

SAFETY DATA SHEET

in compliance with EC Regulation No. 1907/2006 and EC Regulation No. 830/2015

SECTION 1: IDENTIFICATION OF THE SUBSTANCE AND OF THE COMPANY

1.1 PRODUCT IDENTIFIER

Name of the substance: L (+) Tartaric Acid (99+%)
Trade name: Natural Tartaric Acid
CAS Number: 87-69-4 (99+%)
EC Number: 201-766-0
Reach Registration Nr. 01-2119537204-47-0005

1.2 RELEVANT IDENTIFIED USES OF THE SUBSTANCE OR MIXTURE AND USES ADVISED AGAINST

Acidifier, antioxidant, flavour enhancer and stabilising agent.
Food industry (production of tinned food, jam, jelly, confectionery and biscuits in general, soft drinks and table waters; acidifier in wine-making field). Pharmaceutical and Cosmetic Industry (preparation of medicines, effervescent tablets and soluble drugs; excipient and acidifier in syrups and antibiotics; production of natural beauty cream for face and body) and Industrial and Technical (retarding agent in the preparation of gypsum, used in the formulation of waterproof cements and heat-insulator; it is also used in textiles, tannings, ceramics, galvanoplastics, cleaning agents, used as laboratory reagent, mining and offshore industries).

1.3. DETAILS OF THE SUPPLIER OF THE SAFETY DATA SHEET

Company: Glycerin Supplier LLC - 13320 Emmett Rd Houston, TX 77041-832-795-6898 - glycerinsupplier@gmail.com

1.4. EMERGENCY TELEPHONE NUMBER

(800) 222-1222

SECTION 2: HAZARDS IDENTIFICATION

2.1. CLASSIFICATION OF THE SUBSTANCE OR MIXTURE

Classification pursuant to EC REG. No. 1272/2008
Causes serious eye damage, Cat.1 H318

2.2. LABEL ELEMENTS

Classification pursuant to EC REG. No. 1272/2008
Hazard pictograms



CORROSION

Signal Word:
Danger

Hazard statements:
H318: Causes serious eye damage

Precautionary statements:
P280: Wear protective gloves/protective clothing/eye protection/face protection.
P305+P351+P338: IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.
P310: Immediately call a POISON CENTER/doctor

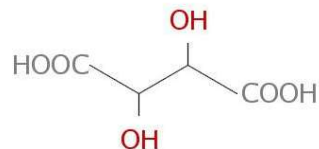
2.3. OTHER HAZARDS

No information available.

SECTION 3: COMPOSITION / INFORMATION ON INGREDIENTS

3.1. SUBSTANCES

CAS Number:	87-69-4 (99+%)
IUPAC Name:	Tartaric Acid
CAS Name:	Butanedioic acid, 2,3-dihydroxy- [R-(R,R)]-
EC Number:	201-766-0
Molecular weight:	150.09 g/mol
Formula:	C ₄ H ₆ O ₆
Chemical formula:	HOOCCH(OH)CH(OH)COOH



3.2. MIXTURES

Not applicable.

SECTION 4: FIRST AID MEASURES

4.1. DESCRIPTION OF FIRST AID MEASURES

Inhalation:	Remove victim from exposure and to open air. Seek medical advice, if necessary.
Skin contact:	Wash off with soap and plenty of water. Take off contaminated garments. If skin irritation persists, consult a specialist.
Eye contact:	Rinse immediately with running water with eyelids held open, for at least 10 minutes. Call an eye specialist, if necessary.
Ingestion:	Make the victim drink plenty of water. Call a doctor, if necessary.

4.2. MOST IMPORTANT SYMPTOMS AND EFFECTS, BOTH ACUTE AND DELAYED

Irritating effects.

4.3. INDICATION OF ANY IMMEDIATE MEDICAL ATTENTION AND SPECIAL TREATMENT NEEDED

Call a doctor in case of exposure.

SECTION 5: FIREFIGHTING MEASURES

5.1. EXTINGUISHING MEDIA

Suitable extinguishing media: Water, CO₂, Foam, Powder.

Extinguishing media not to be used: No limits.

5.2. SPECIAL HAZARDS ARISING FROM THE SUBSTANCE OR MIXTURE

In case of fire, gas and hazardous vapours may be formed.

5.3. ADVICE FOR FIREFIGHTERS

Protective equipment: Do not stay in the hazardous area without a self-contained breathing apparatus.

SECTION 6: ACCIDENTAL RELEASE MEASURES

6.1. PERSONAL PRECAUTIONS, PROTECTIVE EQUIPMENT AND EMERGENCY PROCEDURES

6.1.1. For non-emergency personnel

Avoid generation of dust, do not inhale dust. Avoid contact with the substance. Ensure the supply of fresh air in closed rooms.

6.1.2. For emergency responders

Personal protective clothing, take note of any information in Section 8 on suitable and unsuitable materials.

6.2. ENVIRONMENTAL PRECAUTIONS

Avoid penetration into the sewerage system.

6.3. METHODS AND MATERIAL FOR CONTAINMENT AND CLEANING UP

Collect and place them in a container suitable for recovery. Avoid generation of dust. After collection, flush away traces with water.

6.4. REFERENCE TO OTHER SECTIONS

For instructions on waste treatment, see section 13.

SECTION 7: HANDLING AND STORAGE

7.1. PRECAUTIONS FOR SAFE HANDLING

Use with adequate ventilation system. Minimise dust generation and accumulation. Avoid contact with eyes, skin and clothing. Keep the container tightly closed. Avoid ingestion and inhalation.

7.2. CONDITIONS FOR SAFE STORAGE, INCLUDING ANY INCOMPATIBILITIES

Store in a tightly closed container. Store in a fresh and dry area.

7.3. SPECIFIC END USE(S)

See paragraph 1.2

SECTION 8: EXPOSURE CONTROLS / PERSONAL PROTECTION

8.1. CONTROL PARAMETERS

DN(M)ELs for workers

EXPOSURE PATTERN	ROUTE	DESCRIPTOR	DNEL / DMEL	(CORRECTED) DOSE DESCRIPTOR
Long-term - systemic effects	Dermal	DNEL (Derived No Effect Level)	2.9 mg/kg bw/day	NOAEL: 145 mg/kg bw/day (based on AF of 50)
Long-term - systemic effects	Inhalation	DNEL (Derived No Effect Level)	5,2 mg/m ³	NOAEC: 260.0 mg/m ³ (based on AF of 50)

DN(M)ELs for the general population

EXPOSURE PATTERN	ROUTE	DESCRIPTOR	DNEL / DMEL	(CORRECTED) DOSE DESCRIPTOR
Long-term - systemic effects	Dermal	DNEL (Derived No Effect Level)	1.5 mg/kg bw/day	NOAEL: 150 mg/kg bw/day (based on AF of 100)
Long-term - systemic effects	Inhalation	DNEL (Derived No Effect Level)	1.3 mg/m ³	NOAEC: 130 mg/m ³ (based on AF of 100)
Long-term - systemic effects	Oral	DNEL (Derived No Effect Level)	8.1 mg/kg bw/day	NOAEL: 810 mg/kg bw/day (based on AF of 100)

8.2. EXPOSURE CONTROLS

8.2.1. Appropriate engineering controls

Ensure adequate ventilation, especially in confined areas.

8.2.2. Individual protection measures, such as personal protective equipment

Protective clothing should be selected specifically for the working place and type of work. Take off any contaminated garments. It is advisable to apply protective cream for the skin. Wash hands after handling this substance.

Eyes/face protections

Wear protective goggles against chemicals.

Skin protection

If hands contact is likely to occur, wear suitable gloves tested according to EN374. Suitable gloves and protection garments should be worn.

Respiratory protection

Wear a protective mask in the presence of dust. Use the P2 Filter for solid particles.

8.2.3. Environment exposure controls

Do not pour waste waters directly into the environment.

SECTION 9: PHYSICAL AND CHEMICAL PROPERTIES

9.1. INFORMATION ON BASIC PHYSICAL AND CHEMICAL PROPERTIES

Physical state:	White solid crystalline
Colour:	White
Odour:	Odourless
Odour threshold:	No information available
pH:	2.2 (Solution 0.1 N)
Melting point:	169 °C at 1013 hPa (mbar)
Boiling point:	179.1 °C at 1013 hPa (mbar)
Flash point:	> 100 °C at 102.3 kPa (mbar)
Evaporation rate:	No information available
Flammability (solids, gases):	Non-flammable
Lower flash point:	No information available
Upper flash point:	No information available
Vapour pressure:	< 5 Pa at 20 °C
Vapour density:	No information available
Relative density (water=1):	1.76 g/cm ³ at 20°C

Solubility:	1,390 g/L at 20 °C.
Partition coefficient:	n-Octanol/water: Log Kow (Pow): - 1.91 at 20 °C
Decomposition temperature:	<i>No information available.</i>
Viscosity:	<i>No information available.</i>
Auto-Flammability in clouds:	490 °C a 1013 hPa (mbar)
K _{st} max:	49 bar m/s
(dP/dt) max – increasing pressure:	182 bar/s
Auto-Ignition temperature:	<i>No auto-ignition till the test maximum temperature (400 °C)</i>
Minimum ignition energy:	540 mJ
Oxidising properties:	<i>Not oxidising.</i>

9.2. OTHER INFORMATION

No data

SECTION 10: STABILITY AND REACTIVITY

10.1. REACTIVITY

Stable under normal conditions.

10.2. CHEMICAL STABILITY

The product is chemically stable under standard environmental conditions.

10.3. POSSIBILITY OF HAZARDOUS REACTIONS

Fluorine, metals, silver

10.4. CONDITIONS TO AVOID

Strong heating.

10.5. INCOMPATIBLE MATERIALS

No information available.

10.6. HAZARDOUS DECOMPOSITION PRODUCTS

No information available.

SECTION 11: TOXICOLOGICAL INFORMATION

11.1. INFORMATION ON TOXICOLOGICAL EFFECTS

ACUTE TOXICITY

Oral: LD50: > 2000 mg/kg bw for rat

Dermal: LD50: > 2000 mg/kg bw for rat

Value used for CSA:

LD50 (oral): 2000 mg/kg bw

LD50 (dermal): 2000 mg/kg bw

Justification for classification or non classification

According to Official Journal of the European Union 1272/2008 (CLP) dated December 16th 2008, tartaric acid is non-classified in the acute toxicity hazard categories. But it is necessary to emphasize that tartaric acid is classified in category 5 of acute oral toxicity in the GHS classification system.

SKIN IRRITATION:

A test of the registered substance was performed on skin irritation/corrosion *in vivo* according to OECD Guideline 404: acute dermal irritation/corrosion in a certified GLP lab. The study can be ranked according to the klimisch code as 1: reliable without restrictions. The results showed that no toxic effect was found. And other two *in vitro* studies also support this result. So the irritating effect of tartaric acid can be concluded as no irritating.

Value used for CSA: Skin irritation / corrosion: not irritating.

EYE IRRITATION:

An *in vitro* test of the registered substance was performed on eye irritation complying with OECD Guideline 437: Bovine Corneal Opacity and Permeability Test Method for identifying ocular corrosives and severe irritants. This study is regarded as key study as it can be ranked according to the klimisch code as 1: reliable without restrictions. And the test result showed that tartaric acid is highly irritating.

Value used for CSA: Eye irritation: highly irritating

SKIN SENSITISATION

The following information is taken into account for any hazard / risk assessment:

Skin sensitisation (OECD 429): not sensitising.

Value used for CSA: Not sensitising.

RESPIRATORY SENSITISATION

Values used for CSA: No data available.

REPEATED DOSE TOXICITY

NOAEL of repeated oral dose toxicity of tartaric acid is derived from the key study 004 through read across. In this study, Monosodium L(+) -tartrate was fed to rats in their diet for a total of two years at levels of 25600, 42240, 60160 and 76800 ppm and no adverse effect was observed in the highest concentration of L(+) -tartrate. So it is reasonable to choose 76800 ppm tartrate, which is equal to 2460 mg/kg bw/day, as NOAEL of tartaric acid. Furthermore, in the key study, the test material used was Monosodium L (+) -tartrate, a sodium salt of tartaric acid. It can be served as a read across study, because the basic chemical structures are the same in such two chemicals.

The following information is taken into account for any hazard / risk assessment:

No evidence of an adverse effect was seen in the dose of 3.1 g/kg bw/day and 4.1 g/kg bw/day L(+) -tartrate for male and female rats respectively, corresponding to 2.46 g/kg bw/day and 3.2 g/kg bw/day L(+) -tartaric acid for male and female rats respectively.

Value used for CSA (route: oral):

NOAEL: 2460 mg/kg bw/day (chronic; rat)

Justification for classification or non classification

The DNEL of repeated oral dose toxicity of tartaric acid is 2460 mg/kg bw/day, no specific organ toxicity was found here, so non-classification will be justified.

SINGLE DOSE TOXICITY

No data available

MUTAGENICITY

The FDA report, mutagenic evaluation of compound FDA 71-55, comprises several studies investigating genotoxicity of this substance in vitro and in vivo. In the in vitro studies, 4 host-mediated assays including two bacteria (*S. typhimurium*) and two yeast (*Saccharomyces cerevisiae*) tests, and a mammalian chromosome aberration test (Human embryonic lung cultures) were conducted at different concentration levels. In the in vivo studies, two dominant lethal tests and two mammalian bone marrow chromosome aberration tests were carried out in different series of concentrations in rats. No genetic toxicity was found in those tests in all investigated concentrations. So it can be concluded that L(+) -tartaric acid is non-mutagenic.

The following information is taken into account for any hazard / risk assessment: no genetic toxicity of tartaric acid was found through in vitro and in vivo experiments.

Value used for CSA: Genetic toxicity: negative

CARCINOGENICITY

No data available.

Combined chronic Toxicity/Carcinogenicity study equivalent or similar to OECD Guideline 453 is available under repeated dose toxicity.

REPRODUCTIVE TOXICITY

The FDA report, teratologic evaluation of FDA 71-55, summarised studies of the teratogenicity of tartaric acid in different species: mouse, rat, hamster and rabbit, using prenatal developmental toxicity test. It is found that administrations of the highest dosage, 274 mg/kg bw in mice, 181 mg/kg bw in rats, 225 mg/kg bw in hamsters and 215 mg/kg bw in rabbits, did not generate any teratogenic effects on tested animals. So these dose levels could be set as NOAELs in each individual test. In order to guarantee safety, also considering that the toxicokinetics of tartaric acid in rat is well studied, NOAEL of rat is chosen as the dose descriptor starting point for further calculation.

The following information is taken into account for any hazard/risk assessment: The FDA report, teratologic evaluation of FDA 71-55, includes 4 key studies carried out in different species investigating the developmental toxicity/teratogenicity. No teratogenic effect was found in these studies.

Value used for CSA (route: oral): NOAEL: 181 mg/kg bw/day

ASPIRATION HAZARD

No inhalation toxicity classification

SECTION 12: ECOLOGICAL INFORMATION

12.1 TOXICITY

ACUTE AQUATIC TOXICITY

The fish, daphnia, and algae acute aquatic toxicity levels are greater than 1 mg/L (96h LC50 (fish) > 100 mg/L, 48h EC50 (daphnia) = 93.3mg/L, and 72h ErC50 (algae) =51.4 mg/L). As a result, the substance does not meet the criteria for acute classification according to Regulation (EC) No. 1272/2008, Annex I section 4.1.

CHRONIC AQUATIC TOXICITY

The fish, daphnia, and algae acute aquatic toxicity levels are greater than 10 mg/l and lower than 100 mg/L (96h LC50 (fish) > 100 mg/L, 48h EC50 (daphnia) = 93.3mg/L, and 72h ErC50 (algae) =51.4 mg/L). As well, the substance is very soluble, ready biodegradable and has a Log Kow of -1.91. As a result, the substance does not meet the criteria for chronic classification according to Regulation (EC) No. 1272/2008, Annex I section 4.1.

According to Annex XIII of regulation 1907/2006/EC and according to the Guidance on information requirements and chemical safety assessment Chapter R.11 PBT assessment, a substance does not fulfil the criterion if there is no evidence of chronic toxicity and no classification as carcinogenic (Cat. 1, 2), mutagenic (Cat. 1, 2) or toxic for reproduction (Cat 1, 2, 3) considering human health. As the substance is not toxic and not classified for human health, these criteria are not fulfilled. Furthermore, the substance is not toxic for aquatic organisms.

12.2 PERSISTENCE AND DEGRADABILITY

According to Annex XIII of regulation 1907/2006/EC and according to the Guidance on information requirements and chemical safety assessment Chapter R.11 PBT assessment, a substance does not fulfil the criterion "persistent (P)" and "very persistent (vP)" if it is readily biodegradable. As the substance is shown to be readily biodegradable with a biodegradation of above 80% it is not regarded as persistent or very persistent.

12.3 BIOACCUMULATIVE POTENTIAL

According to Annex XIII of regulation 1907/2006/EC and according to the Guidance on information requirements and chemical safety assessment Chapter R.11 PBT assessment, a substance does not fulfil the criterion "bioaccumulative (B)" or "very bioaccumulative (vB)" if the BCF is below 2000 or the log Kow is below 4.5.

There is no experimental data on BCF. However, the log Kow is negative and below the criterion for bioaccumulation (log Kow 4.5). Therefore, it can be concluded that the substance is neither bioaccumulative nor very bioaccumulative.

12.4 MOBILITY IN SOIL

No data available

12.5 RESULTS OF PBT AND VPvB ASSESSMENT

The substance does not fulfil the criteria for PBT or vPvB properties.

12.6 OTHER ADVERSE EFFECTS

No more data

SECTION 13: DISPOSAL CONSIDERATIONS

13.1. WASTE TREATMENT METHODS

In general, the disposal of chemical residues is regulated in each European country by specific laws and regulations.

In Italy, the disposal must occur according to the laws in force and in compliance with local laws. Therefore, it is recommended to contact the Authorities in charge or specialised Companies authorised to provide indications on how to arrange the disposal.

Packing material must be disposed of in accordance with national regulations. Contaminated packing material must be handled with the same caution used for dangerous substances. Non-contaminated packing material can be treated or recycled as normal residues, unless otherwise indicated.

SECTION 14: TRANSPORT INFORMATION

14.1. UN NUMBER

Not classified as dangerous goods for transport.

14.2. UN PROPER SHIPPING NAME

Not classified as dangerous goods for transport.

14.3. TRANSPORT HAZARD CLASS(ES)

Road and Railway Transport:

Not classified as dangerous goods for transport.

Sea Transport:

Not classified as dangerous goods for transport.

Air Transport:

Not classified as dangerous goods for transport.

14.4. PACKING GROUP

Not classified as dangerous goods for transport.

14.5. ENVIRONMENTAL HAZARDS

Not classified as dangerous goods for transport.

14.6. SPECIAL PRECAUTIONS FOR USER

Not classified as dangerous goods for transport.

14.7. TRANSPORT IN BULK ACCORDING TO ANNEX II OF MARPOL AND THE IBC CODE

Not classified as dangerous goods for transport.

SECTION 15: REGULATORY INFORMATION

15.1. SAFETY, HEALTH AND ENVIRONMENTAL REGULATIONS/LEGISLATION SPECIFIC FOR THE SUBSTANCE OR MIXTURE

Authorisation pursuant to REACH Regulations:

It is not on the list of substances of very high concern (SVHC) applicable for the authorisation.

Restrictions on use pursuant to REACH Regulations:

It is not subject to restrictions pursuant to Title VII (Annex XVII, Appendix 2, paragraph 28)

15.2 CHEMICAL SAFETY ASSESSMENT

An assessment of the chemical safety has been carried out

16 OTHER INFORMATION

List of the relevant H hazard indications:

H318: Causes serious eye damage

Instructions on training:

Properly train those workers potentially exposed to this substance on the basis of the contents of this safety data sheet.

Main bibliographical references and data sources:

Registration Dossier of Tartaric Acid

Key of abbreviations and acronyms:

DNEL = Derived No Effect Level

DMEL = Derived Minimum Effect Level

EC50 = Median effective concentration

IC50 = Inhibition concentration, 50%

LC50 = Lethal concentration, 50%

LD50 = Median lethal dose

PNEC = Predicted No-Effect Concentration

PBT = Persistent, Bioaccumulative and Toxic substance

TLV®/TWA = Threshold limit value – time-weighted average

TLV®STEL = Threshold limit value – short-time exposure limit

vPvB = very Persistent and very Bioaccumulative

Revision date:

revision on 08/08/2019

Reason to change the data sheet:

general update

The data contained herein are the result of the best information at the time of publication. The company shall not be liable for any damage to persons or objects deriving from the improper use of the information disclosed in this document.