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### IDENTIFICATION OF THE SUBSTANCE/PREPARATION AND THE COMPANY/UNDERTAKING

**Product Identifier** 

Material Name: Ethosuximide Capsules (Greenstone LLC)

**Trade Name:** Not applicable **Chemical Family:** Mixture

Relevant Identified Uses of the Substance or Mixture and Uses Advised Against

Intended Use: Pharmaceutical product used as anticonvulsant

Details of the Supplier of the Safety Data Sheet

**Greenstone LLC** 100 Route 206 North Peapack, NJ 07977 800-435-7095

**Emergency telephone number:** CHEMTREC (24 hours): 1-800-424-9300

# **HAZARDS IDENTIFICATION**

# Classification of the Substance or Mixture

**GHS - Classification** 

Germ Cell Mutagenicity: Category 2 Reproductive Toxicity: Category 1B

**EU Classification:** 

EU Indication of danger: Harmful

Toxic to Reproduction: Category 2

Mutagenic: Category 3

EU Risk Phrases:

R22 - Harmful if swallowed.

R61 - May cause harm to the unborn child. R68 - Possible risk of irreversible effects.

**Label Elements** 

Signal Word: Danger

**Hazard Statements:** H360D - May damage the unborn child

H341 - Suspected of causing genetic defects

**Precautionary Statements:** P202 - Do not handle until all safety precautions have been read and understood

P281 - Use personal protective equipment as required

P308 + P313 - IF exposed or concerned: Get medical attention/advice

P405 - Store locked up

P501 - Dispose of contents/container in accordance with all local and national regulations

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Other Hazards

**Australian Hazard Classification** (NOHSC):

No data available

Hazardous Substance. Non-Dangerous Goods.

Note:

This document has been prepared in accordance with standards for workplace safety, which require the inclusion of all known hazards of the product or its ingredients regardless of the potential risk. The precautionary statements and warnings included may not apply in all cases. Your needs may vary depending upon the potential for exposure in your workplace.

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### 3. COMPOSITION/INFORMATION ON INGREDIENTS

#### **Hazardous**

Ingredient	CAS Number	EU EINECS/ELINCS List	EU Classification	GHS Classification	%
Glycerin, USP	56-81-5	200-289-5	Not Listed	Not Listed	*
Ethosuximide	77-67-8	201-048-7	Xn, R22; Repr. Cat.2,R61; Mut. Cat.3,R68	Acute Tox. 4, H302; Repr. 1B, H360D; Muta. 2, H341	32-35

Ingredient	CAS Number	EU EINECS/ELINCS List	EU Classification	GHS Classification	%						
						Gelatin	9000-70-8	232-554-6	Not Listed	Not Listed	*
						D & C yellow No. 10	8004-92-0	Not Listed	Not Listed	Not Listed	*
Sorbitol solution	50-70-4	200-061-5	Not Listed	Not Listed	*						
Polyethylene glycol 400	25322-68-3	Not Listed	Not Listed	Not Listed	*						
FD & C Red No. 3 (E 127)	16423-68-0	240-474-8	Not Listed	Not Listed	*						

**Additional Information:** 

\* Proprietary

Ingredient(s) indicated as hazardous have been assessed under standards for workplace safety. In accordance with 29 CFR 1910.1200, the exact percentage composition of this mixture has been withheld as a trade secret.

For the full text of the CLP/GHS abbreviations mentioned in this Section, see Section 16

### 4. FIRST AID MEASURES

**Description of First Aid Measures** 

**Eye Contact:** 

Flush with water while holding eyelids open for at least 15 minutes. Seek medical attention

immediately.

**Skin Contact:** Remove contaminated clothing. Flush area with large amounts of water. Use soap. Seek

medical attention.

Never give anything by mouth to an unconscious person. Wash out mouth with water. Do not Ingestion:

induce vomiting unless directed by medical personnel. Seek medical attention immediately.

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**Inhalation:** Remove to fresh air and keep patient at rest. Seek medical attention immediately.

Most Important Symptoms and Effects, Both Acute and Delayed

Symptoms and Effects of For information on potential signs and symptoms of exposure, See Section 2 - Hazards

**Exposure:** Identification and/or Section 11 - Toxicological Information.

Medical Conditions None known

Aggravated by Exposure:

Indication of the Immediate Medical Attention and Special Treatment Needed

Notes to Physician: None

### 5. FIRE FIGHTING MEASURES

**Extinguishing Media:** Use carbon dioxide, dry chemical, or water spray.

Special Hazards Arising from the Substance or Mixture

**Hazardous Combustion** Formation of toxic gases is possible during heating or fire.

**Products:** 

Fire / Explosion Hazards: Not applicable

**Advice for Fire-Fighters** 

During all fire fighting activities, wear appropriate protective equipment, including self-contained breathing apparatus.

### 6. ACCIDENTAL RELEASE MEASURES

#### Personal Precautions, Protective Equipment and Emergency Procedures

Personnel involved in clean-up should wear appropriate personal protective equipment (see Section 8). Minimize exposure.

#### **Environmental Precautions**

Place waste in an appropriately labeled, sealed container for disposal. Care should be taken to avoid environmental release.

#### Methods and Material for Containment and Cleaning Up

Measures for Cleaning /

Collecting:

Contain the source of spill if it is safe to do so. Collect spilled material by a method that controls dust generation. A damp cloth or a filtered vacuum should be used to clean spills of

dry solids. Clean spill area thoroughly.

**Additional Consideration for** 

Large Spills:

Non-essential personnel should be evacuated from affected area. Report emergency situations immediately. Clean up operations should only be undertaken by trained personnel.

### 7. HANDLING AND STORAGE

#### **Precautions for Safe Handling**

If tablets or capsules are crushed and/or broken, avoid breathing dust and avoid contact with eyes, skin, and clothing. Wash thoroughly after handling. Releases to the environment should be avoided. Review and implement appropriate technical and procedural waste water and waste disposal measures to prevent occupational exposure or environmental releases. Potential points of process emissions of this material to the atmosphere should be controlled with dust collectors, HEPA filtration systems or other equivalent controls.

Conditions for Safe Storage, Including any Incompatibilities

Storage Conditions: Store as directed by product packaging.

**Specific end use(s):** Pharmaceutical drug product

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### 8. EXPOSURE CONTROLS / PERSONAL PROTECTION

#### **Control Parameters**

Refer to available public information for specific member state Occupational Exposure Limits.

Glycerin, USP

10 mg/m<sup>3</sup> **Australia TWA** 10 mg/m<sup>3</sup> **Belgium OEL - TWA** Czech Republic OEL - TWA 10 ma/m<sup>3</sup> Estonia OEL - TWA 10 mg/m<sup>3</sup> 20 mg/m<sup>3</sup> **Finland OEL - TWA** France OEL - TWA 10 mg/m<sup>3</sup> Germany (DFG) - MAK 50 mg/m<sup>3</sup> **Greece OEL - TWA** 10 mg/m<sup>3</sup> **Ireland OEL - TWAs** 10 mg/m<sup>3</sup> **OSHA - Final PELS - TWAs:** 15 mg/m<sup>3</sup> 10 mg/m<sup>3</sup> **Poland OEL - TWA** Portugal OEL - TWA 10 mg/m<sup>3</sup> 10 mg/m<sup>3</sup> Spain OEL - TWA

Polyethylene glycol 400

 Austria OEL - MAKs
 1000 mg/m³

 Germany - TRGS 900 - TWAs
 1000 mg/m³

**Germany (DFG) - MAK** 1000 mg/m³ average molecular weight 200-600

Slovakia OEL - TWA 1000 mg/m³
Slovenia OEL - TWA 1000 mg/m³

**Ethosuximide** 

Manufacturer OEL: 2mg/m<sup>3</sup>

**Exposure Controls** 

**Engineering Controls:** Engineering controls should be used as the primary means to control exposures. General room ventilation is adequate unless the process generates dust, mist or fumes. Keep airborne

contamination levels below the exposure limits listed above in this section.

Personal Protective Equipment: Refer to applicable national standards and regulations in the selection and use of personal

protective equipment (PPE).

Hands: Impervious gloves are recommended if skin contact with drug product is possible and for bulk

processing operations.

**Eyes:** Wear safety glasses or goggles if eye contact is possible.

**Skin:** Impervious protective clothing is recommended if skin contact with drug product is possible and

for bulk processing operations.

Respiratory protection: If the applicable Occupational Exposure Limit (OEL) is exceeded, wear an appropriate

respirator with a protection factor sufficient to control exposures to below the OEL.

# 9. PHYSICAL AND CHEMICAL PROPERTIES

Physical State: Capsule Color: Orange

Odor: No data available. Odor Threshold: No data available.

Molecular Formula: Mixture Molecular Weight: Mixture

Solvent Solubility: No data available Water Solubility: No data available

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### 9. PHYSICAL AND CHEMICAL PROPERTIES

pH: No data available.

Melting/Freezing Point (°C): No data available

Boiling Point (°C): No data available.

Partition Coefficient: (Method, pH, Endpoint, Value)

D & C yellow No. 10 No data available

FD & C Red No. 3 (E 127)

No data available

Polyethylene glycol 400

No data available

Glycerin, USP

No data available

Sorbitol solution

No data available

Gelatin

No data available **Ethosuximide**No data available

Decomposition Temperature (°C): No data available.

Evaporation Rate (Gram/s):

Vapor Pressure (kPa):

Vapor Density (g/ml):

Relative Density:

No data available

Flammablity:

Autoignition Temperature (Solid) (°C):

Flammability (Solids):

Flash Point (Liquid) (°C):

Upper Explosive Limits (Liquid) (% by Vol.):

Lower Explosive Limits (Liquid) (% by Vol.):

No data available
No data available

# 10. STABILITY AND REACTIVITY

Reactivity: No data available

Chemical Stability: Stable under normal conditions of use.

Possibility of Hazardous Reactions

Oxidizing Properties: No data available

**Conditions to Avoid:** Fine particles (such as dust and mists) may fuel fires/explosions. **Incompatible Materials:** As a precautionary measure, keep away from strong oxidizers

Hazardous Decomposition No data available

**Products:** 

### 11. TOXICOLOGICAL INFORMATION

Information on Toxicological Effects

General Information: The information included in this section describes the potential hazards of the individual

ingredients.

Short Term: May be harmful if swallowed. (based on animal data)

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### 11. TOXICOLOGICAL INFORMATION

#### **Known Clinical Effects:**

Effects reported during clinical use included vomiting and diarrhea. Central nervous system effects such as dizziness, headache, insomnia, irritability and weakness have also been reported. Clinical use of this drug has caused decreased blood cell count, increased eosinophils in blood or tissue (eosinophilia), skin rash, Stevens Johnson Syndrome (epidermal necrosis and exfoliative dermatitis) May cause adverse effects on the developing fetus.

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### Acute Toxicity: (Species, Route, End Point, Dose)

#### D & C yellow No. 10

Rat Oral LD50 2000 mg/kg

### FD & C Red No. 3 (E 127)

Rat Oral LD50 1840 mg/kg Mouse Oral LD50 1264mg/kg

#### Glycerin, USP

Mouse Oral LD50 4090 mg/kg Rat Oral LD50 12.6 g/kg Rabbit Dermal LD50 > 10 g/kg Rat Inhalation LC50 1hr > 570 mg/m $^3$ Rat Dermal LD 50 > 21.9 g/kg

#### Sorbitol solution

Rat Oral LD50 15,900 mg/kg Mouse Oral LD50 17,800mg/kg

#### **Ethosuximide**

Mouse Oral LD50 1530 mg/kg Rat Oral LD50 1950mg/kg Mouse Intravenous LD50 780mg/kg

Mouse Intravenous LD50 780mg/kg
Mouse Intravenous LD50 1070mg/kg

**Acute Toxicity Comments:** 

A greater than symbol (>) indicates that the toxicity endpoint being tested was not achievable at the highest dose used in the test.

### Irritation / Sensitization: (Study Type, Species, Severity)

### Polyethylene glycol 400

Eye Irritation Rabbit Mild Skin Irritation Rabbit Mild

Glycerin, USP

Eye Irritation Rabbit Mild

### Repeated Dose Toxicity: (Duration, Species, Route, Dose, End Point, Target Organ)

### **Ethosuximide**

3 Month(s) Dog Oral 100 mg/kg/day LOAEL Liver 26 Week(s) Rat Oral 676 mg/kg/day NOAEL None identified 26 Week(s) Dog Oral 100 mg/kg/day NOAEL None identified 26 Week(s) Oral 200 mg/kg/day NOAEL None identified Monkey 136 mg/kg/day 1 Year(s) Mouse Oral LOAEL Liver

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### 11. TOXICOLOGICAL INFORMATION

Reproduction & Developmental Toxicity: (Study Type, Species, Route, Dose, End Point, Effect(s))

**Ethosuximide** 

Embryo / Fetal Development Rat 60 mg/kg/day LOEL Teratogenic

2 Generation Reproductive Toxicity Rat Oral 0.2 % LOAEL Not Teratogenic, Embryotoxicity

Embryo / Fetal Development Mouse Oral 60 mg/kg/day LOAEL Teratogenic

Prenatal & Postnatal Development Mouse Oral 50 mg/mL NOAEL Embryotoxicity, Reproductive toxicity,

Developmental toxicity

Genetic Toxicity: (Study Type, Cell Type/Organism, Result)

**Ethosuximide** 

In Vitro Cytogenetics Human Negative

In Vivo Micronucleus Mouse Bone Marrow Positive

Carcinogen Status: None of the components of this formulation are listed as a carcinogen by IARC, NTP or OSHA.

# 12. ECOLOGICAL INFORMATION

Environmental Overview: Environmental properties have not been investigated. Releases to the environment should be

avoided.

**Toxicity:** 

Aquatic Toxicity: (Species, Method, End Point, Duration, Result)

Glycerin, USP

Oncorhynchus mykiss (Rainbow Trout) LC50 96 Hours 50 mg/L Daphnia magna (Water Flea) EC50 24 Hours >500 mg/L

Aquatic Toxicity Comments: A greater than symbol (>) indicates that aquatic toxicity was not observed at the maximum

dose tested.

Persistence and Degradability: No data available

Bio-accumulative Potential: No data available

Mobility in Soil: No data available

### 13. DISPOSAL CONSIDERATIONS

Waste Treatment Methods: Dispose of waste in accordance with all applicable laws and regulations. Member State

specific and Community specific provisions must be considered. Considering the relevant known environmental and human health hazards of the material, review and implement appropriate technical and procedural waste water and waste disposal measures to prevent occupational exposure and environmental release. It is recommended that waste minimization be practiced. The best available technology should be utilized to prevent environmental

releases. This may include destructive techniques for waste and wastewater.

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### 14. TRANSPORT INFORMATION

The following refers to all modes of transportation unless specified below.

Not regulated for transport under USDOT, EUADR, IATA, or IMDG regulations.

# 15. REGULATORY INFORMATION

Safety, Health and Environmental Regulations/Legislation Specific for the Substance or Mixture

Canada - WHMIS: Classifications
WHMIS hazard class:
Class D, Division 2, Subdivision A



Glycerin, USP

CERCLA/SARA 313 Emission reporting

California Proposition 65

Not Listed

Not Listed

Not Listed

Not Present

Inventory - United States TSCA - Sect. 8(b)

Australia (AICS):

Present

**REACH - Annex V - Exemptions from the obligations of Register:**Present if not chemically modified, except they meet the criteria for classification as dangerous according to Directive 67/548/EEC,

except those only classified as flammable [R10], as a skin irritant [R38] or as an eye irritant [R36], except they are persistent,

bioaccumulative, and toxic or very persistent and very bioaccumulative in accordance with the criteria set out in Annex

232-554-6

XIII, except they were identified in accordance with Article 59[1] at least two years previously as substances giving rise to an

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equivalent level of concern

EU EINECS/ELINCS List 200-289-5

Gelatin

CERCLA/SARA 313 Emission reporting

California Proposition 65

Inventory - United States TSCA - Sect. 8(b)

Australia (AICS):

Not Listed

Not Listed

Present

Present

D & C yellow No. 10

**EU EINECS/ELINCS List** 

CERCLA/SARA 313 Emission reporting

California Proposition 65

Inventory - United States TSCA - Sect. 8(b)

Not Listed
Present

Australia (AICS): Present

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### **15. REGULATORY INFORMATION**

EU EINECS/ELINCS List Not Listed

Sorbitol solution

CERCLA/SARA 313 Emission reporting

California Proposition 65

Inventory - United States TSCA - Sect. 8(b)

Australia (AICS):

REACH - Annex IV - Exemptions from the

Not Listed
Present
Present

obligations of Register:

EU EINECS/ELINCS List 200-061-5

Polyethylene glycol 400

CERCLA/SARA 313 Emission reporting

California Proposition 65

Inventory - United States TSCA - Sect. 8(b)

Australia (AICS):

Standard for the Uniform Scheduling

Not Listed

Not Listed

Present

Present

Schedule 3

for Drugs and Poisons:

EU EINECS/ELINCS List Not Listed

FD & C Red No. 3 (E 127)

CERCLA/SARA 313 Emission reporting

California Proposition 65

Inventory - United States TSCA - Sect. 8(b)

Australia (AICS):

Present

EU EINECS/ELINCS List

Not Listed

Not

**Ethosuximide** 

CERCLA/SARA 313 Emission reporting

California Proposition 65

Australia (AICS):

Standard for the Uniform Scheduling

For Proposition 95

Not Listed

Not Listed

Not Listed

Not Listed

Schedule 4

for Drugs and Poisons:

EU EINECS/ELINCS List 201-048-7

### 16. OTHER INFORMATION

# Text of CLP/GHS Classification abbreviations mentioned in Section 3

Acute toxicity, oral-Cat.4; H302 - Harmful if swallowed Reproductive toxicity-Cat.1B; H360D - May damage the unborn child

Germ cell mutagenicity-Cat.2; H341 - Suspected of causing genetic defects

Mutagenic: Category 3

Toxic to Reproduction: Category 2

Xn - Harmful

R22 - Harmful if swallowed.

R61 - May cause harm to the unborn child. R68 - Possible risks of irreversible effects.

**Data Sources:** The data contained in this SDS may have been gathered from confidential internal sources, raw material suppliers, or from the published literature.

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Product Stewardship Hazard Communication

Prepared by: Global Environment, Health, and Safety Operations

It is believed that the information contained in this Material Safety Data Sheet is accurate, and while it is provided in good faith, it is without a warranty of any kind, expressed or implied. If data for a hazard are not included in this document there is no known information at this time

**End of Safety Data Sheet** 

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