



# SAFETY DATA SHEET

Revision date: 05-Oct-2015

Version: 3.0

Page 1 of 10

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

### Product Identifier

**Material Name:** Glipizide XL Extended-Release Tablets (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 antidiabetic agent

### 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

## 2. HAZARDS IDENTIFICATION

### Classification of the Substance or Mixture

**GHS - Classification** Not classified as hazardous

### Label Elements

**Signal Word:** Not required

**Hazard Statements:** Not classified in accordance with international standards for workplace safety.

### Other Hazards

**Note:** No data available  
This document has been prepared in accordance with standards for workplace safety, which requires the inclusion of all known hazards of the product or its ingredients regardless of the potential risk. The precautionary statements and warning included may not apply in all cases. Your needs may vary depending upon the potential for exposure in your workplace.

## 3. COMPOSITION/INFORMATION ON INGREDIENTS

### Hazardous

Ingredient	CAS Number	EU EINECS/ELINCS List	GHS Classification	%
Glipizide	29094-61-9	249-427-6	Not Listed	<5
Ferric oxide red	1309-37-1	215-168-2	Not Listed	*
Magnesium stearate	557-04-0	209-150-3	Not Listed	*

Ingredient	CAS Number	EU EINECS/ELINCS List	GHS Classification	%
Polyethylene oxide NF	25322-68-3	Not Listed	Not Listed	*
Hydroxypropyl methylcellulose	9004-65-3	Not Listed	Not Listed	*

## SAFETY DATA SHEET

Material Name: Glipizide XL Extended-Release Tablets  
(Greenstone LLC)  
Revision date: 05-Oct-2015

Page 2 of 10

Version: 3.0

Sodium chloride	7647-14-5	231-598-3	Not Listed	*
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**Additional Information:** \* Proprietary  
Ingredient(s) indicated as hazardous have been assessed under standards for workplace safety.

### 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.

**Ingestion:** Never give anything by mouth to an unconscious person. Wash out mouth with water. Do not induce vomiting unless directed by medical personnel. Seek medical attention immediately.

**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 Exposure:** For information on potential signs and symptoms of exposure, See Section 2 - Hazards Identification and/or Section 11 - Toxicological Information.

**Medical Conditions Aggravated by Exposure:** None known

#### Indication of the Immediate Medical Attention and Special Treatment Needed

**Notes to Physician:** None

### 5. FIRE FIGHTING MEASURES

**Extinguishing Media:** Extinguish fires with CO<sub>2</sub>, extinguishing powder, foam, or water.

#### Special Hazards Arising from the Substance or Mixture

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

**Fire / Explosion Hazards:** Fine particles (such as dust and mists) may fuel fires/explosions.

#### 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.

## SAFETY DATA SHEET

Material Name: Glipizide XL Extended-Release Tablets  
(Greenstone LLC)  
Revision date: 05-Oct-2015

Page 3 of 10

Version: 3.0

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**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

Minimize dust generation and accumulation. If tablets or capsules are crushed and/or broken, avoid breathing dust and avoid contact with eyes, skin, and clothing. When handling, use appropriate personal protective equipment (see Section 8). 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

## SAFETY DATA SHEET

Material Name: Glipizide XL Extended-Release Tablets  
(Greenstone LLC)  
Revision date: 05-Oct-2015

Page 4 of 10

Version: 3.0

### 8. EXPOSURE CONTROLS / PERSONAL PROTECTION

#### Control Parameters

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

#### Glipizide

Manufacturer OEL: 200ug/m<sup>3</sup>

#### Ferric oxide red

ACGIH Threshold Limit Value (TWA) 5 mg/m<sup>3</sup>  
Australia TWA 5 mg/m<sup>3</sup>  
10 mg/m<sup>3</sup>  
Austria OEL - MAKs 5 mg/m<sup>3</sup>  
10 mg/m<sup>3</sup>  
Belgium OEL - TWA 2 ppm  
5 mg/m<sup>3</sup>  
Bulgaria OEL - TWA 5.0 mg/m<sup>3</sup>  
Denmark OEL - TWA 3.5 mg/m<sup>3</sup>  
Estonia OEL - TWA 3.5 mg/m<sup>3</sup>  
Finland OEL - TWA 5 mg/m<sup>3</sup>  
France OEL - TWA 5 mg/m<sup>3</sup>  
Greece OEL - TWA 10 mg/m<sup>3</sup>  
Hungary OEL - TWA 6 mg/m<sup>3</sup>  
Ireland OEL - TWAs 5 mg/m<sup>3</sup>  
10 mg/m<sup>3</sup>  
4 mg/m<sup>3</sup>  
Lithuania OEL - TWA 3.5 mg/m<sup>3</sup>  
OSHA - Final PELs - TWAs: 10 mg/m<sup>3</sup>  
15 mg/m<sup>3</sup>  
Poland OEL - TWA 5 mg/m<sup>3</sup>  
Portugal OEL - TWA 5 mg/m<sup>3</sup>  
Romania OEL - TWA 5 mg/m<sup>3</sup>  
Slovakia OEL - TWA 1.5 mg/m<sup>3</sup>  
Spain OEL - TWA 5 mg/m<sup>3</sup>  
Sweden OEL - TWAs 3.5 mg/m<sup>3</sup>

#### Polyethylene oxide NF

Austria OEL - MAKs 1000 mg/m<sup>3</sup>  
Germany - TRGS 900 - TWAs 1000 mg/m<sup>3</sup>  
Germany (DFG) - MAK 1000 mg/m<sup>3</sup> average molecular weight 200-600  
Slovakia OEL - TWA 1000 mg/m<sup>3</sup>  
Slovenia OEL - TWA 1000 mg/m<sup>3</sup>

#### Magnesium stearate

ACGIH Threshold Limit Value (TWA) 10 mg/m<sup>3</sup>  
Lithuania OEL - TWA 5 mg/m<sup>3</sup>  
Sweden OEL - TWAs 5 mg/m<sup>3</sup>

#### Sodium chloride

Latvia OEL - TWA 5 mg/m<sup>3</sup>  
Lithuania OEL - TWA 5 mg/m<sup>3</sup>

#### Exposure Controls

## SAFETY DATA SHEET

Material Name: Glipizide XL Extended-Release Tablets  
(Greenstone LLC)  
Revision date: 05-Oct-2015

Page 5 of 10

Version: 3.0

**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

<b>Physical State:</b>	Tablet	<b>Color:</b>	Blue (2.5 mg) White (5 and 10 mg)
<b>Odor:</b>	No data available.	<b>Odor Threshold:</b>	No data available.
<b>Molecular Formula:</b>	Mixture	<b>Molecular Weight:</b>	Mixture
<b>Solvent Solubility:</b>	No data available		
<b>Water Solubility:</b>	No data available		
<b>pH:</b>	No data available.		
<b>Melting/Freezing Point (°C):</b>	No data available		
<b>Boiling Point (°C):</b>	No data available.		
<b>Partition Coefficient: (Method, pH, Endpoint, Value)</b>			
<b>Ferric oxide red</b>			
No data available			
<b>Magnesium stearate</b>			
No data available			
<b>Polyethylene oxide NF</b>			
No data available			
<b>Sodium chloride</b>			
No data available			
<b>Glipizide</b>			
Predicted 7.4 Log D 0.046			
<b>Hydroxypropyl methylcellulose</b>			
No data available			
<b>Decomposition Temperature (°C):</b>	No data available.		
<b>Evaporation Rate (Gram/s):</b>	No data available		
<b>Vapor Pressure (kPa):</b>	No data available		
<b>Vapor Density (g/ml):</b>	No data available		
<b>Relative Density:</b>	No data available		
<b>Viscosity:</b>	No data available		
<b>Flammability:</b>			
<b>Autoignition Temperature (Solid) (°C):</b>	No data available		
<b>Flammability (Solids):</b>	No data available		
<b>Flash Point (Liquid) (°C):</b>	No data available		
<b>Upper Explosive Limits (Liquid) (% by Vol.):</b>	No data available		
<b>Lower Explosive Limits (Liquid) (% by Vol.):</b>	No data available		
<b>Polymerization:</b>	Will not occur		

## SAFETY DATA SHEET

Material Name: Glipizide XL Extended-Release Tablets  
(Greenstone LLC)  
Revision date: 05-Oct-2015

Page 6 of 10

Version: 3.0

### 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 Products:** No data available

### 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:** Antidiabetic drug: has blood-sugar lowering properties  
**Known Clinical Effects:** Ingestion of this material may cause effects similar to those seen in clinical use including effects on gastrointestinal disturbances, allergic skin reactions, blood system changes, liver effects, kidney effects, and endocrine reactions. Overdosage of sulfonylureas can produce hypoglycemia which characterized by hunger, nervousness, profuse sweating, faintness, and sometimes convulsions.

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

##### Magnesium stearate

Rat Oral LD50 > 2000 mg/kg  
Rat Inhalation LC50 > 2000 mg/m<sup>3</sup>

##### Sodium chloride

Rat Oral LD50 3000 mg/kg  
Mouse Oral LD50 4000 mg/kg

##### Glipizide

Mouse Oral LD50 > 5000 mg/kg  
Rat Oral LD50 > 4000mg/kg

##### Hydroxypropyl methylcellulose

Rat Oral LD50 > 10,000 mg/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 oxide NF

Eye Irritation Rabbit Mild  
Skin Irritation Rabbit Mild

##### Sodium chloride

Eye Irritation Rabbit Moderate  
Skin Irritation Rabbit Mild

## SAFETY DATA SHEET

Material Name: Glipizide XL Extended-Release Tablets  
(Greenstone LLC)  
Revision date: 05-Oct-2015

Page 7 of 10

Version: 3.0

### 11. TOXICOLOGICAL INFORMATION

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

##### Glipizide

6 Month(s)	Rat	Oral 8 mg/kg/day	NOAEL	No effects at maximum dose
10 Month(s)	Dog	Oral 8 mg/kg/day	NOAEL	No effects at maximum dose
15 Month(s)	Rat	Oral 8 mg/kg/day	NOAEL	No effects at maximum dose
40 Month(s)	Dog	Oral 8 mg/kg/day	NOAEL	No effects at maximum dose

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

##### Glipizide

Reproductive & Fertility	Rat	Oral 50 mg/kg/day	NOAEL	No effects at maximum dose
Embryo / Fetal Development	Rat	Oral 2000 mg/kg/day	NOAEL	No effects at maximum dose
Embryo / Fetal Development	Rabbit	Oral 10 mg/kg/day	NOAEL	No effects at maximum dose
Prenatal & Postnatal Development	Rat	Oral 50 mg/kg/day	NOAEL	No effects at maximum dose

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

##### Glipizide

Bacterial Mutagenicity (Ames)	<i>Salmonella</i>	Negative
<i>In Vivo</i> Cytogenetics	Mouse	Negative
Dominant Lethal Assay	Mouse	Negative

#### Carcinogenicity: (Duration, Species, Route, Dose, End Point, Effect(s))

##### Glipizide

24 Month(s)	Rat	Oral 50 mg/kg/day	NOAEL	Not carcinogenic
18 Month(s)	Mouse	Oral 50 mg/kg/day	NOAEL	Not carcinogenic

#### Carcinogen Status:

None of the components of this formulation are listed as a carcinogen by IARC, NTP or OSHA. See below

#### Ferric oxide red

##### IARC:

Group 3 (Not Classifiable)

### 12. ECOLOGICAL INFORMATION

**Environmental Overview:** Environmental properties have not been investigated.

#### **Toxicity:**

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

##### Glipizide

*Daphnia magna* (Water Flea) LC50 48 Hours > 370 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

## SAFETY DATA SHEET

Material Name: Glipizide XL Extended-Release Tablets  
(Greenstone LLC)  
Revision date: 05-Oct-2015

Page 8 of 10

Version: 3.0

### Bio-accumulative Potential:

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

#### Glipizide

Predicted 7.4 Log D 0.046

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.

## 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:

None required

This product has been classified in accordance with the hazard criteria of the CPR and the MSDS contains all of the information required by the CPR.

#### Glipizide

CERCLA/SARA 313 Emission reporting	Not Listed
California Proposition 65	Not Listed
Standard for the Uniform Scheduling for Drugs and Poisons:	Schedule 4
EU EINECS/ELINCS List	249-427-6

#### Ferric oxide red



## SAFETY DATA SHEET

Material Name: Glipizide XL Extended-Release Tablets  
(Greenstone LLC)  
Revision date: 05-Oct-2015

Page 9 of 10

Version: 3.0

### 15. REGULATORY INFORMATION

CERCLA/SARA 313 Emission reporting	Not Listed
California Proposition 65	Not Listed
Inventory - United States TSCA - Sect. 8(b)	Present
Australia (AICS):	Present
EU EINECS/ELINCS List	215-168-2

#### Polyethylene oxide NF

CERCLA/SARA 313 Emission reporting	Not Listed
California Proposition 65	Not Listed
Inventory - United States TSCA - Sect. 8(b)	Present
Australia (AICS):	Present
Standard for the Uniform Scheduling for Drugs and Poisons:	Schedule 3
EU EINECS/ELINCS List	Not Listed

#### Hydroxypropyl methylcellulose

CERCLA/SARA 313 Emission reporting	Not Listed
California Proposition 65	Not Listed
Inventory - United States TSCA - Sect. 8(b)	Present
Australia (AICS):	Present
Standard for the Uniform Scheduling for Drugs and Poisons:	Schedule 4
EU EINECS/ELINCS List	Not Listed

#### Magnesium stearate

CERCLA/SARA 313 Emission reporting	Not Listed
California Proposition 65	Not Listed
Inventory - United States TSCA - Sect. 8(b)	Present
Australia (AICS):	Present
EU EINECS/ELINCS List	209-150-3

#### Sodium chloride

CERCLA/SARA 313 Emission reporting	Not Listed
California Proposition 65	Not Listed
Inventory - United States TSCA - Sect. 8(b)	Present
Australia (AICS):	Present
EU EINECS/ELINCS List	231-598-3

### 16. OTHER INFORMATION

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

**Reasons for Revision:** Updated Section 1 - Identification of the Substance/Preparation and the Company/Undertaking. Updated Section 2 - Hazard Identification. Updated Section 11 - Toxicology Information. Updated Section 7 - Handling and Storage. Updated Section 3 - Composition / Information on Ingredients.

**Revision date:** 05-Oct-2015  
Product Stewardship Hazard Communication

**Prepared by:** Global Environment, Health, and Safety Operations

## **SAFETY DATA SHEET**

**Material Name: Glipizide XL Extended-Release Tablets  
(Greenstone LLC)  
Revision date: 05-Oct-2015**

**Page 10 of 10**

**Version: 3.0**

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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**