

SAFETY DATA SHEET**TrypAMINO® L-Tryptophan, Feed Grade 98%**

Material no.		Version	2.0 / US
Specification	159641	Revision date	04/22/2015
Order Number		Print Date	04/22/2015
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1. Identification**1.1. Product identifier**

Trade name TrypAMINO®
 L-Tryptophan, Feed Grade 98%

CAS-No. 73-22-3

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

Teletax 973-929-8040

Email address Product-Regulatory-Services@Evonik.com

1.4. 24 HOUR EMERGENCY TELEPHONE NUMBERS:

CHEMTREC - US & CANADA: 800-424-9300

CHEMTREC MEXICO: 01-800-681-9531

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

Product Regulatory Services : 973-929-8060

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.

2.3. Other hazards

May cause eye and skin irritation.

Inhalation No hazard expected in normal use.

Skin Possibly irritating.

Eyes Possibly irritating.

Ingestion No hazard expected in normal use.

Dust can form an explosive mixture in air.

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3. Composition/information on ingredients**• L-Tryptophan** >= 98%

CAS-No. 73-22-3

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.

If symptoms persist, seek medical advice (show this MSDS where possible).

Skin contact

Wash off with plenty of water.

Consult a doctor in the event of permanent skin irritation.

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

If symptoms persist, seek medical advice (show this MSDS where possible).

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**

Formation of flammable or explosive dust/air mixtures possible.

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

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

flammable smouldering gases

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.

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

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

Do not allow the product into the following compartments:, groundwater, surface water

6.3. Methods and material for containment and cleaning up

Absorb mechanically avoiding production of dust. Keep away from sources of ignition - No smoking. To absorb spilled substance an approved industrial vacuum cleaner is recommended. If necessary, the spilled substance should be moistened.

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.

Advice on common storage

Incompatible with strong acids and strong oxidizing agents.

German storage class

11 - Combustible Solids

Dust explosion class

St1

Method: VDI Guideline 2263 sheet 1

Maximum rate of pressure rise: 122 bar/s

Standardized max. rate of pressure increase, KSt: 122bar·m/s

8. Exposure controls/personal protection**8.1. Control parameters**

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

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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. 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 measures to prevent the build up of electrostatic charge.

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

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**

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physical state	solid		
Colour	white until light grey		
Odour	characteristic		
Odour Threshold	not determined		
pH	5.5 - 7.0	(25 g/l)	(25 °C)
Melting point/range	279 - 292 °C		
	Method: DIN 51004 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	dust:	30 g/m ³	(34 µm)
	Method:	VDI 2263	
Upper explosion limit	not to be determined		
Vapour pressure	< 0.0000001 hPa	(25 °C)	
	Method:	calculated literature	
Vapour density	No data available		
Relative vapour density	no data available		
Relative density	1.39	(22 °C)	
	Method:	OECD Test Guideline 109	
Water solubility	11.4 g/l	(25 °C)	
	Related to substance:	pure substance	
Partition coefficient: n-octanol/water	log Pow:	-1.11	
	Related to substance:	pure substance	
Autoignition temperature	490 °C		
	Method:	VDI Guideline 2263 sheet 1 for dust whirled up (BAM-furnace) grain size < 63µm	

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Thermal decomposition 290 - 292 °C
TG (thermal gravimetric analysis)
(literature value)

Viscosity, dynamic not applicable
solid

9.2. Other Information

Explosiveness The product is susceptible to dust explosion.

Minimum ignition energy 3 - 10 mJ (25 °C)
Classification: Remarkable combustability
Method: VDI Guideline 2263 sheet 1
mean grain size: 34 µm
sieve fraction
without inductance

maximum absolute explosive pressure 8.5 bar
Method: VDI 2263
grain size
< 63µm

Metal corrosion no data available

Burning number BZ 2 - briefly ignites and rapidly extinguishes.
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 reactions Dust can form an explosive mixture in air.

10.4. Conditions to avoid

See chapter

7.2. Conditions for safe storage, including any incompatibilities

10.5. Incompatible materials

No further information available

10.6. Hazardous decomposition products

No hazardous decomposition products known.

11. Toxicological information**11.1. Information on toxicological effects**

Acute oral toxicity NOAEL Rat: 2000 mg/kg

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	Method:	OECD TG 423
	LD50 Rat:	> 5000 mg/kg
	Method:	OECD TG 423
Acute inhalation toxicity	NOAEL Rat:	5.17 mg/l / 4 h
	Method:	OECD Test Guideline 403
Acute dermal toxicity	Assessment:	no data available
Skin irritation	Rabbit:	500 mg / 4 h
	No skin irritation	
	Method:	OECD Test Guideline 404
Eye irritation	Rabbit:	100 mg
	No eye irritation	
	Method:	OECD Test Guideline 405
Sensitization	Local Lymphnode Assay Mouse:	Does not cause skin sensitisation.
	Method:	OECD TG 429
Repeated dose toxicity	Oral Rat(male/female)	
	Testing period:	90 d
	Subsequent observation period:	28 day
	NOAEL:	> 3764 mg/kg
	Method:	OECD TG 408
Assessment of STOT single exposure	Assessment:	no data available
Assessment of STOT repeat exposure	Assessment:	no data available
Risk of aspiration toxicity		no data available
Gentoxicity in vitro	Cytogenetic test V79 V 79 cells (Chinese hamster)	312,5 - 5000 µg/ml
	negative	
	Metabolic activation:	with
	Method:	OECD TG 476
	HGPRT-Test	
	Cytogenetic test V79 V 79 cells (Chinese hamster)	187,5 - 3000 µg/ml
	negative	
	Metabolic activation:	without
	Method:	OECD TG 476
	HGPRT-Test	
	Ames test Salmonella typhimurium 0,316 - 5000 µg/plate	
	negative	
	Metabolic activation:	with or without
	Method:	OECD TG 471
	Chromosome aberration test in vitro human lymphocytes	625 - 5000 µg/ml
	negative	
	Metabolic activation:	with or without
	Method:	OECD TG 473

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Carcinogenicity Oral Rat(male/female): 546 day
 Number of exposures: 5 days/weeks
 Subsequent observation period: 182 - 189 day
 Method: OECD Test Guideline 451
 Did not show carcinogenic effects in animal experiments.
 literature

carcinogenicity assessment Animal testing did not show any carcinogenic effects.

 Contains no carcinogenic substances as defined by NTP, IARC and/or OSHA.

Toxicity to reproduction no evidence of reproductiontoxic properties
 90 days study
 OECD TG 408

Teratogenicity Oral Rat / 20 days
 Number of exposures: continuous
 Test period: 20 days
 NOAEL (No Observed Adverse Effect Level) 7150 mg/kg
 teratogenesis:
 NOAEL maternal (No Observed Adverse Effect Level): 2850 mg/kg
 Method: OECD TG 414
 literature

Toxicological information on components**L-Tryptophan**

Acute oral toxicity LD50 Rat(female): > 2000 mg/kg
 Method: OECD TG 423
 Assessment: The substance or mixture has no acute oral toxicity
 No deaths observed.
 (limit test)

LD50 Mouse: > 5000 mg/kg
 No deaths occurred.

Acute inhalation toxicity LC50 Rat(male and female): > 5.17 mg/l / 4 h / dust/mist
 Method: OECD Test Guideline 403
 Assessment: The substance or mixture has no acute inhalation toxicity
 No deaths observed.

Acute dermal toxicity Assessment: no data available

Skin irritation Rabbit: 500 mg
 No skin irritation
 Method: OECD Test Guideline 404

Eye irritation Rabbit: 100 mg
 No eye irritation
 Method: OECD Test Guideline 405

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Sensitization	Local Lymphnode Assay Mouse: Did not cause sensitisation on laboratory animals. Method: OECD TG 429
Repeated dose toxicity	Oral Rat(male and female) / 90-day Number of exposures: continuous NOAEL: 3764 mg/kg Method: OECD TG 408
	Oral Rat(male and female) / 28-day Number of exposures: continuous NOAEL: 1188 - 1271 mg/kg Method: OECD Test Guideline 407
Gentoxicity in vitro	chromosomal aberration Human lymphocytes negative Metabolic activation: with or without Method: OECD TG 473
Carcinogenicity	Oral Rat(male and female) Number of exposures: 5 days/weeks Subsequent observation period: 182 - 189 day Method: OECD Test Guideline 451
carcinogenicity assessment	Animal testing did not show any carcinogenic effects.
Teratogenicity	Oral Rat / 20 days Number of exposures: continuous Test period: 20 days NOAEL (No Observed Adverse Effect Level) teratogenesis: ca. 7150 mg/kg NOAEL maternal (No Observed Adverse Effect Level): ca. 2850 mg/kg Method: OECD TG 414 literature

12. Ecological information**12.1. Toxicity**

Toxicity in aquatic invertebrates	NOEC Daphnia magna: 100 mg/l / 48 h Method: OECD TG 202
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12.2. Persistence and degradability

Biodegradability	Result: rapidly biodegradable Method: OECD TG 301 B
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12.3. Bioaccumulative potential

Bioaccumulation	No data available
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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**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

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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
Canada (DSL)	listed/registered
Australia (AICS)	listed/registered
Japan (MITI)	listed/registered
Korea (TCCL)	listed/registered
Philippines (PICCS)	listed/registered
China	listed/registered
Switzerland	not 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

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Health : 1
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 Hygienists
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
GHS	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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voc	volatile organic compounds
WHMIS	Workplace Hazardous Materials Information System
WHO	World Health Organization