, N.O.S. (contains potassium hydroxide)			
14.3. Transport hazard class(es)	Class Subsidiary risk	8 Not Applicab	P		
14.4. Packing group					
14.5. Environmental hazard	Not Applicable				
	Hazard identifica	tion (Kemler)	80		
	Classification code		C5		
14.6. Special precautions	Hazard Label		8		
for user	Special provisions		274		
	Limited quantity		5 L		
	Tunnel Restrictic	on Code	3 (E)		

Air transport (ICAO-IATA / DGR)

14.1. UN number	1719				
14.2. UN proper shipping name	Caustic alkali liquid, n.o.s. * (contains potassium hydroxide)				
14.3 Transport bazard	ICAO/IATA Class 8				
class(es)	ICAO / IATA Subsidiary Hazard	ICAO / IATA Subsidiary Hazard Not Applicable			
	ERG Code	8L			
14.4. Packing group					
14.5. Environmental hazard	Not Applicable				
	Special provisions		A3 A803		
	Cargo Only Packing Instructions		856		
	Cargo Only Maximum Qty / Pack		60 L		
14.6. Special precautions for user	Passenger and Cargo Packing Ir	nstructions	852		
	Passenger and Cargo Maximum	Qty / Pack	5 L		
	Passenger and Cargo Limited Qu	uantity Packing Instructions	Y841		
	Passenger and Cargo Limited Ma	aximum Qty / Pack	1 L		

Sea transport (IMDG-Code / GGVSee)

14.1. UN number	1719		
14.2. UN proper shipping name	CAUSTIC ALKALI LIQUID, N.O.S. (contains potassium hydroxide)		
14.3. Transport hazard	IMDG Class 8		
class(es)	IMDG Subrisk Not Applicable		
14.4. Packing group	Ш		
14.5 Environmental hazard	Not Applicable		

	EMS Number	F-A, S-B
14.6. Special precautions	Special provisions	223 274
for user	Limited Quantities	5 L

Inland waterways transport (ADN)

14.1. UN number	1719			
14.2. UN proper shipping name	CAUSTIC ALKALI LIQUID, N.O.S. (contains potassium hydroxide)			
14.3. Transport hazard class(es)	8 Not Applicable			
14.4. Packing group	Ш			
14.5. Environmental hazard	Not Applicable			
	Classification code	C5		
	Special provisions	274		
14.6. Special precautions for user	Limited quantity	5 L		
	Equipment required	PP, EP		
	Fire cones number	0		

14.7. Maritime transport in bulk according to IMO instruments

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

Not Applicable

14.7.2. Transport in bulk in accordance with MARPOL Annex V and the IMSBC Code

Product name	Group
potassium hydroxide	Not Available
sodium metasilicate, pentahydrate	Not Available
C12-14- alkyl(hydroxyethyl)dimethyl chloride, ethoxylated	Not Available
Fatty alcohol ethoxylates*	Not Available
sodium linear-(C12-14)alkyl ether sulfate	Not Available
cocamide diethanolamide.	Not Available
N,N-dimethyldecanamide	Not Available
hydrocarbons, C10-13- n-alkanes, isoalkanes, cyclics, < 2% aromatics	Not Available
ethylene glycol monobutyl ether	Not Available

14.7.3. Transport in bulk in accordance with the IGC Code

Product name	Ship Type
potassium hydroxide	Not Available
sodium metasilicate, pentahydrate	Not Available
C12-14- alkyl(hydroxyethyl)dimethyl chloride, ethoxylated	Not Available
Fatty alcohol ethoxylates*	Not Available
sodium linear-(C12-14)alkyl ether sulfate	Not Available
cocamide diethanolamide.	Not Available

Product name	Ship Type
N,N-dimethyldecanamide	Not Available
hydrocarbons, C10-13- n-alkanes, isoalkanes, cyclics, < 2% aromatics	Not Available
ethylene glycol monobutyl ether	Not Available

SECTION 15 Regulatory information

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

Europe EC Inventory European Union - European Inventory of Existing Commercial Chemical Substances (EINECS) European Union (EU) Regulation (EC) No 1272/2008 on Classification, Labelling and Packaging of Substances and Mixtures - Annex VI Sodium metasilicate, pentahydrate is found on the following regulatory lists European Union - European Inventory of Existing Commercial Chemical factors in the work environment and infection risk groups for biological factors (Norwegian) Sodium metasilicate, pentahydrate is found on the following regulatory lists European Chemicals Agency (ECHA) Community Rolling Action Plan (CoRAP) List of Substances Europe EC Inventory European Union (EU) Regulation (EC) No 1272/2008 on Classification, Labelling and Packaging of Substances and Mixtures - Annex VI C12-14-alkyl(hydroxyethyl)dimethyl chloride, ethoxylated is found on the following regulatory lists
European Union - European Inventory of Existing Commercial Chemical Labelling and Packaging of Substances and Mixtures - Annex VI Substances (EINECS) dNorway regulations om action values and limit values for physical chemical factors in the work environment and infection risk groups for biological factors (Norwegian) sodium metasilicate, pentahydrate is found on the following regulatory lists EU European Chemicals Agency (ECHA) Community Rolling Action Plan (CoRAP) List of Substances European Union - European Inventory of Existing Commercial Chemical Substances (EINECS) Europe EC Inventory European Union (EU) Regulation (EC) No 1272/2008 on Classification, Labelling and Packaging of Substances and Mixtures - Annex VI C12-14-alkyl(hydroxyethyl)dimethyl chloride, ethoxylated is found on the following regulatory lists European Union (EU) Regulation (EC) No 1272/2008 on Classification, Labelling and Packaging of Substances and Mixtures - Annex VI
sodium metasilicate, pentahydrate is found on the following regulatory lists EU European Chemicals Agency (ECHA) Community Rolling Action Plan (CoRAP) List of Substances European Union - European Inventory of Existing Commercial Chemical Substances (EINECS) Europe EC Inventory European Union (EU) Regulation (EC) No 1272/2008 on Classification, Labelling and Packaging of Substances and Mixtures - Annex VI C12-14-alkyl(hydroxyethyl)dimethyl chloride, ethoxylated is found on the following regulatory lists
EU European Chemicals Agency (ECHA) Community Rolling Action Plan (CoRAP) List of Substances European Union - European Inventory of Existing Commercial Chemical Substances (EINECS) Europe EC Inventory European Union (EU) Regulation (EC) No 1272/2008 on Classification, Labelling and Packaging of Substances and Mixtures - Annex VI C12-14-alkyl(hydroxyethyl)dimethyl chloride, ethoxylated is found on the following regulatory lists
Europe EC Inventory European Union (EU) Regulation (EC) No 1272/2008 on Classification, Labelling and Packaging of Substances and Mixtures - Annex VI C12-14-alkyl(hydroxyethyl)dimethyl chloride, ethoxylated is found on the following regulatory lists
C12-14-alkyl(hydroxyethyl)dimethyl chloride, ethoxylated is found on the following regulatory lists
Net Appliceble
Not Applicable
Fatty alcohol ethoxylates* is found on the following regulatory lists
Not Applicable
sodium linear-(C12-14)alkyl ether sulfate is found on the following regulatory lists
Europe EC Inventory
cocamide diethanolamide. is found on the following regulatory lists
Europe EC Inventory European Union - European Inventory of Existing Commercial Chemical Substances (EINECS)
N,N-dimethyldecanamide is found on the following regulatory lists
Europe EC Inventory European Union - European Inventory of Existing Commercial Chemical Substances (EINECS)
hydrocarbons, C10-13- n-alkanes, isoalkanes, cyclics, < 2% aromatics is found on the following regulatory lists
Not Applicable
ethylene glycol monobutyl ether is found on the following regulatory lists
EU Consolidated List of Indicative Occupational Exposure Limit Values (IOELVs) European Union (EU) Regulation (EC) No 1272/2008 on Classification, Labelling and Packaging of Substances and Mixtures - Annex VI
Europe EC Inventory International Agency for Research on Cancer (IARC) - Agents Classified by
European Union - European Inventory of Existing Commercial Chemical the IARC Monographs - Not Classified as Carcinogenic
chemical factors in the work environment and infection risk groups for

This safety data sheet is in compliance with the following EU legislation and its adaptations - as far as applicable - : Directives 98/24/EC, - 92/85/EEC, - 94/33/EC, - 2008/98/EC, - 2010/75/EU; Commission Regulation (EU) 2020/878; Regulation (EC) No 1272/2008 as updated through ATPs.

Information according to 2012/18/EU (Seveso III):

Seveso Category	Not Available

15.2. Chemical safety assessment

No Chemical Safety Assessment has been carried out for this substance/mixture by the supplier.

ECHA SUMMARY

Ingredient	CAS number Index No		EC		Dossier
potassium hydroxide	1310-58-3 019-002-00-8		Not A		vailable
Harmonisation (C&L Inventory)	Hazard Class and Category Code(s)		Pictograms Signal Word Code(s)		Hazard Statement Code(s)
1	Acute Tox. 4; Skin Corr. 1A		GHS05; Dgr		H302; H314
2	Skin Corr. 1A; Met. Corr. 1; Acute Tox. 4; Eye Dam. 1; Acute Tox. 3; STOT SE 1; Asp. Tox. 1; Flam. Liq. 2; STOT SE 3; Acute Tox. 3; Aquatic Chronic 3; Expl. 1.1; STOT RE 1		GHS05; Dgr; GHS GHS06; GHS09; GHS01	S08;	H314; H290; H312; H318; H301; H370; H304; H317; H335; H332; H412; H201; H372

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

Ingredient	CAS number	Index No	ECHA Dossier
sodium metasilicate, pentahydrate	10213-79-3	014-010-00-8	Not Available

Harmonisation (C&L Inventory)	Hazard Class and Category Code(s)	Pictograms Signal Word Code(s)	Hazard Statement Code(s)
1	Skin Corr. 1B; STOT SE 3	GHS05; Dgr	H314; H335
2	Skin Corr. 1B; STOT SE 3; Met. Corr. 1; Acute Tox. 4; Eye Dam. 1	GHS05; Dgr	H314; H335; H290; H302; H318

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

Ingredient	CAS number	Index No	ECHA Dossier
C12-14- alkyl(hydroxyethyl)dimethyl chloride, ethoxylated	1554325-20-0	Not Available	Not Available

Harmonisation (C&L Inventory)	Hazard Class and Category Code(s)	Pictograms Signal Word Code(s)	Hazard Statement Code(s)
1	Acute Tox. 4; Skin Irrit. 2; Eye Dam. 1; Aquatic Acute 1	GHS05; GHS09; Dgr	H302; H315; H318; H400
2	Acute Tox. 4; Skin Irrit. 2; Eye Dam. 1; Aquatic Acute 1	GHS05; GHS09; Dgr	H302; H315; H318; H400

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

Ingredient	CAS number	Index No	ECHA Dossier
Fatty alcohol ethoxylates*	160875-66-1*	Not Available	Not Available

Harmonisation (C&L Inventory)	Hazard Class and Category Code(s)	Pictograms Signal Word Code(s)	Hazard Statement Code(s)
1	Acute Tox. 4; Eye Dam. 1	GHS05; Dgr	H302; H318
2	Acute Tox. 4; Eye Dam. 1; Skin Irrit. 2; STOT SE 3; Aquatic Chronic 2	GHS05; Dgr; GHS09	H302; H318; H315; H202; H335; H411

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

Ingredient	CAS number	Index No	ECHA Dossier
sodium linear-(C12-14)alkyl ether sulfate	68891-38-3	Not Available	Not Available

Harmonisation (C&L Inventory)	Hazard Class and Category Code(s)	Pictograms Signal Word Code(s)	Hazard Statement Code(s)
1	Skin Irrit. 2; Eye Dam. 1; Aquatic Chronic 3	GHS05; Dgr	H315; H318; H412
2	Skin Irrit. 2; Eye Dam. 1; Aquatic Chronic 2; Skin Sens. 1	GHS05; Dgr; GHS09	H315; H318; H411; H317
1	Acute Tox. 4; Eye Irrit. 2	GHS07; Wng	H302; H319
2	Acute Tox. 4; Skin Irrit. 2; Eye Dam. 1; Acute Tox. 3; Skin Sens. 1B; Flam. Sol. 1; STOT SE 3; Aquatic Chronic 2	GHS05; Dgr; GHS06; GHS02; GHS09	H302; H315; H318; H311; H317; H335; H228; H411

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

Ingredient	CAS number	Index No	ECHA Dossier
cocamide diethanolamide.	68155-07-7*	Not Available	Not Available

Harmonisation (C&L Inventory)	Hazard Class and Category Code(s)	Pictograms Signal Word Code(s)	Hazard Statement Code(s)
1	Skin Irrit. 2; Eye Dam. 1; Aquatic Chronic 2	GHS05; GHS09; Dgr	H315; H318; H411
2	Skin Irrit. 2; Eye Dam. 1; Aquatic Chronic 2; Repr. 2	GHS05; GHS09; Dgr; GHS08	H315; H318; H411; H361
Harmonisation Code 1 = The m	ost prevalent classification. Harmonisation Code 2 = The r	nost severe classification.	

Ingredient	CAS number	Index No	ECHA Dossier
N,N-dimethyldecanamide	14433-76-2	Not Available	Not Available

Harmonisation (C&L Inventory)	Hazard Class and Category Code(s)	Pictograms Signal Word Code(s)	Hazard Statement Code(s)
1	Skin Irrit. 2; Eye Irrit. 2; STOT SE 3; Aquatic Chronic 3	GHS07; Wng	H315; H319; H335; H412
2	Skin Irrit. 2; STOT SE 3; Aquatic Chronic 2; Eye Dam. 1; Acute Tox. 4; Repr. 2	GHS09; GHS05; Dgr	H315; H335; H411; H318

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

Ingredient	CAS number	Index No		ECHA	ECHA Dossier	
ethylene glycol monobutyl ether	111-76-2	603-014-00-0 N		Not A	ot Available	
Harmonisation (C&L Inventory)	Hazard Class and Category Code(s)		Pictograms Si Word Code(s)	gnal	Hazard Statement Code(s)	
1	Acute Tox. 4; Acute Tox. 4; Skin Irrit. 2; Eye Irrit. 2; Acute Tox. 4		GHS07; Wng		H302; H312; H315; H319; H332	
2	Skin Irrit. 2; Flam. Liq. 2; Skin Sens. 1; Aquatic Chronic 2; Acute Tox. 2; Acute Tox. 2; Repr. 2; STOT SE 1; STOT RE 2; Acute Tox. 3; Eye Dam. 1; Muta. 2; Carc. 2		GHS06; Dgr; GHS08; GHS09	5	H315; H310; H330; H361; H370; H373; H412; H301; H317; H318; H341; H351	

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

National Inventory Status

National Inventory	Status	
Australia - AIIC / Australia Non-Industrial Use	No (C12-14-alkyl(hydroxyethyl)dimethyl chloride, ethoxylated)	
Canada - DSL	No (C12-14-alkyl(hydroxyethyl)dimethyl chloride, ethoxylated; Fatty alcohol ethoxylates*; N,N-dimethyldecanamide)	
Canada - NDSL	No (potassium hydroxide; sodium metasilicate, pentahydrate; C12-14-alkyl(hydroxyethyl)dimethyl chloride, ethoxylated; Fatty alcohol ethoxylates*; sodium linear-(C12-14)alkyl ether sulfate; cocamide diethanolamide.; ethylene glycol monobutyl ether)	
China - IECSC	No (C12-14-alkyl(hydroxyethyl)dimethyl chloride, ethoxylated)	
Europe - EINEC / ELINCS / NLP	No (C12-14-alkyl(hydroxyethyl)dimethyl chloride, ethoxylated; Fatty alcohol ethoxylates*)	
Japan - ENCS	No (C12-14-alkyl(hydroxyethyl)dimethyl chloride, ethoxylated)	
Korea - KECI	No (C12-14-alkyl(hydroxyethyl)dimethyl chloride, ethoxylated)	
New Zealand - NZIoC	No (C12-14-alkyl(hydroxyethyl)dimethyl chloride, ethoxylated; cocamide diethanolamide.)	
Philippines - PICCS	No (C12-14-alkyl(hydroxyethyl)dimethyl chloride, ethoxylated; cocamide diethanolamide.; N,N-dimethyldecanamide)	
USA - TSCA	Yes	
Taiwan - TCSI	No (C12-14-alkyl(hydroxyethyl)dimethyl chloride, ethoxylated)	
Mexico - INSQ	No (C12-14-alkyl(hydroxyethyl)dimethyl chloride, ethoxylated; Fatty alcohol ethoxylates*; sodium linear-(C12-14)alkyl ether sulfate; cocamide diethanolamide.; N,N-dimethyldecanamide)	
Vietnam - NCI	No (C12-14-alkyl(hydroxyethyl)dimethyl chloride, ethoxylated)	
Russia - FBEPH	No (C12-14-alkyl(hydroxyethyl)dimethyl chloride, ethoxylated; Fatty alcohol ethoxylates*)	
Legend:	Yes = All CAS declared ingredients are on the inventory No = One or more of the CAS listed ingredients are not on the inventory. These ingredients may be exempt or will require registration.	

SECTION 16 Other information

Revision Date	28/02/2023
Initial Date	03/05/2019

CONTACT POINT

- For quotations contact your local Customer Services - http://wssdirectory.wilhelmsen.com/#/customerservices - - Responsible for safety data sheet Wilhelmsen Ships Service AS - Prepared by: Product HSE Manager, - Email: Email: WSS.GLOBAL.SDSINFO@wilhelmsen.com - Telephone: Tel.: +31 10 4877775

Full text Risk and Hazard codes

H201	Explosive; mass explosion hazard.		
H202	Explosive, severe projection hazard.		
H226	Flammable liquid and vapour.		
H228	Flammable solid.		
H301	Toxic if swallowed.		
H302	Harmful if swallowed.		
H304	May be fatal if swallowed and enters airways.		
H310	Fatal in contact with skin.		
H311	Toxic in contact with skin.		
H312	Harmful in contact with skin.		
H315	Causes skin irritation.		
H317	May cause an allergic skin reaction.		
H318	Causes serious eye damage.		
H319	Causes serious eye irritation.		
H330	Fatal if inhaled.		
H332	Harmful if inhaled.		
H335	May cause respiratory irritation.		
H336	May cause drowsiness or dizziness.		
H341	Suspected of causing genetic defects.		
H351	Suspected of causing cancer.		
H361	Suspected of damaging fertility or the unborn child.		
H370	Causes damage to organs.		
H372	Causes damage to organs through prolonged or repeated exposure.		
H373	May cause damage to organs through prolonged or repeated exposure.		
H400	Very toxic to aquatic life.		
H411	Toxic to aquatic life with long lasting effects.		
H412	Harmful to aquatic life with long lasting effects.		

SDS Version Summary

Version	Date of Update	Sections Updated
4.23	28/02/2023	Hazards identification - Classification, Composition / information on ingredients - Ingredients

Other information

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

The SDS is a Hazard Communication tool and should be used to assist in the Risk Assessment. Many factors determine whether the reported Hazards are Risks in the workplace or other settings. Risks may be determined by reference to Exposures Scenarios. Scale of use, frequency of use and current or available engineering controls must be considered.

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

PC - STEL: Permissible Concentration-Short Term Exposure Limit

IARC: International Agency for Research on Cancer

ACGIH: American Conference of Governmental Industrial Hygienists

STEL: Short Term Exposure Limit TEEL: Temporary Emergency Exposure Limit. IDLH: Immediately Dangerous to Life or Health Concentrations ES: Exposure Standard 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** AIIC: Australian Inventory of Industrial Chemicals DSL: Domestic Substances List NDSL: Non-Domestic Substances List IECSC: Inventory of Existing Chemical Substance in China EINECS: European INventory of Existing Commercial chemical Substances ELINCS: European List of Notified Chemical Substances NLP: No-Longer Polymers ENCS: Existing and New Chemical Substances Inventory KECI: Korea Existing Chemicals Inventory NZIoC: New Zealand Inventory of Chemicals PICCS: Philippine Inventory of Chemicals and Chemical Substances TSCA: Toxic Substances Control Act TCSI: Taiwan Chemical Substance Inventory INSQ: Inventario Nacional de Sustancias Químicas NCI: National Chemical Inventory FBEPH: Russian Register of Potentially Hazardous Chemical and Biological Substances

Classification and procedure used to derive the classification for mixtures according to Regulation (EC) 1272/2008 [CLP]

Classification according to regulation (EC) No 1272/2008 [CLP] and amendments	Classification Procedure
Corrosive to Metals Category 1, H290	On basis of test data
Skin Corrosion/Irritation Category 1A, H314	Minimum classification
, EUH208	Calculation method

Note:

"This composition meets the criteria for not being harmful to the marine environment according to MARPOL Annex V and may be discharged into the sea when used to clean cargo holds and external surfaces on ships."

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