ThreAMINO® L-Threonine, Feed Grade 98,5%



 Material no.
 Version
 2.0 / US

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1. Identification

1.1. Product identifier

Trade name ThreAMINO®

L-Threonine, Feed Grade 98,5%

CAS-No. 72-19-5

1.2. Recommended use of the chemical and restrictions on use

Relevant applications identified Feed additive

1.3. Details of the supplier of the safety data sheet

Company Evonik Corporation USA

299 Jefferson Road

Parsippany, NJ 07054-0677

USA

Telephone 973-929-8000

Telefax 973-929-8040

Email address Product-Regulatory-Services@Evonik.com

1.4. 24 HOUR EMERGENCY TELEPHONE NUMBERS:

CHEMTREC - US &

800-424-9300

CANADA:

CHEMTREC MEXICO: 01-800-681-9531

CHEMTREC +1 703-527-3887 (collect calls accepted)

INTERNATIONAL:

Product Regulatory

973-929-8060

Services

2. Hazards identification

2.1. Classification of the substance or mixture

Classification according to Regulation 29CFR 1910.1200

Remarks

Not a hazardous substance or mixture.

2.2. Label elements

Statutory basis Classification according to Regulation 29CFR 1910.1200

Remarks Not a hazardous substance or mixture.

Supplemental hazard information / Label elements

Contains Residual , L-Threonine

The following percentage of the mixture consists of ingredient(s) with unknown acute toxicity: 99.5 %

2.3. Other hazards

May form explosive dust-air mixture.

Inhalation No hazard expected in normal use.
Skin No hazard expected in normal use.

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Eyes No hazard expected in normal use. Ingestion No hazard expected in normal use.

3. Composition/information on ingredients

• L-Threonine >= 98.5%

CAS-No. 72-19-5

Other information

This material is classified as not hazardous under OSHA regulations.

This product is intended for FDA regulated uses only.

4. First aid measures

4.1. Description of first aid measures

Inhalation

In case product dust is released: Possible discomfort: cough, sneezing

Move victims into fresh air.

Skin contact

No hazards which require special first aid measures.

Eye contact

Possible discomfort is due to foreign substance effect.

Rinse thoroughly with plenty of water keeping eyelid open.

In case of persistent discomfort: Consult an ophthalmologist.

Ingestion

Have the mouth rinsed with water.

After absorbing large amounts of substance

Consult a physician.

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

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

After absorbing large amounts of substance:

Acceleration of gastrointestinal passage

5. Fire-fighting measures

5.1. Extinguishing media

Suitable extinguishing media: Water, mist, Foam

Unsuitable extinguishing media: quenching powder, Carbon dioxide (CO2)

5.2. Special hazards arising from the substance or mixture

In the case of fire, the following hazardous smoke fumes may

be produced: carbon monoxide, carbon dioxide, nitric oxides, hydrocyanic acid.

In the event of fire and/or explosion do not breathe fumes.

5.3. Advice for firefighters

Contaminated fire-extinguishing water must be disposed of in accordance with the regulations issued by the appropriate local authorities.

Fire residues should be disposed of in accordance with the regulations.

In the event of fire, wear self-contained breathing apparatus.

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6. Accidental release measures

6.1. Personal precautions, protective equipment and emergency procedures

Wear personal protective equipment. Keep unauthorized persons away.

6.2. Environmental precautions

Obey relevant local, state, provincial and federal laws and regulations. Do not contaminate any lakes, streams, ponds, groundwater or soil.

6.3. Methods and material for containment and cleaning up

Absorb mechanically avoiding production of dust.

7. Handling and storage

7.1. Precautions for safe handling

Handle in accordance with good industrial hygiene and safety practice.

7.2. Conditions for safe storage, including any incompatibilities

Advice on protection against fire and explosion

Take precautionary measures against static charges, keep away from sources of ignition. Avoid dust formation.

Storage

Store in a cool and shaded area.

Keep containers dry and tightly closed to avoid moisture absorption and contamination.

German storage class

11 - Combustible Solids

Dust explosion class

St1

Method: VDI 3673
Maximum rate of pressure rise: 66 bar/s

Standardized max. rate of pressure increase, KSt: 66bar·m/s

8. Exposure controls/personal protection

8.1. Control parameters

exposure limit for dust		
CAS-No. Control parameters type of exposure	3 mg/m3 Respirable fraction. Suitable measuring processes are:	Time Weighted Average (TWA):(ACGIH)
	NIOSH method 0500 NIOSH method 0600	
Control parameters type of exposure	10 mg/m3 Inhalable particulate.	Time Weighted Average (TWA):(ACGIH)
Control parameters	15 mg/m3	Time Weighted Average (TWA) Permissible Exposure Limit (PEL):(OSHA Z1)
type of exposure	Total dust.	
Control parameters	5 mg/m3	Time Weighted Average (TWA) Permissible Exposure Limit (PEL):(OSHA Z1)
type of exposure	Respirable fraction.	

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Suitable measuring processes are:

NIOSH method 0500 NIOSH method 0600

DNEL/DMEL values

Remarks No substance-related safety assessment is necessary / has been conducted

for this product.

PNEC values

Remarks No substance-related safety assessment is necessary / has been conducted

for this product.

8.2. Exposure controls

Engineering measures

Use process enclosures, local exhaust ventilation or other engineering controls to control airborne exposure.

Take precautionary measures against static discharges. Earthing of equipment.

Personal protective equipment

Respiratory protection

A respiratory protection program that meets OSHA 1910.134 and ANSI Z88.2 or applicable federal/provincial requirements must be followed whenever workplace conditions warrant respirator use. NIOSH's "Respirator Decision Logic" may be useful in determining the suitability of various types of respirators.

Hand protection

Glove material Natural rubber (NR), for example, Cama Clean 708, Kächele-Cama Latex GmbH (KCL),

Germany

Material thickness 0.5 mm Break through time 8 h

Method DIN EN 374

Glove material Nitrile, for example, Dermatril 740, Kächele-Cama Latex GmbH (KCL), Germany

Material thickness 0.11 mm Break through time 8 h

Method DIN EN 374

The above mentioned hand protection is based on knowledge of the chemistry and anticipated uses of this product but it may not be appropriate for all workplaces. A hazard assessment should be conducted prior to use to ensure suitability of gloves for specific work environments and processes prior to use.

Eye protection

Safety glasses

Skin and body protection

No special protective equipment required.

Hygiene measures

Wash face and/or hands before break and end of work.

Cleanse and apply cream to skin after work.

Protective measures

Handle in accordance with good industrial hygiene and safety practice.

If there is the possibility of skin/eye contact, the indicated hand/eye/body protection should be used.

9. Physical and chemical properties

9.1. Information on basic physical and chemical properties

physical state solid

Colour white until light grey

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Form solid

Odour characteristic

Odour Threshold not determined

рΗ 5.0 - 6.5(25 °C) (25 g/l)

253 °C Melting point/range

decomposition

Boiling point/range not applicable

Flash point not applicable

solid

Evaporation rate No data available

Flammability (solid, gas) not highly flammable

Method: UN method N.1

Lower explosion limit 60 g/m³ dust:

VDI 3673 Method:

grain size < 63µm

Upper explosion limit No data available

Vapour pressure not applicable

Vapour density No data available

Relative vapour density no data available

Relative density No data available

Water solubility 85.7 g/l (20 °C)

> Method: **OECD Test Guideline 105**

Partition coefficient: n-

log Pow: -2.94

octanol/water Related to substance: pure substance

370 °C Autoignition temperature

> Method: VDI Guideline 2263 sheet 1

(BAM-furnace) for dust whirled up mean grain size

49µm

253 °C Thermal decomposition

TG (thermal gravimetric analysis)

Viscosity, dynamic not applicable

solid

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9.2. Other information

Explosiveness The product is susceptible to dust explosion.

Sublimation point 200 °C

Bulk density 585 - 715 kg/m3

glow temperature > 400 °C

Method: VDI 2263

Minimum ignition energy > 10 mJ (25 °C)

Classification: Normal combustability
Method: VDI Guideline 2263 sheet 1

mean grain size: 18 μm

sieve fraction with inductance

maximum absolute

9.6 bar

explosive pressure (with 1000 g/m³)

grain size < 63µm

Metal corrosion no data available

Burning number BZ 3 - local burning or smouldering with little or no spreading.

Method: Combustibility test in accordance with VDI 2263

10. Stability and reactivity

10.1. Reactivity

No further information available

10.2. Chemical stability

Stable under recommended storage conditions.

10.3. Possibility of hazardous reactions

Possibility of hazardous Dust can form an explosive mixture in air.

reactions

10.4. Conditions to avoid

See chapter

7.2. Conditions for safe storage, including any incompatibilities

10.5. Incompatible materials

Avoid contact with oxidizing substances.

10.6. Hazardous decomposition products

No hazardous decomposition products known.

11. Toxicological information

11.1. Information on toxicological effects

Acute oral toxicity NOEL Rat: 2000 mg/kg

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Method: OECD TG 423
Test substance: comparable product

LD50 Rat: > 5000 mg/kg

Method: OECD TG 423

Test substance: comparable product

Acute inhalation toxicity NOAEL Rat: 5.15 mg/l / 4 h

Method: OECD Test Guideline 403

Acute dermal toxicity Assessment: no data available

Skin irritation Rabbit

No skin irritation

Method: OECD Test Guideline 404

Eye irritation Rabbit

No eye irritation

Method: OECD Test Guideline 405

Sensitization Magnusson & Kligman Guinea pig: Does not cause skin sensitisation.

Method: OECD Test Guideline 406

Repeated dose toxicity Oral Rat(male/female)

Testing period: 28 d Subsequent observation 42 day

period:

NOAEL: > 1000 mg/kg

target organ/effect: no pathological changes

no data available

Method: OECD 407

Assessment of STOT single

exposure

Assessment: no data available

Assessment of STOT repeat

exposure

Risk of aspiration toxicity no data available

Gentoxicity in vitro Chromosome aberration test in vitro Human lymphocytes 625 - 5000 µg/ml

negative

Assessment:

Metabolic activation: with or without Method: OECD TG 473

Ames test Salmonella typhimurium <= 5000 μg/plate

negative

Metabolic activation: with or without
Method: OECD TG 471
Test substance: comparable product

Carcinogenicity no data available

carcinogenicity assessment Contains no carcinogenic substances as defined by NTP, IARC and/or

OSHA.

Toxicity to reproduction no data available

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Toxicological information on components

L-Threonine

Acute oral toxicity LD50 Rat: > 5000 mg/kg

Method: OECD Test Guideline 401

Acute inhalation toxicity LC0 Rat(male/female): > 5.15 mg/l / 4 h

Method: OECD Test Guideline 403

limit test (maximum concentration attainable in experiments) - No deaths

occurred.

Acute dermal toxicity Assessment: no data available

Skin irritation Rabbit

No skin irritation

Method: OECD Test Guideline 404

Eve irritation Rabbit

No eye irritation

Method: OECD Test Guideline 405

Sensitization Magnusson & Kligman Guinea pig: Does not cause skin sensitisation.

Method: OECD Test Guideline 406

Repeated dose toxicity Oral Rat(male/female)

Testing period: 28 d Subsequent observation 42 day

period:

NOAEL: > 1000 mg/kg

target organ/effect: no pathological changes

Method: OECD 407

Gentoxicity in vitro Chromosome aberration test in vitro Human lymphocytes 625 - 5000 µg/ml

negative

Metabolic activation: with or without Method: OECD TG 473

Ames test Salmonella typhimurium <= 5000 μg/plate

negative

Metabolic activation: with or without

Method: OECD TG 471

Test substance: comparable product

12. Ecological information

12.1. Toxicity

Toxicity in aquatic NOEC Daphnia magna: > 1000 mg/l / 48 h

invertebrates Method: OECD TG 202

Toxicity to algae EC50 static test Desmodesmus subspicatus (green algae): > 1000 mg/l

/ 72 h

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Toxicity to bacteria EC 80 nitrobacteria: 119 mg/l / 69 h

Method: literature

Method: literature

12.2. Persistence and degradability

Biodegradability Result: rapidly biodegradable

Method: QSAR-Method

12.3. Bioaccumulative potential

Bioaccumulation No data available

12.4. Mobility in soil

Mobility No data available

12.5. Other adverse effects

Further Information No further information available

13. Disposal considerations

13.1. Waste treatment methods

Product

Waste must be disposed of in accordance with federal, provincial and local regulations.

Uncleaned packaging

Packaging material should be recycled or disposed of in accordance with federal, state and local regulations.

14. Transport information

Not dangerous according to transport regulations.

14.1. UN number:

14.2. UN proper shipping name: --

14.3. Transport hazard class(es): --

14.4. Packing group: --

14.5. Environmental hazards (Marine --

pollutant):

14.6. Special precautions for user: Yes

Not dangerous according to transport regulations.

15. Regulatory information

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US Federal Regulations

OSHA

If listed below, chemical specific standards apply to the product or components:

None listed

Clean Air Act Section (112)

If listed below, components present at or above the de minimus level are hazardous air pollutants:

None listed

CERCLA Reportable Quantities

If listed below, a reportable quantity (RQ) applies to the product based on the percent of the named component:

None listed

SARA Title III Section 311/312 Hazard Categories

The product meets the criteria only for the listed hazard classes:

No SARA Hazards

SARA Title III Section 313 Reportable Substances

If listed below, components are subject to the reporting requirements of Section 313 of Title III of the Superfund Amendments and Reauthorization Act of 1986 and 40 CFR Part 372:

None listed

Toxic Substances Control Act (TSCA)

If listed below, non-proprietary substances are subject to export notification under Section 12 (b) of TSCA:

None listed

State Regulations

California Proposition 65

A warning under the California Drinking Water Act is required only if listed below:

None listed

International Chemical Inventory Status

Unless otherwise noted, this product is in compliance with the inventory listing of the countries shown below. For information on listing for countries not shown, contact the Product Regulatory Services Department.

Europe (EINECS/ELINCS) listed/registered USA (TSCA) listed/registered

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Canada (DSL) listed/registered
Australia (AICS) listed/registered
Japan (MITI) listed/registered
Korea (TCCL) listed/registered
Philippines (PICCS) listed/registered
China listed/registered
Switzerland listed/registered

An employer using HMIS/NFPA labeling must through training ensure that its employees are fully aware of the hazards of the chemicals used.

HMIS Ratings

Health: 0
Flammability: 1
Physical Hazard: 0

16. Other information

Further information

Revision date 04/22/2015

Changes since the last version are highlighted in the margin. This version replaces all previous versions.

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Legend

ACC American Chemistry Council

ACGIH American Conference of Governmental Industrial Hygenists

ACS Advisory Committee on Sustainability

ADI Acceptable Daily Intake

ASTM American Society for Testing and Materials

ATP Adaptation to Technical Progress

BCF Bioconcentration factor
BOD Biochemical oxygen demand

c.c. closed cup
CAO Cargo Aircraft Only
Carc Carcinogen

CAS Chemical Abstract Services

CDN Canada

CEPA Canadian Environmental Protection Act

CERCLA Comprehensive Environmental Response – Compensation and Liability Act

CFR Code of Federal Regulations

CMR carcinogenic-mutagenic-toxic for reproduction

COD Chemical oxygen demand

DIN German Institute for Standardization
DMEL Derived minimum effect level

DNEL Derived no effect level
DOT Department of Transportation
EC50 half maximal effective concentration
EPA Environmental Protection Agency
ErC50 Reduction of Growth Rate
ERG Emergency Response Guide Book
FDA Food and Drug Administration

Globally Harmonized System of Classification and Labelling of Chemicals (GHS)

GLP Good Laboratory Practice
GMO Genetic Modified Organism
HCS Hazard Communication Standard
HMIS Hazardous Materials Identification System
IARC International Agency for Research on Cancer
IATA International Air Transport Association

IBC Intermediate Bulk Container

ICAO-TI International Civil Aviation Organization- Technical Instructions

ICCA International Council of Chemical Association

ID Identification number

IMDG International Maritime Dangerous Goods
IUPAC International Union of Pure and Applied Chemistry
ISO International Organization For Standardization

LC50 50 % Lethal Concentration

LD50 50 % Lethal Dose **L(E)C50** LC50 or EC50

LOAEL Lowest observed adverse effect level

LOEL Lowest observed effect level

MARPOL International Convention for the Prevention of Pollution from Ships

NFPA National Fire Protection Association
NOAEL No observed adverse effect level
NOEC no observed effect concentration

NOEL no observed effect level

o. c. open cup

OECD Organisation for Economic Cooperation and Development

OEL Occupational Exposure Limit

OSHA Occupational Safety and Health Administration

PBT Persistent, bioaccumulative, toxic
PEC Predicted effect concentration
PNEC Predicted no effect concentration

RQ Reportable Quantity SDS Safety Data Sheet

STOT Specific Target Organ Toxicity

UN United Nations

vPvB very persistent, very bioaccumulative

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volatile organic compounds Workplace Hazardous Materials Information System World Health Organization voc WHMIS WHO