

Version No: 1

Issue date: 30/APR//2024

Safety Data Sheet (Conforms to Annex II of REACH (1907/2006) - Regulation 2020/878)

Revision date: Not Applicable

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

1.1. Product Identifier

Product name	Clean Buffer 02	
Synonyms	Not Available	
Other means of identification	Not Available	

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

Relevant identified uses	Laboratory use.
Uses advised against	Not Applicable

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

Registered company name	CleanNA
Address	Coenecoop 75, 2741 PH, Waddinxveen, The Netherlands
Telephone	+31 (0) 182 22 33 50
Fax	+31 (0) 182 22 33 98
Website	www.cleanna.com
Email	info@cleanna.com

1.4. Emergency telephone number

Emergency telephone numbers 112 (European emergency number)

SECTION 2. Hazards identification

2.1. Classification of the substance or mixture

Classification according to regulation (EC) No 1272/2008 [CLP] and amendments	H315 – Skin Corrosion/irritation – category 2 H319 – Serious eye damage / Eye irritation – Category 2
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2.2. Label elements

Hazard pictogram(s)	
Signal word	Warning
Hazard statement(s)	
H315	Causes skin irrittaion
H319	Causes serious eye irritation
Supplementary statement(s)	
	Not Applicable
Precautionary statement(s) Prev	rention
P264	Wash all exposed external body areas thoroughly after handling
P280	Wear protective gloves & clothing, eye & face protection

Precautionary statement(s) Response

P337 + P313	eye irritation persist get medical advice	
P302 + P352	IF ON SKIN: wash with plenty of water and soap	
P332 + P313	If skin irritation occurs get medical attention / advice	
P362 + P364	Take off contaminated clothing and wash it before re-use	
P305 + P351 + P338	IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses if present and easy to do. Continue rinsing.	

Precautionary statement(s) Storage

Not Applicable

Precautionary statement(s) Disposal

Not Applicable

2.3. Other hazards

Cumulative effects may result following exposure * Possible respiratory and skin sensitizer *

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

SECTION 3. Composition / information on ingredients

3.1 Substances

See 'Composition on ingredients' in Section 3.2

3.2 Mixtures

1. CAS No 2. EC No 3. Index No 4. REACH No	%[weight]	Name	Classification according to regulation (EC) No 1272/2008 [CLP] and amendments	SCL / M-Factor	Nanoform Particle Characteristics
 Not Available Not Available Not Available Not Available 	1 - 5	Anionic detergent	Flammable Solids – Category 1 Acute Toxicity (Oral, Dermal and Inhalation) – Category 4 Skin Corrosion/Irritation – Category 2 Serious Eye damage/Eye irritation – Category 1 Specific target Organ Toxicity – Single Exposure (resp. tract Irritation) – Category 3 H228, H302, H312, H315, H318, H332, H335	Not Available	Not Available
1. 1310-73-2 2. 215-185-5 3. 011-002-00-6 4. Not Available	0.1 - 1	Sodium hydroxide	Skin Corrosion / Irritation – category 1A H314	Skin Corr. 1A; H314: C >= 5% Skin Corr. 1B: H314: 2% <= C <= 5% Skin Inti2, 2H315: 0,5% <= C <= 2% Eye Inti. 2: H319: 0,5% <= C <= 2%	Not Available
Legend:	Legend: 2. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from C&L * EU IOELVs available; [e] Substance identified as having endocrine disrupting properties				

SECTION 4. First aid measures

4.1. Description of first aid measures

Eye Contact	 If this product comes in contact with the eyes: Wash out immediately with fresh running water Ensure complete irrigation of the eye by keeping eyelids apart and away from eye moving the eyelids by occasionally lifting the upper and lower lids Seek medical attention without delay if pain persists or recurs Removal of contact lenses after an eye injury should only be undertaken by skilled personnel
Skin Contact	If skin contact occurs: - Immediately remove all contaminated clothing including footwear - Flush skin and hair with running water (and soap if available) - Seek medical attention in case of irritation
Inhalation	 If fumes or combustion products are inhaled remove from contaminated area. Other measures are usually unnecessary.
Ingestion	 Immediately give a glass of water First aid is no generally required. If in doubt, contact a POISONS INFORMATION CENTER

4.2 Most important symptoms and effects, both acute and delayed

See Section 11

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

Treat symptomatically

SECTION 5. Firefighting measures

5.1 Extinguishing media

- There is no restriction on the type of extinguisher which may be used.

- use extinguishing media suitable for surrounding area.

5.2 Special hazards arising from the substrate or mixture

Fire Incompatibility	None known.
5.3. Advice for firefighters	
Fire Fighting	 Alert Fire Brigade and tell them location and nature of hazard Wear breathing apparatus plus protective gloves in the event of a fire Prevent, by any means available, spillage from entering drains or water courses Use fire fighting procedures suitable for surrounding area DO NOT approach container suspected to be hot Cool fire exposed containers from path of fire If safe to do so, remove containers from path of fire Equipment should be thoroughly decontaminated after use
Fire/Explosion Hazard	 Non Combustible Not considered a significant fire risk, however containers may burn May emit 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	 Environmental hazard – contain spillage Clean up all spills immediately. Avoid breathing vapours and contact with skin and eyes. Control personal contact with the substance, by using protective equipment. Contain and absorb spill with sand, earth, inert material or vermiculite Wipe up. Place in a suitable, labelled container for waste disposal.
Major Spills	 Environmental hazard – contain spillage Moderate hazard Clear area of personnel and move upwind. Alert Fire Brigade and tell them location and nature of hazard Wear breathing apparatus plus protective gloves Prevent, by any means available, spillage from entering drains or water course Stop leak if safe to do so Contain spill with sand, earth of vermicuite Collect recoverable product into labelied containers for recycling Neutralize / decontaminate residue (see section 13 for specific agent) Collect solid residues and seal in labelled drums for disposal Wash area and prevent runoff into drains After clean up operations decontaminate and launder all protective clothing and equipment before storing an d reuse If contamination of drains or waterways occurs, advise emergency services

6.4. Reference to other sections

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

SECTION 7. Handling and storage

7.1. Precautions for safe handling	
 Avoid all personal contact including inhalation Wear protective clothing when risk of exposure occurs Use in a well-ventilated area Avoid contact with moisture Avoid contact with incompatible materials When handling DO NOT eat, drink or smoke Keep containers securely sealed when not in use Avoid physical damage to containers Always was hands with soap and water after handling Work clothes should be laundered separately. Launder contaminated clothing before reuse Use good occupational work practice Observe manufacturer's storage and handling recommendations contained within this SDS Atmosphere should be regularly against established exposure standards to ensure working conditions a DO NOT allow clothing wet with material to stay in contact with skin 	tre maintained
Fire and explosion protection See section 5	
Other information None known	

7.2. Conditions for safe storage, including any incompatibilities

Suitable container	 Polyethylene of polypropylene container Packaging as recommended by manufacturer Check all containers are clearly labelled and free from leaks 	
Storage incompatibility	e known	
Hazard categories in accordance with Regulation (EC) No 1272/2008	lot available	
Qualifying quantity (tonnes) of dangerous substances as referred to in Article 3(10) for the application of	Not available	

7.3. Specific end use(s)

See section 1.2

SECTION 8. Exposure controls / personal protection

8.1. Control parameters

5.1. Control parameters				
Ingredient	DNELs Exposure Pattern V	Vorker		PNECs Compartment
	Dermal	33.8 mg/kg bw/day	(Systemic, Chronic)	
	Inhalation	4.6 mg/m ³	(Systemic, Chronic)	
	Dermal	0.79 mg/m ³	(Local, Chronic)	
	Inhalation	1 mg/m ³	(Local, Chronic)	
	Dermal	40 mg/kg bw/day	(Systemic, Acute)	
	Inhalation	4.6 mg/m ³	(Systemic, Acute)	
	Dermal	0.79 mg/m ³	(Local, Acute)	
	Inhalation	4.6 mg/m ³	(Local, Acute)	
Sodium hydroxide	Dermal	16.9 mg/kg bw/day	(Systemic, Chronic) *	Not Available
Sodium nyuroxide	Inhalation	2.3 mg/m ³	(Systemic, Chronic) *	
	Oral	10 mg/kg bw/day	(Systemic, Chronic) *	
	Dermal	0.394 mg/m ³	(Local, Chronic) *	
	Inhalation	1 mg/m ³	(Local, Chronic) *	
	Dermal	20 mg/kg bw/day	(Systemic, Acute) *	
	Inhalation	2.3 mg/m ³	(Systemic, Acute) *	
	Oral	10 mg/kg bw/day	(Systemic, Acute) *	
	Dermal	0.394 mg/m ³	(Local, Acute) *	
	Inhalation	2.3 mg/m ³	(Local, Acute) *	

* Values for General Population

Occupational Exposure Limits (OE	iL)					
INGREDIENT DATA						
Source	Ingredient	Material name	TWA	STEL	Peak	Notes
Not Available	Not Available	Not Available	Not Available	Not Available	Not Available	Not Available
Emergency Limits	'					
Ingredient	TEEL-1	r	EEL-2		TEEL-3	
Sodium hydroxide	Not Available	١	lot Available		Not Available	
Ingredient	Original IDLH		Rev	ised IDLH		
Sodium hydroxide	10 mg/m ³		Not Available			

Occupational Exposure Banding

Ingredient	Occupational Exposure Band Rating Occupational Exposure band Limit	
Sodium hydroxide	> 0.1 to <= milligrams per cubic meter of air (mg/m ³)	
Notes:	Occupational exposure banding is a process of assigning chemicals into specific categories or bands based on a chemical's potency and the adverse heal 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.	

8.2. Exposure controls Engineering controls are used to remove a hazard or place a barrier between the worker and the hazard. Well-designed engineering 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. General exhaust is adequate under normal operating conditions. If risk of overexposure exists, wear SAA approved respirator. Correct fit is essential to obtain adequate protection. Provide adequate ventilation in warehouse or closed storage areas. Air contaminants generated in the workplace possess varying "escape" velocities which, in turn, determine the "capture velocities" of fresh circulating air required to effectively remove the contaminant. Type of contaminant Air Speed Solvent, vapours, degreasing etc. evaporating from tank (in still air) 0.25 - 0.5 m/s (50 - 100 f/min) Aerosols, fumes form pouring operations, intermittent container filling, low speed conveyer transfers, 0.5 - 1 m/s (100 - 200 f/min) welding, spray drift, plating acid fumes, pickling (released at low velocity into zone of active generation) Direct spray, spray painting in shallow boots, drum filling, conveyer loading, crusher debts, gas discharge 1 - 2.5 m/s (200 - 500 f/min) 8.2.1 Appropriate engineering (active generation into zone of rapid air motion) controls Grinding, abrasive blasting, tumbling, high speed generated dusts (released at high velocity into zone of very 2.5 - 10 m/s (500 - 200 f/min) high rapid air motion) Within each range the appropriate value depends to: Lower end of range Upper end of range 1: Room air currents minimal of favourable to capture 1: Disturbing room air currents 2: Contaminants of low toxicity or of nuisance value only 2: Contaminants of high toxicity 3: Intermittent, low production 3: High Production, heavy use 4: Large hood or large air mass in motion 4: Small hood - local control only Simple theory shows that air velocity falls rapidly with distance away from the opening of a simple extraction pipe. Velocity generally decreases with the square of distance from the extraction point (in simple cases). Therefore the air speed at the extraction point should be adjusted, accordingly, after reference to distance from the contaminating source. The air velocity at the extraction fan, for example, should be a minimum of 1-2 m/s (200-400 f/min) for extraction of solvents generated in a tank 2 meters distant from the extraction point. Other mechanical considerations, producing performance deficits within the extraction apparatus, make it essential that theoretical air velocities are multiplied by factors of 10 or more when extraction systems are installed or used 8.2.2. Personal protection Chemical goggles. Safety glasses with side 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. This should include a review of lens absorption and adsorption for the class of chemicals in use and an account of injury experience. Medical and first-aid personnel should be trained in their removal and suitable equipment should be readily available. In the event of chemical exposure, begin eye irrigation immediately and remove contact lens as soon as practicable. Lens should be removed at the first signs of eye redness or irritation - lens should be removed in a clean environment only after workers have washed hands thoroughly. [CDC NIOSH Current Intelligence Bulletin 59], [AS/NZS 1336 or national equivalent] Eye and face protection Skin protection See Hand protection below Wear chemical protective gloves, e.g. PVC Wear safety footwear/gumboots, e.g. Rubber The selection of suitable gloves does not only depend on the material, but also on further marks of quality which vary from manufacturer to manufacturer. Where the chemical is a preparation of several substances, the resistance of the glove material can not be calculated in advance and has therefore to be checked prior to the application. The exact break through time for substances has to be obtained from the manufacturer of the protective gloves and has to be observed when making a final choice Personal hygiene is a key element of effective hand care. Gloves must only be worn on clean hands. After using gloves, hands should be washed and dried thoroughly. Application of a non-perfumed moisturiser is recommended. Suitability and durability of glove type is dependent on usage. Important factors in the selection of gloves include: · frequency and duration of contact, chemical resistance of glove material, glove thickness and dexterity Hand / feet protection Select gloves tested to a relevant standard (e.g. Europe EN 374, US F739, AS/NZS 2161.1 or national equivalent). When prolonged or frequently repeated contact may occur, a glove with a protection class of 5 or higher (breakthrough time greater than 240 minutes according to EN 374, AS/NZS 2161.10.1 or national equivalent) is recommended. When only brief contact is expected, a glove with a protection class of 3 or higher (breakthrough time greater than 60 minutes according to EN 374, AS/NZS 2161.10.1 or national equivalent) is recommended. · Some glove polymer types are less affected by movement and this should be taken into account when considering gloves for long-term use. · Contaminated gloves should be replaced. As defined in ASTM F-739-96 in any application, gloves are rated as: · Excellent when breakthrough time > 480 min Good when breakthrough time > 20 min · Fair when breakthrough time < 20 min · Poor when glove material degrades For general applications, gloves with a thickness typically greater than 0.35 mm, are recommended.

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

	Glove thickness may also vary depending on the glove manufacturer, the glove type and the glove model. Therefore, the manufacturers technical data should always be taken into account to ensure selection of the most appropriate glove for the task. Note: Depending on the activity being conducted, gloves of varying thickness may be required for specific tasks. For example: • Thinner gloves (down to 0.1 mm or less) may be required where a high degree of manual dexterity is needed. However, these gloves are only likely to give short duration protection and would normally be just for single use applications, then disposed of. • Thicker gloves (up to 3 mm or more) may be required where there is a mechanical (as well as a chemical) risk i.e. where there is abrasion or puncture potential Gloves must only be worn on clean hands. After using gloves, hands should be washed and dried thoroughly. Application of a non-perfumed moisturiser is recommended.
Body protection	See Other protection below
Other protection	- Overalls - PVC apron - Barrier cream - Skin cleansing cream - Eye wash unit

Respiratory protection

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: Clean Buffer 02

Material	Performance Index
BUTYL	A
NAT_NEOPR+NITRILE	A
NATURAL RUBBER	A
NATUREL + NEOPRENE	A
NEOPRENE	A
MEOPRENE/NATURAL	A
NITRILE	A
NITRILE+PVC	A
PE	A
PE/EVAL/PE	A
PVC	A
SARANEX-23	A
SARANEX-23 2-PLY	A
TEFLON	A
VITON/CHLOROBUTYL	A

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 solution must be based on detailed observation.

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	Not Available		
Physical state	Liquid	Relative density (Water = 1)	Not Available
Odour	Not Available	Partition coefficient n-octanol / water	Not Available
Odour threshold	Not Available	Auto-ignition temperature (°C)	Not Available
pH (as supplied)	Not Available	Decomposition temperature (°C)	Not Available
Melting point / freezing point (°C)	Not Available	Viscosity (cSt)	Not Available
Initial boiling point and boiling range (°C)	Not Available	Molecular weight (g/mol)	Not Available
Flash point (°C)	Not Available	Taste	Not Available
Evaporation rate	Not Available	Explosive properties	Not Available
Flammability	Not Available	Oxidising properties	Not Available
Upper Explosive Limit (%)	Not Available	Surface Tension (dyn/cm or mN/m)	Not Available
Lower Explosive Limit (%)	Not Available	Volatile Component (%vol)	Not Available
Vapour pressure (kPa)	Not Available	Gas group	Not Available
Solubility in water	Immiscible	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 presence of incompatible materials Product is considered stable Hazardous polymerisation will not occur
10.3. Possibility of hazardous reactions	See section 7.2
10.4. Conditions to avoid	See section 7.2
10.5. Incompatible materials	See section 7.2
10.6. Hazardous decomposition products	See section 5.3

SECTION 11. Toxicological information

11.1. Information on toxicological effects The material is not thought to produce adverse health effects or irritation of the respiratory tract (as classified by EC Directives using animal models). Nevertheless, adverse systemic effects have been produced following exposure of animals by at least on other route and good hygiene practice Inhaled requires that exposure be kept to a minimum and that suitable control measures be used in an occupational setting. The material has NOT been classified by ED Directives or other classification systems as "harmful by ingestion". This is because of the lack of Ingestion corroborating animal or human evidence. This material can cause inflammation of the skin on contact in some persons. The material may accentuate any pre-existing dermatitis condition. Skin contact is not thought to have harmful health effects (as classified under EC Directives) the material may still produce health damage following entry through wounds, lesions or abrasions. Skin Contact Eye This material can cause eye irritation and damage in some persons Long-term exposure to the product is not thought to produce chronic effects adverse to the health (as classified by EC Directives using animal models). Chronic Nevertheless exposure by all routes should be minimized as a matter of course. ΤΟΧΙΟΙΤΥ IRRITATION Clean Buffer 02 Not Available Not Available TOXICITY IRRITATION Dermal (rabbit) LD50 1350 mg/kg ² Eye (rabbit) 0.05 mg/24h SEVERE SEVERE Oral (rabbit) LD50 1 ma/24h 325 ma/ka 1 Eve (rabbit) 1 mg/30s rinsed SEVERE Sodium hydroxide Eve (rabbit) Adverse effect observed Eye Irritating 1 Skin (rabbit) 500 mg/24h SEVERE Skin Adverse effect observed Corrosive 1 1. Value obtained from Europe ECHA Registered Substances - Acute toxicity 2. Value obtained from manufacturer's SDS. Unless otherwise Legend specified data extracted from RTECS - Register of Toxic Effect of chemical Substances For alkyl sulfates, alkane sulfonates and alpha-olefin sulfonates: Most chemicals of this category are not defined substances but mixtures of homologues wit different alkyl side chains. Common physical and/or biological pathways result in structurally similar breakdown products and are, together with the surfactant properties, responsible for similar environmental behavior and essentially identical hazard profiles with regard to human health. Acute toxicity: These substances are well absorbed after ingestion, penetration through the skin is however, poor. After absorption, these chemicals are distributed mainly to the liver. In animals, signs of poisoning by mouth include lethargy, hair standing up, decreased motor activity and breathing rate, and diarrhea. Poisoning form skin contact caused irritation, tremor, tonic-clonic convulsions, breathing failure, and weight loss. The C-12-akyl sulfate sodium salt caused the greatest effect. In eye irritation tests, C-12 containing alkyl sulfates ate greater than 10% concentration were severely irritating and produced irreversible effects on the cornea. With increasing alkyl chain length, the irritating potential decreases, and the longer species are only mildly irritant. Animal studies have not shown alkyl sulfates and C14-18 alpha=olefin sulfonates to cause skin sensitization. However there is anecdotal evidence to suggest Clean Buffer 02 sodium lauryl sulfate causes sensitization of the lung, resulting in hyperactive airway dysfunction and lung allergy, accompanied by fatigue, malaise and aching. Significant symptoms of exposure can persist for more than two years and can be activated by a variety of non-specific environmental situation and activated by a variety of non-specific environmental situation and exposure can be activated by a variety of non-specific environmental situation and exposure can be exposed and passive smoking. Airborne sulfonates may be responsible for respiratory allergies an in some cases minor skin allergies. Repeated skin contact with some sulfonated surfactants has produced skin inflammation was sensitizing I predisposed individuals. <u>Repeat dose toxicity</u>: The liver seems to be the only organ that is affected by repeated exposure with elevated levels of liver enzymes, an increase in liver weight and enlargement of liver cells being seen Genetic toxicity: Alkyl sulfates and alkyl-olefin sulfonates do not appear to cause mutations or genetic toxicity

	Cancer-causing potential: Animal testing suggested that alkyl sulfates and alpha-ole <u>Reproductive toxicity</u> : In animal testing these substances only caused harm to th <u>Developmental toxicity</u> : Alkane sulfonates are not considered to be toxic to develo	ne foetus and/or offspring at levels which	
Sodium hydroxide	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 frequent 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. The material may produce severe irritation to the eye causing pronounced inflammation. Repeated or prolonged exposure to irritants may produce conjunctivitis. The material may cause severe skin irritation after prolonged or repeated exposure and may produce on contact kin redness, swelling, the production of vesicles, scaling and thickening of the skin. Repeated exposures may produce severe ulceration		
Acute Toxicity	×	Carcinogenicity	×
Skin Irritation/Corrosion	*	Reproductivity	×
Serious Eye Damage/Irritation	*	STOT - Single Exposure	×
Respiratory or Skin sensitisation	×	STOT - Repeated Exposure	×
Mutagenicity	×	Aspiration Hazard	×
		Legend: 🗙 – Data eit	her not available or does not fill the criteria for classification

- Data available to make classification

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11.2. Information on other hazards

1. Endocrine Disruption Properties Not Available

2. Other Information

See Section 11.1

SECTION 12. Ecological information

· · · · · · · · · · · · · · · · · · ·	Endpoint	Test duration (hr)	Species	Value	Source
Clean Buffer 02	Not available	Not available	Not available	Not available	Not available
	LC50	96	Fish	144 – 267 mg/L	4
Sodium hydroxide	EC50	48	Crustacea	34.59 – 47.13 mg/L	4
	EC50(ECx)	48	Crustacea	34.59 – 47.13 mg/L	4
Legend:	Ecotox databas		ECHA Registered Substances - Ecotoxicological Info OC Aquatic Hazard Assessment Data 6. NITE (Japan)		

12.2. Persistence and degradability

Ingredient	Persistence: Water/Soil	Persistence: Air
Sodium hydroxide	LOW	LOW

12.3. Bio accumulative potential

Ingredient	Bioaccumulation
Sodium hydroxide	LOW (LogKOW = -3.8796)

12.4. Mobility in soil

Ingredient	Mobility
Sodium hydroxide	LOW (KOC = 14.3)

12.5. Results of PBT and vPvB assessment

	Р	В	т	
Relevant available data	Not Available	Not Available	Not Available	
PBT	×	×	×	
vPvB	×	×	×	
PBT Criteria fulfilled?			No	
vPvB			No	
PBT Criteria fulfilled? No vPvB No				

12.6. Endocrine Disruption Properties

Not Available

12.7. Other adverse effects

Not Available

SECTION 13. Disposal considerations

Marine Pollutant

NO

Product / Packaging disposal	 Return to supplier for reuse/recycling, if possible Otherwise: If container cannot be cleaned sufficiently well to ensure that residuals do not remain of if the container cannot be used to store the same product, then puncture containers to prevent reuse and bury at an authorized landfill Where possible retain label warnings and SDS and observe all notices pertaining to the product. Legislation addressing waste disposal requirements may differ by country, state and/ or territory. Each user must refer to laws operating in their area. In some areas, certain wastes must be tracked. A hierarchy of Controls seems to be common - the user should investigate: Reduction Reuse Recycling Disposal (if all else fails) This material may be recycled if unused, or if it has not been contaminated so as to make it unsuitable for its intended use. If it has been contaminated, it may be possible to reclaim the product by filtration, distillation or some other means. Shelf life considerations should also be applied in making decisions of this type. Note that properties of a material may change in use, and recycling or reuse may not always be applied in. In all cases disposal to sewer may be subject to local laws and regulations and these should be considered first. Where in doubt contact the responsible authority. Recycle wherever possible or consult manufacturer for recycling options. Consult State Land Waste Authority for disposal. Bury or incinerate residue at an approved site Recycle ontainers for possible or incidence in authorized inardifill
Waste treatment options	Not Available
Sewage disposal options	Not Available

Land transport (ADR): NOT REGULATED FOR TRANSPORT OF DANGEROUS GOODS

14.1. UN number	Not Applicable	Not Applicable		
14.2. UN proper shipping name	Not Applicable			
14.3. Transport hazard class(es)	Class Not Applicable Sub risk Not Applicable			
14.4. Packing group	Not Applicable			
14.5. Environmental hazard	Not Applicable			
14.6. Special precautions for user	Hazard identification (Kemler)	Not Applicable		
	Classification code	Not Applicable		
	Hazard Label	Not Applicable		
	Special provisions	Not Applicable		
	Limited quantity	Not Applicable		
	Tunnel Restriction Code	Not Applicable		

Air transport (ICAO-IATA / DGR): NOT REGULATED FOR TRANSPORT OF DANGEROUS GOODS

14.1. UN number	Not Applicable	Not Applicable		
14.2. UN proper shipping name	Not Applicable			
	ICAO/IATA Class	Not Applicable		
14.3. Transport hazard	ICAO / IATA Sub risk	Not Applicable		
class(es)	ERG Code	Not Applicable		
14.4. Packing group	Not Applicable			
14.5. Environmental hazard	Not Applicable			
	Special provisions		Not Applicable	
	Cargo Only Packing Instructions		Not Applicable	
	Cargo Only Maximum Qty / Pack		Not Applicable	
14.6. Special precautions for	Passenger and Cargo Packing Instructions		Not Applicable	
user	Passenger and Cargo Maximum Qty / Pack		Not Applicable	
	Passenger and Cargo	Limited Quantity Packing Instructions	Not Applicable	
	Passenger and Cargo	Limited Maximum Qty / Pack	Not Applicable	

Sea transport (IMDG-Code / GGVSee): NOT REGULATED FOR TRANSPORT OF DANGEROUS GOODS

14.1. UN number	Not Applicable		
14.2. UN proper shipping name	Not Applicable		
14.3. Transport hazard class(es)	IMDG Class Not Applicable IMDG Sub risk Not Applicable		
14.4. Packing group	Not Applicable		
14.5. Environmental hazard	Not Applicable		
14.6. Special precautions for user	EMS NumberNot ApplicableSpecial provisionsNot ApplicableLimited QuantitiesNot Applicable		

Inland waterways transport (ADN): NOT REGULATED FOR TRANSPORT OF DANGEROUS GOODS

14.1. UN number	Not Applicable	
14.2. UN proper shipping name	Not Applicable	
14.3. Transport hazard class(es)	Not applicable	Not Applicable
14.4. Packing group	Not applicable	
14.5. Environmental hazard	Not applicable	
	Classification code	Not Applicable
	Special provisions	Not Applicable
14.6 Special precautions for user	Limited quantity	Not Applicable
	Equipment required	Not Applicable
	Fire cones number	Not Applicable

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

Not Applicable

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

Product name	Group
Anionic detergent	Not Available
Sodium hydroxide	Not Available

14.9. Transport in bulk in accordance with the ICG Code

Product name	Ship Type
Anionic detergent	Not Available
Sodium hydroxide	Not Available

SECTION 15. Regulatory information

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

Sodium hydroxide is found on the following regulatory lists Europe EC Inventory

European Union – (EU) regulation (EC) No 1272/2008 on Classification, Labelling and packaging of Substance and Mixtures – Annex VI

European Union – European Inventory of Existing Commercial Chemical Substances (EINECS)

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	ECHA Dossier
Sodium hydroxide	1310-73-2	011-002-00-6	Not Available

Harmonisation (C&L Inventory)	Hazard Class and Category Code(s)	Pictograms Signal Word Code(s)	Hazard Statement Code(s)
1	Skin Corr. 1A	GHS05; Dgr	H314
2	Skin Corr. 1A	GHS05; Dgr	H314
1	Skin Corr. 1A	GHS05; Dgr	H314
2	Met. Corr. 1; Skin Corr. 1A; Eye Dam. 1; STOT SE 3; Acute Tox. 4 ; Acute Tox. 4 ; Aquatic Chronic 3 ; STOT SE 1; Aquatic Acute 3	GHS05; Dgr; GHS06; GHS08	H290; H302; H312; H314; H318; H335; H370; H402; H412;

Harmonization code 1 = The most prevalent classification. Harmonization code 2 = The most sever classification.

National Inventory Status

National Inventory	Status
Australia - AIIC / Australia Non-Industrial Use	Yes
Canada - DSL	Yes
Canada - NDSL	No (sodium hydroxide)
China - IECSC	Yes
Europe - EINEC / ELINCS / NLP	Yes
Japan - ENCS	Yes
Korea - KECI	Yes
New Zealand - NZIoC	Yes
Philippines - PICCS	Yes
USA - TSCA	Yes
Taiwan - TCSI	Yes
Mexico - INSQ	Yes
Vietnam - NCI	Yes
Russia - FBEPH	Yes
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

Full text Risk and Hazard codes		
H290	May be corrosive to metals	
H302	Harmful if swallowed	
H312	Harmful in contact with skin	
H314	Causes severe skin burn and eye damage	
H318	Causes serious eye damage	
H335	May cause respiratory irritation	
H370	Causes damage to organs	
H402	Harmful to aquatic life	
H412	Harmful to aquatic life with long lasting effects	

Version Summary

Version	Date of Update	Sections Updated
1	30/APR/2024	Initial version

Other information

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