

FW-CLD N&A Cloud Nine Flavor Flavor West Manufacturing, LLC.

Version No: 1.1.4.7

Safety Data Sheet accordtog OSHA HazCom Standard (2012) requirements

Chemwatch Hazard Alert Code: 4

Issue Date: **06/18/2021**Print Date: **06/18/2021**L.GHS.USA.EN

SECTION 1 Identification

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Product name	FW-CLD N&A Cloud Nine Flavor
Synonyms	Not Available
Proper shipping name	Extracts, flavoring, liquid
Other means of identification	Not Available

Recommended use of the chemical and restrictions on use

Name, address, and telephone number of the chemical manufacturer, importer, or other responsible party

Registered company name	Flavor West Manufacturing, LLC.		
Address	29400 Hunco Way, Lake Elsinore CA 92530 United States		
Telephone	(951) 893-5120		
Fax	(714) 276-1621		
Website	www.FlavorWest.com		
Email	Flavor@FlavorWest.com		

Emergency phone number

Association / Organisation	Chemwatch	CHEMWATCH EMERGENCY RESPONSE	
Emergency telephone numbers	see below	+61 2 9186 1132	
Other emergency telephone numbers	see below	+1 855-237-5573	

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

Una vez conectado y si el mensaje no está en su idioma preferido, por favor marque 02

SECTION 2 Hazard(s) identification

Classification of the substance or mixture

NFPA 704 diamond



Note: The hazard category numbers found in GHS classification in section 2 of this SDSs are NOT to be used to fill in the NFPA 704 diamond. Blue = Health Red = Fire Yellow = Reactivity White = Special (Oxidizer or water reactive substances)

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Classification

Flammable Liquid Category 3, Eye Irritation Category 2A, Skin Corrosion/Irritation Category 2, Skin Sensitizer Category 1

Label elements

Hazard pictogram(s)





Signal word

Warning

Hazard statement(s)

H226	Flammable liquid and vapour.	
H319	Causes serious eye irritation.	
H315	Causes skin irritation.	
H317	May cause an allergic skin reaction.	

Hazard(s) not otherwise classified

Not Applicable

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 label before use.	

Precautionary statement(s) Prevention

P210	Keep away from heat/sparks/open flames/hot surfaces No smoking.
P233	Keep container tightly closed.
P280	Wear protective gloves, protective clothing, eye protection and face protection.
P240	Ground/bond container and receiving equipment.
P241	Use explosion-proof electrical/ventilating/lighting/intrinsically safe equipment.
P242	Use only non-sparking tools.
P243	Take precautionary measures against static discharge.
P261	Avoid breathing mist/vapours/spray.
P264	Wash all exposed external body areas thoroughly after handling.
P272	Contaminated work clothing should not be allowed out of the workplace.

Precautionary statement(s) Response

P362	Take off contaminated clothing and wash before reuse.
P370+P378	In case of fire: Use alcohol resistant foam or normal protein foam for extinction.
P302+P352	IF ON SKIN: Wash with plenty of water.
P305+P351+P338	IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.
P333+P313	If skin irritation or rash occurs: Get medical advice/attention.
P337+P313	If eye irritation persists: Get medical advice/attention.
P303+P361+P353	IF ON SKIN (or hair): Remove/Take off immediately all contaminated clothing. Rinse skin with water/shower.

Precautionary statement(s) Storage

P403+P235 Store in a well-ventilated place. Keep cool.

Precautionary statement(s) Disposal

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

SECTION 3 Composition / information on ingredients

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Substances

See section below for composition of Mixtures

Mixtures

CAS No	%[weight]	Name
57-55-6	60-70	propylene glycol
56-81-5	10-20	glycerol
64-17-5	10-20	ethanol
7732-18-5	1-10	water

The specific chemical identity and/or exact percentage (concentration) of composition has been withheld as a trade secret.

SECTION 4 First-aid measures

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 and moving the eyelids by occasionally lifting the upper and lower lids. • Seek medical attention without delay; if pain persists or recurs seek medical attention. • Removal of contact lenses after an eye injury should only be undertaken by skilled personnel.
Skin Contact	If skin contact occurs: Immediately remove all contaminated clothing, including footwear. Flush skin and hair with running water (and soap if available). Seek medical attention in event of irritation.
Inhalation	 If fumes 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.
Ingestion	 Immediately give a glass of water. First aid is not generally required. If in doubt, contact a Poisons Information Centre or a doctor.

Most important symptoms and effects, both acute and delayed

See Section 11

Indication of any immediate medical attention and special treatment needed

- ▶ Polyethylene glycols are generally poorly absorbed orally and are mostly unchanged by the kidney.
- Dermal absorption can occur across damaged skin (e.g. through burns) leading to increased osmolality, anion gap metabolic acidosis, elevated calcium, low ionised calcium, CNS depression and renal failure.
- ► Treatment consists of supportive care.

[Ellenhorn and Barceloux: Medical Toxicology]

Propylene glycol is primarily a CNS depressant in large doses and may cause hypoglycaemia, lactic acidosis and seizures.

- The usual measures are supportive care and decontamination (Ipecac/ lavage/ activated charcoal/ cathartics), within 2 hours of exposure should suffice.
- Check the anion gap, arterial pH, renal function and glucose levels.

Ellenhorn and Barceloux: Medical Toxicology

SECTION 5 Fire-fighting measures

- Extinguishing media

 Alcohol stable foam.
 - Dry chemical powder.
 - ► BCF (where regulations permit).
 - ► Carbon dioxide.

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Special hazards arising from the substrate or mixture

Fire Incompatibility

Avoid contamination with oxidising agents i.e. nitrates, oxidising acids, chlorine bleaches, pool chlorine etc. as ignition may result

Special protective equipment and precautions for fire-fighters

Fire Fighting

- Alert Fire Brigade and tell them location and nature of hazard.
- ▶ May be violently or explosively reactive.
- Wear breathing apparatus plus protective gloves.
- ▶ Prevent, by any means available, spillage from entering drains or water course.

▶ Liquid and vapour are flammable.

- ▶ Moderate fire hazard when exposed to heat or flame.
- Vapour forms an explosive mixture with air.
- Moderate explosion hazard when exposed to heat or flame.

Fire/Explosion Hazard

Combustion products include:

carbon monoxide (CO) carbon dioxide (CO2)

acrolein

other pyrolysis products typical of burning organic material.

SECTION 6 Accidental release measures

Personal precautions, protective equipment and emergency procedures

See section 8

Environmental precautions

See section 12

Methods and material for containment and cleaning up

Minor Spills

- ▶ Remove all ignition sources.
- Clean up all spills immediately.
- Avoid breathing vapours and contact with skin and eyes. ▶ Control personal contact with the substance, by using protective equipment.

Chemical Class: alcohols and glycols For release onto land: recommended sorbents listed in order of priority.

SORBENT TYPE	RANK	APPLICATION	COLLECTION	LIMITATIONS

LAND SPILL - SMALL

cross-linked polymer - particulate	1	shovel	shovel	R, W, SS
cross-linked polymer - pillow	1	throw	pitchfork	R, DGC, RT
sorbent clay - particulate	2	shovel	shovel	R,I, P
wood fiber - pillow	3	throw	pitchfork	R, P, DGC, RT
treated wood fiber - pillow	3	throw	pitchfork	DGC, RT
foamed glass - pillow	4	throw	pichfork	R, P, DGC, RT

Major Spills

LAND SPILL - MEDIUM

cross-linked polymer - particulate	1	blower	skiploader	R,W, SS
polypropylene - particulate	2	blower	skiploader	W, SS, DGC
sorbent clay - particulate	2	blower	skiploader	R, I, W, P, DGC
polypropylene - mat	3	throw	skiploader	DGC, RT
expanded mineral - particulate	3	blower	skiploader	R, I, W, P, DGC
polyurethane - mat	4	throw	skiploader	DGC, RT

Legend

DGC: Not effective where ground cover is dense

R; Not reusable

I: Not incinerable

P: Effectiveness reduced when rainy

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RT:Not effective where terrain is rugged

SS: Not for use within environmentally sensitive sites

W: Effectiveness reduced when windy

Reference: Sorbents for Liquid Hazardous Substance Cleanup and Control;

R.W Melvold et al: Pollution Technology Review No. 150: Noyes Data Corporation 1988

- Clear area of personnel and move upwind.
- ▶ Alert Fire Brigade and tell them location and nature of hazard.
- ▶ May be violently or explosively reactive.
- Wear breathing apparatus plus protective gloves.

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

SECTION 7 Handling and storage

Precautions for safe handling

Safe handling

- Containers, even those that have been emptied, may contain explosive vapours.
- ▶ Do NOT cut, drill, grind, weld or perform similar operations on or near containers.
- ▶ Avoid all personal contact, including inhalation.
- Wear protective clothing when risk of overexposure occurs.
- Use in a well-ventilated area.
- Prevent concentration in hollows and sumps.
- DO NOT allow clothing wet with material to stay in contact with skin

Other information

Consider storage under inert gas.

- ▶ Store in original containers in approved flammable liquid storage area.
- Store away from incompatible materials in a cool, dry, well-ventilated area.
- ▶ DO NOT store in pits, depressions, basements or areas where vapours may be trapped.
- ▶ No smoking, naked lights, heat or ignition sources.
- ▶ Material is hygroscopic, i.e. absorbs moisture from the air. Keep containers well sealed in storage.

Conditions for safe storage, including any incompatibilities

Suitable container

- Packing as supplied by manufacturer.
- ▶ Plastic containers may only be used if approved for flammable liquid.
- Check that containers are clearly labelled and free from leaks
- For low viscosity materials (i): Drums and jerry cans must be of the non-removable head type. (ii): 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)
- For manufactured product having a viscosity of at least 250 cSt.

Glycerol:

- reacts violently with strong oxidisers, acetic anhydride, alkali metal hydrides, calcium hypochlorite, calcium oxychloride, chlorine, chromic anhydride, chromium oxides, ethylene oxide, hydrogen peroxide, phosphorous triiodide, potassium chlorate, potassium permanganate, potassium peroxide, silver perchlorate, sodium hydride, sodium peroxide, sodium triiodide, sodium tetrahydroborate,is incompatible with strong acids, caustics, aliphatic amines, isocyanates, uranium fluoride
- is able to polymerise above 145 C
- Glycols and their ethers undergo violent decomposition in contact with 70% perchloric acid. This seems likely to involve formation of the glycol perchlorate esters (after scission of ethers) which are explosive, those of ethylene glycol and 3-chloro-1,2-propanediol being more powerful than glyceryl nitrate, and the former so sensitive that it explodes on addition of water.

Storage incompatibility

Alcohols

- are incompatible with strong acids, acid chlorides, acid anhydrides, oxidising and reducing agents.
- reacts, possibly violently, with alkaline metals and alkaline earth metals to produce hydrogen
- react with strong acids, strong caustics, aliphatic amines, isocyanates, acetaldehyde, benzoyl peroxide, chromic acid, chromium oxide, dialkylzincs, dichlorine oxide, ethylene oxide, hypochlorous acid, isopropyl chlorocarbonate, lithium tetrahydroaluminate, nitrogen dioxide, pentafluoroguanidine, phosphorus halides, phosphorus pentasulfide, tangerine oil, triethylaluminium, triisobutylaluminium
- ▶ should not be heated above 49 deg. C. when in contact with aluminium equipment

SECTION 8 Exposure controls / personal protection

Control parameters

Occupational Exposure Limits (OEL)

INGREDIENT DATA

Source	Ingredient	Material name	TWA	STEL	Peak	Notes
US OSHA Permissible	glycerol	Glycerin (mist)- Respirable	5 mg/m3	Not	Not	Not Available

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Source	Ingredient	Material name	TWA	STEL	Peak	Notes
Exposure Limits (PELs) Table Z-1		fraction		Available	Available	
US OSHA Permissible Exposure Limits (PELs) Table Z-1	glycerol	Glycerin (mist)- Total dust	15 mg/m3	Not Available	Not Available	Not Available
US NIOSH Recommended Exposure Limits (RELs)	glycerol	Glycerin (mist)	Not Available	Not Available	Not Available	See Appendix D
US OSHA Permissible Exposure Limits (PELs) Table Z-1	ethanol	Ethyl alcohol (Ethanol)	1000 ppm / 1900 mg/m3	Not Available	Not Available	Not Available
US NIOSH Recommended Exposure Limits (RELs)	ethanol	Ethyl alcohol	1000 ppm / 1900 mg/m3	Not Available	Not Available	Not Available
US ACGIH Threshold Limit Values (TLV)	ethanol	Ethanol	Not Available	1000 ppm	Not Available	A3

Emergency Limits

Ingredient	TEEL-1	TEEL-2	TEEL-3
propylene glycol	30 mg/m3	330 mg/m3	2,000 mg/m3
propylene glycol	30 mg/m3	1,300 mg/m3	7,900 mg/m3
glycerol	45 mg/m3	180 mg/m3	1,100 mg/m3
ethanol	Not Available	Not Available	15000* ppm

Ingredient	Original IDLH	Revised IDLH
propylene glycol	Not Available	Not Available
glycerol	Not Available	Not Available
ethanol	3,300 ppm	Not Available
water	Not Available	Not Available

Occupational Exposure Banding

Ingredient	Occupational Exposure Band Rating	Occupational Exposure Band Limit
propylene glycol	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.	

MATERIAL DATA

For ethanol:

Odour Threshold Value: 49-716 ppm (detection), 101 ppm (recognition)

Eye and respiratory tract irritation do not appear to occur at exposure levels of less than 5000 ppm and the TLV-TWA is thought to provide an adequate margin of safety against such effects. Experiments in man show that inhalation of 1000 ppm caused slight symptoms of poisoning and 5000 ppm caused strong stupor and morbid sleepiness. Subjects exposed to 5000 ppm to 10000 ppm experienced smarting of the eyes and nose and coughing. Symptoms disappeared within minutes.

for propylene glycol:

Saturated vapour concentration @ 20 deg C.= 65.8 ppm, 204.6 mg/m3; i.e higher concentrations can only occur as aerosols or at higher temperatures. Odour Threshold: Practically odourless.

A small number of individuals show skin irritation or sensitisation from repeated or prolonged exposure to propylene glycol. A workplace environmental exposure limit (WEEL) has been established by AIHA and is thought to be protective against systemic effects.

Exposure controls

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.

Personal protection









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Eye and face protection

- ► Safety glasses with side shields.
- Chemical goggles.
- 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

- ▶ Wear chemical protective gloves, e.g. PVC.
- Wear safety footwear or safety gumboots, e.g. Rubber

NOTE:

Hands/feet protection

- The material may produce skin sensitisation in predisposed individuals. Care must be taken, when removing gloves and other protective equipment, to avoid all possible skin contact.
- Contaminated leather items, such as shoes, belts and watch-bands should be removed and destroyed.

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.

Body protection

See Other protection below

Body protection

Other protection

- Employees working with confirmed human carcinogens should be provided with, and be required to wear, clean, full body protective clothing (smocks, coveralls, or long-sleeved shirt and pants), shoe covers and gloves prior to entering the regulated area. [AS/NZS ISO 6529:2006 or national equivalent]
- Employees engaged in handling operations involving carcinogens should be provided with, and required to wear and use half-face filter-type respirators with filters for dusts, mists and fumes, or air purifying canisters or cartridges. A respirator affording higher levels of protection may be substituted. [AS/NZS 1715 or national equivalent]
- Emergency deluge showers and eyewash fountains, supplied with potable water, should be located near, within sight of, and on the same level with locations where direct exposure is likely.

Prior to each exit from an area containing confirmed human carcinogens, employees should be required to remove and leave protective clothing and equipment at the point of exit and at the last exit of the day, to place used clothing and equipment in impervious containers at the point of exit for purposes of decontamination or disposal. The contents of such impervious containers must be identified with suitable labels. For maintenance and decontamination activities, authorized employees entering the area should be provided with and required to wear clean, impervious garments, including gloves, boots and continuous-air supplied hood.

- Prior to removing protective garments the employee should undergo decontamination and be required to shower upon removal of the garments and hood.
- Overalls.
- ▶ PVC Apron.
- ▶ PVC protective suit may be required if exposure severe.
- Eyewash unit.
- Some plastic personal protective equipment (PPE) (e.g. gloves, aprons, overshoes) are not recommended as they may produce static electricity.
- For large scale or continuous use wear tight-weave non-static clothing (no metallic fasteners, cuffs or pockets).
- Non sparking safety or conductive footwear should be considered. Conductive footwear describes a boot or shoe with a sole made from a conductive compound chemically bound to the bottom components, for permanent control to electrically ground the foot an shall dissipate static electricity from the body to reduce the possibility of ignition of volatile compounds.

Recommended material(s)

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:

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Material	СРІ
BUTYL	С
NATURAL RUBBER	С
NATURAL+NEOPRENE	С
NEOPRENE	С
NITRILE	С
NITRILE+PVC	С
PE/EVAL/PE	С
PVA	С
PVC	С

Respiratory protection

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

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С

VITON

* 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.

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

^ - 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

SECTION 9 Physical and chemical properties

Information on basic physical and chemical properties

Appearance	Clear yellow		
Physical state	Liquid	Relative density (Water = 1)	1.03
Odour	Characteristic	Partition coefficient n-octanol / water	Not Available
Odour threshold	Not Available	Auto-ignition temperature (°C)	Not Available
pH (as supplied)	Not Available	Decomposition temperature	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)	42.7	Taste	Mixed fruit
Evaporation rate	Not Available	Explosive properties	Not Available
Flammability	Flammable.	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 (%)	Not Available
Vapour density (Air = 1)	Not Available	VOC g/L	Not Available

SECTION 10 Stability and reactivity

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Reactivity	See section 7
Chemical stability	 Unstable in the presence of incompatible materials. Product is considered stable. Hazardous polymerisation will not occur.
Possibility of hazardous reactions	See section 7
Conditions to avoid	See section 7
Incompatible materials	See section 7
Hazardous decomposition products	See section 5

SECTION 11 Toxicological information

Information on toxicological 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.

The most common signs of inhalation overexposure to ethanol, in animals, include ataxia, incoordination and drowsiness for those surviving narcosis. The narcotic dose for rats, after 2 hours of exposure, is 19260 ppm.

Inhaled

Exposure to aliphatic alcohols with more than 3 carbons may produce central nervous system effects such as headache, dizziness, drowsiness, muscle weakness, delirium, CNS depression, coma, seizure, and neurobehavioural changes. Symptoms are more acute with higher alcohols. Respiratory tract involvement may produce irritation of the mucosa, respiratory insufficiency, respiratory depression secondary to CNS depression, pulmonary oedema, chemical pneumonitis and bronchitis. Cardiovascular involvement may result in arrhythmias and hypotension.

The material has NOT been classified by EC Directives or other classification systems as "harmful by inhalation". This is because of the lack of corroborating animal or human evidence. In the absence of such evidence, care should be taken nevertheless to ensure exposure is kept to a minimum and that suitable control measures be used, in an occupational setting to control vapours, fumes and aerosols.

Inhalation hazard is increased at higher temperatures.

Inhalation of vapours or aerosols (mists, fumes), generated by the material during the course of normal handling, may be damaging to the health of the individual.

Ingestion of ethanol may produce nausea, vomiting, gastrointestinal bleeding, abdominal pain and diarrhoea. Systemic effects:

Blood concentration:	Effects:
<1.5 g/l	Mild: Impaired visual acuity, coordination and reaction time, emotional lability
1.5-3.0 g/l	Moderate: Slurred speech, confusion, ataxia, emotional lability, perceptual and sensation disturbances possible blackout spells, and incoordination with impaired objective performance in standardised tests. Possible diplopia, flushing, tachycardia, sweating and incontinence. Bradypnoea may occur early and tachypnoea may develop in cases of metabollic acidosis, hypoglycaemia and hypokalaemia.

Ingestion

Ingestion of propylene glycol produced reversible central nervous system depression in humans following ingestion of 60 ml. Symptoms included increased heart-rate (tachycardia), excessive sweating (diaphoresis) and grand mal seizures in a 15 month child who ingested large doses (7.5 ml/day for 8 days) as an ingredient of vitamin preparation.

Excessive repeated ingestions may cause hypoglycaemia (low levels of glucose in the blood stream) among susceptible individuals; this may result in muscular weakness, incoordination and mental confusion.

Very high doses given during feeding studies to rats and dogs produce central nervous system depression (although one-third of that produced by ethanol), haemolysis and insignificant kidney changes.

The toxic effects of glycols (dihydric alcohols), following ingestion are similar to those of alcohol, with depression of the central nervous system (CNS), nausea, vomiting and degenerative changes in liver and kidney.

Effects on the nervous system characterise over-exposure to higher aliphatic alcohols. These include headache, muscle weakness, giddiness, ataxia, (loss of muscle coordination), confusion, delirium and coma. Gastrointestinal effects may include nausea, vomiting and diarrhoea. In the absence of effective treatment, respiratory arrest is the most common cause of death in animals acutely poisoned by the higher alcohols.

The material has NOT been classified by EC Directives or other classification systems as "harmful by ingestion". This is because of the lack of corroborating animal or human evidence. The material may still be damaging to the health of the individual, following ingestion, especially where pre-existing organ (e.g liver, kidney) damage is evident. Present definitions of harmful or toxic substances are generally based on doses producing mortality rather than those producing morbidity (disease, ill-health). Accidental ingestion of the material may be damaging to the health of the individual.

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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.

A single prolonged exposure is not likely to result in the material being absorbed in harmful amounts. However the material may be absorbed in potentially harmful amounts when applied in large quantities to severe burns (second or third degree) over large areas of the body as part of a cream, other topical application or by prolonged contact with clothing accidentally wetted by the material. Absorption under such circumstances can elevated serum osmolality and may result in osmotic shock.

Most liquid alcohols appear to act as primary skin irritants in humans. Significant percutaneous absorption occurs in rabbits but not apparently in man.

Skin Contact
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.

The material produces moderate skin irritation; evidence exists, or practical experience predicts, that the material either

- roduces moderate inflammation of the skin in a substantial number of individuals following direct contact, and/or
- produces significant, but moderate, inflammation when applied to the healthy intact skin of animals (for up to four hours), such inflammation being present twenty-four hours or more after the end of the exposure period.

Skin irritation may also be present after prolonged or repeated exposure; this may result in a form of contact dermatitis (nonallergic). The dermatitis is often characterised by skin redness (erythema) and swelling (oedema) which may progress to blistering (vesiculation), scaling and thickening of the epidermis. At the microscopic level there may be intercellular oedema of the spongy layer of the skin (spongiosis) and intracellular oedema of the epidermis.

Eye

Direct contact of the eye with ethanol may cause immediate stinging and burning with reflex closure of the lid and tearing, transient injury of the corneal epithelium and hyperaemia of the conjunctiva. Foreign-body type discomfort may persist for up to 2 days but healing is usually spontaneous and complete.

Irritation of the eyes may produce a heavy secretion of tears (lachrymation).

Limited evidence or practical experience suggests, that the material may cause eye irritation in a substantial number of individuals. Repeated or prolonged eye contact may cause inflammation characterised by temporary redness (similar to windburn) of the conjunctiva (conjunctivitis); temporary impairment of vision and/or other transient eye damage/ulceration may occur.

Long-term exposure to respiratory irritants may result in disease of the airways involving difficult breathing and related systemic problems.

Practical experience shows that skin contact with the material is capable either of inducing a sensitisation reaction in a substantial number of individuals, and/or of producing a positive response in experimental animals.

Substances that can cause occupational asthma (also known as asthmagens and respiratory sensitisers) can induce a state of specific airway hyper-responsiveness via an immunological, irritant or other mechanism. Once the airways have become hyper-responsive, further exposure to the substance, sometimes even to tiny quantities, may cause respiratory symptoms. These symptoms can range in severity from a runny nose to asthma.

On the basis of epidemiological data, the material is regarded as carcinogenic to humans. There is sufficient data to establish a causal association between human exposure to the material and the development of cancer.

Toxic: danger of serious damage to health by prolonged exposure through inhalation, in contact with skin and if swallowed. Serious damage (clear functional disturbance or morphological change which may have toxicological significance) is likely to be caused by repeated or prolonged exposure. As a rule the material produces, or contains a substance which produces severe lesions. Such damage may become apparent following direct application in subchronic (90 day) toxicity studies or following sub-acute (28 day) or chronic (two-year) toxicity tests.

Limited evidence suggests that repeated or long-term occupational exposure may produce cumulative health effects involving organs or biochemical systems.

Long-term exposure to ethanol may result in progressive liver damage with fibrosis or may exacerbate liver injury caused by other agents

Repeated ingestion of ethanol by pregnant women may adversely affect the central nervous system of the developing foetus, producing effects collectively described as foetal alcohol syndrome. These include mental and physical retardation, learning disturbances, motor and language deficiency, behavioural disorders and reduced head size.

Consumption of ethanol (in alcoholic beverages) may be linked to the development of Type I hypersensitivities in a small number of individuals.

Propylene glycol is though, by some, to be a sensitising principal following the regular use of topical creams by eczema patients. A study of 866 persons using a formulation containing propylene glycol in a patch test indicated that propylene glycol caused primary irritation in 16% of exposed individuals probably caused by dehydration. Undiluted propylene glycol was tested on 1556 persons in a 24 hour patch test. 12.5% showed reactions which were largely toxic (70%) or allergic in nature (30%).

Chronic

FW-CLD N&A Cloud Nine Flavor

TOXICITY IRRITATION Not Available Not Available

propylene glycol

TOXICITY	IRRITATION
Dermal (rabbit) LD50: >2000 mg/kg ^[1]	Eye (rabbit): 100 mg - mild
Inhalation(Rat) LC50; >44.9 mg/L4h ^[2]	Eye (rabbit): 500 mg/24h - mild
Oral(Rat) LD50; >10400 mg/kg ^[2]	Eye: no adverse effect observed (not irritating) ^[1]
	Skin(human):104 mg/3d Intermit Mod
	Skin(human):500 mg/7days mild

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		Skin: no adverse			
	TOXICITY	IRRITATION			
glycerol	dermal (guinea pig) LD50: 58500 mg/kg ^[1]	Not Available			
	Oral(Rat) LD50; >20<39800 mg/kg ^[1]				
	TOXICITY IRRITATION				
	Dermal (rabbit) LD50: 17100 mg/kg ^[1]	Eye (rabbit): 500	0 mg SEVERE		
	Inhalation(Mouse) LC50; 39 mg/L4h ^[2]	mg/24hr-moderate			
ethanol	Oral(Rat) LD50; >7692 mg/kg ^[1]	Eye: adverse eff	fect observed (irritating) ^[1]		
		Skin (rabbit):20	mg/24hr-moderate		
		Skin (rabbit):400 mg (open)-mild			
		Skin: no adverse	e effect observed (not irritating) ^[1]		
	TOXICITY	IRRITATION			
water	Oral(Rat) LD50; >90000 mg/kg ^[2]	Not Available			
Legend:	Value obtained from Europe ECHA Registered Sul	ostances - Acute toxicity 2.* V	alue obtained from manufacturer's SDS.		
og	Unless otherwise specified data extracted from RTEG				
FW-CLD N&A Cloud Nine Flavor	The following information refers to contact allergens as a group and may not be specific to this product. Contact allergies quickly manifest themselves as contact eczema, more rarely as urticaria or Quincke's oedema. The pathogenesis of contact eczema involves a cell-mediated (T lymphocytes) immune reaction of the delayed type. Other allergic skin reactions, e.g. contact urticaria, involve antibody-mediated immune reactions. The significance of the contact allergen is not simply determined by its sensitisation potential: the distribution of the substance and the opportunities for contact with it are equally important.				
GLYCEROL	For glycerol: Acute toxicity: Glycerol is of a low order of acute ora high dose levels, the signs of toxicity include tremor a indicate that glycerol has low potential to irritate the sl very widespread potential for exposure and the abser sensitiser. Repeat dose toxicity: Repeated oral exposure to gly gastro-intestinal tract.	nd hyperaemia of the gastro- kin and the eye. The available ace of case reports of sensitis	intestinal -tract. Skin and eye irritation studies a human and animal data, together with the ation, indicate that glycerol is not a skin		
GLYCEROL	Acute toxicity: Glycerol is of a low order of acute oral high dose levels, the signs of toxicity include tremor a indicate that glycerol has low potential to irritate the slycery widespread potential for exposure and the absersensitiser. Repeat dose toxicity: Repeated oral exposure to gly	nd hyperaemia of the gastro- kin and the eye. The available ace of case reports of sensitis cerol does not induce advers	intestinal -tract. Skin and eye irritation studies a human and animal data, together with the ation, indicate that glycerol is not a skin		
	Acute toxicity: Glycerol is of a low order of acute ora high dose levels, the signs of toxicity include tremor a indicate that glycerol has low potential to irritate the si very widespread potential for exposure and the abser sensitiser. Repeat dose toxicity: Repeated oral exposure to gly gastro-intestinal tract.	nd hyperaemia of the gastro- kin and the eye. The available ace of case reports of sensitis- cerol does not induce adverse rature search. The ren years after exposure to the refunction syndrome (RADS) and diagnosis of RADS include the ent asthma-like symptoms with a spirometry, with the presence	intestinal -tract. Skin and eye irritation studies a human and animal data, together with the ation, indicate that glycerol is not a skin e effects other than local irritation of the e material ceases. This may be due to a which can occur following exposure to high the absence of preceding respiratory disease, thin minutes to hours of a documented ce of moderate to severe bronchial		
WATER FW-CLD N&A Cloud	Acute toxicity: Glycerol is of a low order of acute ora high dose levels, the signs of toxicity include tremor a indicate that glycerol has low potential to irritate the si very widespread potential for exposure and the absersensitiser. Repeat dose toxicity: Repeated oral exposure to gly gastro-intestinal tract. No significant acute toxicological data identified in lite. Asthma-like symptoms may continue for months or exponentially included in lite. Asthma-like symptoms may continue for months or exponentially included in lite. As the included in lite in a non-atopic individual, with abrupt onset of persist exposure to the irritant. A reversible airflow pattern, on hyperreactivity on methacholine challenge testing and	and hyperaemia of the gastro-icin and the eye. The available ice of case reports of sensitistice of case reports of sensitistic of case of cas	intestinal -tract. Skin and eye irritation studies a human and animal data, together with the ation, indicate that glycerol is not a skin e effects other than local irritation of the e material ceases. This may be due to a which can occur following exposure to high the absence of preceding respiratory disease, thin minutes to hours of a documented ce of moderate to severe bronchial ytic inflammation, without eosinophilia, have irred to cause perceptible health damage in which requires extremely high intake over a reconsuming foods or supplements, which red to either inappropriate intravenous		
WATER FW-CLD N&A Cloud Nine Flavor & GYCEROL FW-CLD N&A Cloud Nine Flavor & PROPYLENE	Acute toxicity: Glycerol is of a low order of acute ora high dose levels, the signs of toxicity include tremor a indicate that glycerol has low potential to irritate the si very widespread potential for exposure and the absersensitiser. Repeat dose toxicity: Repeated oral exposure to glygastro-intestinal tract. No significant acute toxicological data identified in lite. Asthma-like symptoms may continue for months or exponsible or highly irritating compound. Key criteria for the in a non-atopic individual, with abrupt onset of persist exposure to the irritant. A reversible airflow pattern, on hyperreactivity on methacholine challenge testing and also been included in the criteria for diagnosis of RAD. The acute oral toxicity of propylene glycol is very low, humans. Serious toxicity generally occurs only at plas relatively short period of time. It would be nearly important at most 1 g/kg of PG. Cases of propylene glycol.	and hyperaemia of the gastro- kin and the eye. The available lice of case reports of sensitis- cerol does not induce adverse rature search. The available reports of sensitis- cerol does not induce adverse rature search. The years after exposure to the refunction syndrome (RADS) The diagnosis of RADS include the rest asthma-like symptoms with the spirometry, with the presence of the lack of minimal lymphocy and large quantities are request that are requested to reach toxic levels by the potential for the potential for or repeated exposure and mate and swelling and swelling endings.	intestinal -tract. Skin and eye irritation studies a human and animal data, together with the ation, indicate that glycerol is not a skin e effects other than local irritation of the effects other than local irritation of the e material ceases. This may be due to a which can occur following exposure to high the absence of preceding respiratory disease, thin minutes to hours of a documented se of moderate to severe bronchial ytic inflammation, without eosinophilia, have irred to cause perceptible health damage in the consuming foods or supplements, which red to either inappropriate intravenous or long-term oral toxicity is also low. ay produce a contact dermatitis (nonallergic). ing the epidermis. Histologically there may be		
WATER FW-CLD N&A Cloud Nine Flavor & GYCEROL FW-CLD N&A Cloud Nine Flavor & PROPYLENE GLYCOL PROPYLENE GLYCOL &	Acute toxicity: Glycerol is of a low order of acute ora high dose levels, the signs of toxicity include tremor a indicate that glycerol has low potential to irritate the si very widespread potential for exposure and the absersensitiser. Repeat dose toxicity: Repeated oral exposure to gly gastro-intestinal tract. No significant acute toxicological data identified in lite. Asthma-like symptoms may continue for months or exponentially included in the continuation of the initial annon-atopic individual, with abrupt onset of persist exposure to the irritant. A reversible airflow pattern, on hyperreactivity on methacholine challenge testing and also been included in the criteria for diagnosis of RAD. The acute oral toxicity of propylene glycol is very low, humans. Serious toxicity generally occurs only at plas relatively short period of time. It would be nearly important at most 1 g/kg of PG. Cases of propylene glycal administration or accidental ingestion of large quantitic. The material may cause skin irritation after prolonged.	and hyperaemia of the gastro- kin and the eye. The available lice of case reports of sensitis- cerol does not induce adverse rature search. The available reports of sensitis- cerol does not induce adverse rature search. The years after exposure to the refunction syndrome (RADS) The diagnosis of RADS include the rest asthma-like symptoms with the spirometry, with the presence of the lack of minimal lymphocy and large quantities are request that are requested to reach toxic levels by the potential for the potential for or repeated exposure and mate and swelling and swelling endings.	intestinal -tract. Skin and eye irritation studies a human and animal data, together with the ation, indicate that glycerol is not a skin e effects other than local irritation of the effects other than local irritation of the e material ceases. This may be due to a which can occur following exposure to high the absence of preceding respiratory disease, thin minutes to hours of a documented se of moderate to severe bronchial ytic inflammation, without eosinophilia, have irred to cause perceptible health damage in the consuming foods or supplements, which red to either inappropriate intravenous or long-term oral toxicity is also low. ay produce a contact dermatitis (nonallergic). ing the epidermis. Histologically there may be		
WATER FW-CLD N&A Cloud Nine Flavor & GYCEROL FW-CLD N&A Cloud Nine Flavor & PROPYLENE GLYCOL PROPYLENE GLYCOL & ETHANOL	Acute toxicity: Glycerol is of a low order of acute ora high dose levels, the signs of toxicity include tremor a indicate that glycerol has low potential to irritate the sivery widespread potential for exposure and the absersensitiser. Repeat dose toxicity: Repeated oral exposure to gly gastro-intestinal tract. No significant acute toxicological data identified in lite. Asthma-like symptoms may continue for months or exponentially included in the continuation of the initial and included in the criteria for the initial and included in the criteria for diagnosis of RAD. The acute oral toxicity of propylene glycol is very low, humans. Serious toxicity generally occurs only at plas relatively short period of time. It would be nearly imponential at most 1 g/kg of PG. Cases of propylene glycadministration or accidental ingestion of large quantitic. The material may cause skin irritation after prolonged This form of dermatitis is often characterised by skin intercellular oedema of the spongy layer (spongiosis)	and hyperaemia of the gastro- kin and the eye. The available lice of case reports of sensitis. It cerol does not induce adverse rature search. It is gastro- lice of case reports of sensitis. It is gastro- lice of case reports of sensitis. It is gastro- lice of case reports of sensitis. It is gastro- lice of case reports of sensitis. It is gastro- lice of search to the research of the lack of minimal lymphocy. It is gastro- lice of search toxic levels by the selection of the lack of minimal lymphocy. It is gastro- lice of search toxic levels by the selection of the lack of minimal lymphocy. It is gastro- lice of case reports of the lack of minimal lymphocy. It is gastro- lice of case reports of the lack of minimal lymphocy. It is gastro- lice of case reports of sensitis. It is gastro- li	intestinal -tract. Skin and eye irritation studies a human and animal data, together with the ation, indicate that glycerol is not a skin e effects other than local irritation of the effects other than local irritation of the e material ceases. This may be due to a which can occur following exposure to high the absence of preceding respiratory disease, thin minutes to hours of a documented ce of moderate to severe bronchial ytic inflammation, without eosinophilia, have irred to cause perceptible health damage in which requires extremely high intake over a consuming foods or supplements, which ed to either inappropriate intravenous or long-term oral toxicity is also low. The produce a contact dermatitis (nonallergic) ing the epidermis. Histologically there may be the epidermis.		
WATER FW-CLD N&A Cloud Nine Flavor & GYCEROL FW-CLD N&A Cloud Nine Flavor & PROPYLENE GLYCOL PROPYLENE GLYCOL & ETHANOL Acute Toxicity	Acute toxicity: Glycerol is of a low order of acute ora high dose levels, the signs of toxicity include tremor a indicate that glycerol has low potential to irritate the sivery widespread potential for exposure and the absersensitiser. Repeat dose toxicity: Repeated oral exposure to gly gastro-intestinal tract. No significant acute toxicological data identified in lite. Asthma-like symptoms may continue for months or exponsible acute in a non-allergenic condition known as reactive airways delevels of highly irritating compound. Key criteria for the in a non-atopic individual, with abrupt onset of persist exposure to the irritant. A reversible airflow pattern, on hyperreactivity on methacholine challenge testing and also been included in the criteria for diagnosis of RAD. The acute oral toxicity of propylene glycol is very low, humans. Serious toxicity generally occurs only at plas relatively short period of time. It would be nearly important at most 1 g/kg of PG. Cases of propylene glycadministration or accidental ingestion of large quantitic. The material may cause skin irritation after prolonged. This form of dermatitis is often characterised by skin intercellular oedema of the spongy layer (spongiosis).	and hyperaemia of the gastro- kin and the eye. The available lice of case reports of sensitis cerol does not induce adverse rature search. The search are exposure to the refunction syndrome (RADS) and diagnosis of RADS included the licent asthma-like symptoms with a spirometry, with the presence of the lack of minimal lymphocy and large quantities are requested to poisoning are usually related the syndrometry of the search toxic levels by the logical poisoning are usually related the syndrometry of the search toxic levels by the poisoning are usually related the syndrometry of the syndrometry of the syndrometry of the syndrometry the syndrometry the syndrometry of the syndr	intestinal -tract. Skin and eye irritation studies a human and animal data, together with the ation, indicate that glycerol is not a skin e effects other than local irritation of the effects other than local irritation of the e material ceases. This may be due to a which can occur following exposure to high the absence of preceding respiratory disease, thin minutes to hours of a documented to of moderate to severe bronchial ytic inflammation, without eosinophilia, have irred to cause perceptible health damage in the which requires extremely high intake over a reconsuming foods or supplements, which red to either inappropriate intravenous or long-term oral toxicity is also low. The appropriate intravenous are produce a contact dermatitis (nonallergic). In the epidermis. Histologically there may be the epidermis.		
WATER FW-CLD N&A Cloud Nine Flavor & GYCEROL FW-CLD N&A Cloud Nine Flavor & PROPYLENE GLYCOL PROPYLENE GLYCOL & ETHANOL Acute Toxicity Skin Irritation/Corrosion Serious Eye Damage/Irritation	Acute toxicity: Glycerol is of a low order of acute ora high dose levels, the signs of toxicity include tremor a indicate that glycerol has low potential to irritate the sivery widespread potential for exposure and the absersensitiser. Repeat dose toxicity: Repeated oral exposure to glygastro-intestinal tract. No significant acute toxicological data identified in lite. Asthma-like symptoms may continue for months or exponsible acute airways dylevels of highly irritating compound. Key criteria for the in a non-atopic individual, with abrupt onset of persist exposure to the irritant. A reversible airflow pattern, on hyperreactivity on methacholine challenge testing and also been included in the criteria for diagnosis of RAD. The acute oral toxicity of propylene glycol is very low, humans. Serious toxicity generally occurs only at plas relatively short period of time. It would be nearly important at most 1 g/kg of PG. Cases of propylene glycadministration or accidental ingestion of large quantitity. The material may cause skin irritation after prolonged This form of dermatitis is often characterised by skin intercellular oedema of the spongy layer (spongiosis)	and hyperaemia of the gastro- kin and the eye. The available lice of case reports of sensitis cerol does not induce adverse rature search. The years after exposure to the refunction syndrome (RADS) and diagnosis of RADS included and asthma-like symptoms with a spirometry, with the presence of the lack of minimal lymphocy and large quantities are reques and large quantities are r	intestinal -tract. Skin and eye irritation studies a human and animal data, together with the ation, indicate that glycerol is not a skin e effects other than local irritation of the e material ceases. This may be due to a which can occur following exposure to high the absence of preceding respiratory disease, thin minutes to hours of a documented ce of moderate to severe bronchial ytic inflammation, without eosinophilia, have irred to cause perceptible health damage in which requires extremely high intake over a consuming foods or supplements, which ed to either inappropriate intravenous or long-term oral toxicity is also low. ay produce a contact dermatitis (nonallergic). ing the epidermis. Histologically there may be the epidermis.		
WATER FW-CLD N&A Cloud Nine Flavor & GYCEROL FW-CLD N&A Cloud Nine Flavor & PROPYLENE GLYCOL PROPYLENE GLYCOL & ETHANOL Acute Toxicity Skin Irritation/Corrosion Serious Eye	Acute toxicity: Glycerol is of a low order of acute ora high dose levels, the signs of toxicity include tremor a indicate that glycerol has low potential to irritate the sivery widespread potential for exposure and the absersensitiser. Repeat dose toxicity: Repeated oral exposure to gly gastro-intestinal tract. No significant acute toxicological data identified in lite. Asthma-like symptoms may continue for months or exponentially included in the continued of the initial and non-atlopic individual, with abrupt onset of persist exposure to the irritant. A reversible airflow pattern, on hyperreactivity on methacholine challenge testing and also been included in the criteria for diagnosis of RAL. The acute oral toxicity of propylene glycol is very low, humans. Serious toxicity generally occurs only at plas relatively short period of time. It would be nearly important at most 1 g/kg of PG. Cases of propylene glycal administration or accidental ingestion of large quantitic. The material may cause skin irritation after prolonged This form of dermatitis is often characterised by skin intercellular oedema of the spongy layer (spongiosis)	and hyperaemia of the gastro- kin and the eye. The available lice of case reports of sensitis cerol does not induce adverse rature search. The search are exposure to the refunction syndrome (RADS) and diagnosis of RADS included the licent asthma-like symptoms with a spirometry, with the presence of the lack of minimal lymphocy and large quantities are requested to poisoning are usually related the syndrometry of the search toxic levels by the logical poisoning are usually related the syndrometry of the search toxic levels by the poisoning are usually related the syndrometry of the syndrometry of the syndrometry of the syndrometry the syndrometry the syndrometry of the syndr	intestinal -tract. Skin and eye irritation studies a human and animal data, together with the ation, indicate that glycerol is not a skin e effects other than local irritation of the e material ceases. This may be due to a which can occur following exposure to high the absence of preceding respiratory disease, thin minutes to hours of a documented ce of moderate to severe bronchial ytic inflammation, without eosinophilia, have irred to cause perceptible health damage in which requires extremely high intake over a consuming foods or supplements, which red to either inappropriate intravenous or long-term oral toxicity is also low. ay produce a contact dermatitis (nonallergic). ing the epidermis. Histologically there may be the epidermis.		

Legend: X − Data either not available or does not fill the criteria for classification

✓ – Data available to make classification

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SECTION 12 Ecological information

Toxicity

FIM OLD NOA Claud	Endpoint	Test Duration (hr)	Species	Value	Source	
FW-CLD N&A Cloud Nine Flavor	Not Available	Not Available	Not Available	Not Available	Not Available	
	Endpoint	Test Duration (hr)	Species	Value	Source	
	NOEC(ECx)	336h	Algae or other aquatic plants	<5300mg/l	1	
	EC50	72h	Algae or other aquatic plants	19300mg/l	2	
propylene glycol	LC50	96h	Fish	>10000mg/l	2	
	EC50	48h	Crustacea	>114.4mg/L	4	
	EC50	96h	Algae or other aquatic plants	19000mg/l	2	
	Endpoint	Test Duration (hr)	Species	Value	Source	
glycerol	EC0(ECx)	24h	Crustacea	>500mg/l	1	
	LC50	96h	Fish	885mg/l	2	
	Endpoint	Test Duration (hr)	Species	Value	Sourc	
	EC50(ECx)	96h	Algae or other aquatic plants	<0.001mg/L	4	
- 41	EC50	72h	Algae or other aquatic plants	Algae or other aquatic plants 275mg/l		
ethanol	LC50	96h	Fish	>100mg/l	2	
	EC50	48h	Crustacea	>79mg/L	4	
	EC50	96h	Algae or other aquatic plants	<0.001mg/L	4	
	Endpoint	Test Duration (hr)	Species	Value	Source	
water	Not Available	Not Available	Not Available	Not Available	Not Available	
Legend:	3. EPIWIN Sui	te V3.12 (QSAR) - Aquatic Toxicit	e ECHA Registered Substances - Ecotoxicologi y Data (Estimated) 4. US EPA, Ecotox database IITE (Japan) - Bioconcentration Data 7. METI (J	e - Aquatic Toxicity Da	ata 5.	

When ethanol is released into the soil it readily and quickly biodegrades but may leach into ground water; most is lost by evaporation. When released into water the material readily evaporates and is biodegradable.

Ethanol does not bioaccumulate to an appreciable extent.

The material is readily degraded by reaction with photochemically produced hydroxy radicals; release into air will result in photodegradation and wet deposition. Propylene glycol is known to exert high levels of biochemical oxygen demand (BOD) during degradation in surface waters. This process can adversely affect aquatic life by consuming oxygen needed by aquatic organisms for survival. Large quantities of dissolved oxygen (DO) in the water column are consumed when microbial populations decompose propylene glycol.

Sufficient dissolved oxygen levels in surface waters are critical for the survival of fish, macro-invertebrates, and other aquatic organisms.

For glycerol

log Kow : -2.66- -2.47 BOD 5: 0.617-0.87,31-51% COD : 1.16,82-95% ThOD : 1.217-1.56 Completely biodegradable.

Environmental fate:

Based on the relevant physical-chemical properties and the fact that glycerol is readily biodegradable, glycerol will partition primarily to water.

Biodegradability: Glycerol is considered to be readily biodegradable in the aquatic environment. Pre-adapted microorganisms can degrade glycerol rapidly under both aerobic and anaerobic conditions.

DO NOT discharge into sewer or waterways.

Persistence and degradability

Ingredient	Persistence: Water/Soil	Persistence: Air
propylene glycol	LOW	LOW
glycerol	LOW	LOW
ethanol	LOW (Half-life = 2.17 days)	LOW (Half-life = 5.08 days)
water	LOW	LOW

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Bioaccumulative potential

Ingredient	Bioaccumulation
propylene glycol	LOW (BCF = 1)
glycerol	LOW (LogKOW = -1.76)
ethanol	LOW (LogKOW = -0.31)

Mobility in soil

Ingredient	Mobility
propylene glycol	HIGH (KOC = 1)
glycerol	HIGH (KOC = 1)
ethanol	HIGH (KOC = 1)

SECTION 13 Disposal considerations

Waste treatment methods

Product / Packaging

disposal

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

Otherwise:

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

Legislation addressing waste disposal requirements may differ by country, state and/ or territory. Each user must refer to laws operating in their area. In some areas, certain wastes must be tracked.

A Hierarchy of Controls seems to be common - the user should investigate:

- Reduction
- ▶ 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.

- 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.
- Dispose of 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).
- Decontaminate empty containers.

SECTION 14 Transport information

Labels Required



Marine Pollutant

Land transport (DOT)

UN number	1197			
UN proper shipping name	Extracts, flavoring, liquid			
Transport hazard class(es)	Class Subrisk	3 Not Applicable		
Packing group	III			
Environmental hazard	Not Applica	Not Applicable		

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Hazard identification (Kemler) Not Applicable Classification code Not Applicable Hazard Label Special precautions for Special provisions B1, IB3, T2, TP1 Limited quantity Not Applicable Tunnel Restriction Code Not Applicable

Air transport (ICAO-IATA / DGR)

UN number	1197		
UN proper shipping name	Extracts, flavouring, liquid		
	ICAO/IATA Class	3	
Transport hazard class(es)	ICAO / IATA Subrisk	Not Applicable	
	ERG Code	3L	
Packing group	III		
Environmental hazard	Not Applicable		
	Special provisions		А3
	Cargo Only Packing Instructions		366
	Cargo Only Maximum	Qty / Pack	220 L
Special precautions for user	Passenger and Cargo Packing Instructions 355		
user	Passenger and Cargo Maximum Qty / Pack		60 L
	Passenger and Cargo	Limited Quantity Packing Instructions	Y344
	Passenger and Cargo Limited Maximum Qty / Pack		10 L

Sea transport (IMDG-Code / GGVSee)

UN number	1197				
UN proper shipping name	EXTRACTS, FLAVO	EXTRACTS, FLAVOURING, LIQUID			
Transport hazard class(es)	IMDG Class 3 IMDG Subrisk Not Applicable				
Packing group	III				
Environmental hazard	Not Applicable				
Special precautions for user	EMS Number F-E , S-D Special provisions 223 955 Limited Quantities 5 L				
	Limited Quantities	5 L			

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

Not Applicable

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

Product name	Group
propylene glycol	Not Available
glycerol	Not Available
ethanol	Not Available
water	Not Available

Transport in bulk in accordance with the ICG Code

Product name	Ship Type
propylene glycol	Not Available
glycerol	Not Available
ethanol	Not Available

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Product name	Ship Type
water	Not Available

SECTION 15 Regulatory information

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

propylene glycol is found on the following regulatory lists

US AIHA Workplace Environmental Exposure Levels (WEELs) US ATSDR Minimal Risk Levels for Hazardous Substances (MRLs) US DOE Temporary Emergency Exposure Limits (TEELs)

US EPA Integrated Risk Information System (IRIS)

US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory US Toxicology Excellence for Risk Assessment (TERA) Workplace Environmental Exposure Levels (WEEL)

glycerol is found on the following regulatory lists

US DOE Temporary Emergency Exposure Limits (TEELs) US NIOSH Recommended Exposure Limits (RELs) US OSHA Permissible Exposure Limits (PELs) Table Z-1

US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory US TSCA Chemical Substance Inventory - Interim List of Active Substances

US TSCA Chemical Substance Inventory - Interim List of Active Substances

ethanol is found on the following regulatory lists

US ACGIH Threshold Limit Values (TLV) US ACGIH Threshold Limit Values (TLV) - Carcinogens US DOE Temporary Emergency Exposure Limits (TEELs) US NIOSH Recommended Exposure Limits (RELs)

US OSHA Permissible Exposure Limits (PELs) Table Z-1 US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory US TSCA Chemical Substance Inventory - Interim List of Active Substances

water is found on the following regulatory lists

US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory

US TSCA Chemical Substance Inventory - Interim List of Active Substances

Federal Regulations

Superfund Amendments and Reauthorization Act of 1986 (SARA)

Section 311/312 hazard categories

Flammable (Gases, Aerosols, Liquids, or Solids)	Yes
Gas under pressure	No
Explosive	No
Self-heating	No
Pyrophoric (Liquid or Solid)	No
Pyrophoric Gas	No
Corrosive to metal	No
Oxidizer (Liquid, Solid or Gas)	No
Organic Peroxide	No
Self-reactive	No
In contact with water emits flammable gas	No
Combustible Dust	No
Carcinogenicity	No
Acute toxicity (any route of exposure)	No
Reproductive toxicity	No
Skin Corrosion or Irritation	Yes
Respiratory or Skin Sensitization	Yes
Serious eye damage or eye irritation	Yes
Specific target organ toxicity (single or repeated exposure)	No
Aspiration Hazard	No
Germ cell mutagenicity	No
Simple Asphyxiant	No
Hazards Not Otherwise Classified	No

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None Reported

State Regulations

US. California Proposition 65

None Reported

National Inventory Status

National Inventory	Status		
Australia - AIIC / Australia Non-Industrial Use	Yes		
Canada - DSL	Yes		
Canada - NDSL	No (propylene glycol; glycerol; ethanol; water)		
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 and are not exempt from listing(see specific ingredients in brackets)		

SECTION 16 Other information

Revision Date	06/18/2021
Initial Date	06/19/2021

SDS Version Summary

Version	Date of Update	Sections Updated
0.0.3.1	05/10/2021	Regulation Change
0.0.4.1	05/24/2021	Regulation Change
0.0.4.2	05/30/2021	Template Change
0.0.4.3	06/04/2021	Template Change
0.0.4.4	06/05/2021	Template Change
0.0.4.5	06/09/2021	Template Change
0.0.4.6	06/11/2021	Template Change
0.0.4.7	06/15/2021	Template Change

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.

Definitions and abbreviations

 ${\sf PC-TWA: Permissible \ Concentration-Time \ Weighted \ Average}$

PC-STEL: Permissible Concentration-Short Term Exposure Limit

IARC: International Agency for Research on Cancer

ACGIH: American Conference of Governmental Industrial Hygienists

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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

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