

Revision date: 07-Aug-2014

Version: 1.0

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

**Product Identifier** 

Material Name: Amoxicillin Trihydrate and Clavulanate Potassium Chewable Tablets

Trade Name: Synonyms: Chemical Family: Clavamox, Synulox Clavamox Chewable Tablets; Synulox Chewable Tablets Mixture

Relevant Identified Uses of the Substance or Mixture and Uses Advised Against Intended Use: Veterinary product used as antibiotic agent Restrictions on Use: Not for human use

Details of the Supplier of the Safety Data Sheet

Zoetis Inc. 100 Campus Drive, P.O. Box 651 Florham Park, New Jersey 07932 (USA) Rocky Mountain Poison Control Center Phone: 1-866-531-8896 Product Support/Technical Services Phone: 1-800-366-5288

Emergency telephone number: CHEMTREC (24 hours): 1-800-424-9300 Contact E-Mail: VMIPSrecords@zoetis.com Zoetis Belgium S.A. Mercuriusstraat 20 1930 Zaventem Belgium

Emergency telephone number: International CHEMTREC (24 hours): +1-703-527-3887

# 2. HAZARDS IDENTIFICATION

Appearance:

Brown tablets

#### Classification of the Substance or Mixture GHS - Classification

Respiratory Sensitization: Category 1 Skin Sensitization: Category 1

### **EU Classification:**

EU Indication of danger: Harmful

Xn

EU Symbol: EU Risk Phrases:

R42/43 - May cause sensitization by inhalation and skin contact.

### Label Elements

Signal Word:	Danger
Hazard Statements:	H317 - May cause an allergic skin reaction
	H334 - May cause allergy or asthma symptoms or breathing difficulties if inhaled

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Precautionary Statements:P261 - Avoid breathing dust/fume/gas/mist/vapors/spray P280 - Wear protective gloves/protective clothing/eye protection/face protection P285 - In case of inadequate ventilation wear respiratory protection P272 - Contaminated work clothing should not be allowed out of the workplace P304 + P340 - IF INHALED: Remove victim to fresh air and keep at rest in a positic comfortable for breathing P342 + P311 - If experiencing respiratory symptoms: Call a POISON CENTRE or doctor/physician P302+ P352 - IF ON SKIN: Wash with plenty of soap and water P333 + P313 - If skin irritation or rash occurs: Get medical advice/attention P362 - Take off contaminated clothing and wash before reuse P501 - Dispose of contents/container in accordance with all local and national regulation
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Other Hazards Short Term:

**Known Clinical Effects:** 

Australian Hazard Classification (NOHSC):

Note:

Individuals who are allergic to penicillin antibiotics could have allergic reaction, possibly severe. If an allergic reaction occurs, the worker should be removed to the nearest emergency room and the appropriate therapy instituted.

May cause effects similar to those generally seen in clinical use of antibiotics including gastrointestinal irritation, vomiting, transient diarrhea, nausea, and abdominal pain. Hazardous Substance. Non-Dangerous Goods.

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.

### **3. COMPOSITION/INFORMATION ON INGREDIENTS**

#### Hazardous

Ingredient	CAS Number	EU EINECS/ELINCS List	EU Classification	GHS Classification	%
Microcrystalline cellulose	9004-34-6	232-674-9	Not Listed	Not Listed	15 - 40
Amoxicillin trihydrate	61336-70-7	Not Listed	Xn;R42/43	Skin Sens. 1 (H317) Resp. Sens. 1 (H334)	15
Potassium clavulanate	61177-45-5	262-640-9	Not Listed	Not Listed	4
Colloidal silicon dioxide	7631-86-9	231-545-4	Not Listed	Not Listed	<2
Magnesium stearate	557-04-0	209-150-3	Not Listed	Not Listed	<2

**Additional Information:** 

Ingredient(s) indicated as hazardous have been assessed under standards for workplace safety.

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

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# **4. FIRST AID MEASURES**

### **Description of First Aid Measures**

Immediately flush eyes with water for at least 15 minutes. If irritation occurs or persists, get Eve Contact: medical attention. **Skin Contact:** Remove contaminated clothing and shoes. Wash skin with soap and water. If irritation occurs or persists, get 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 For information on potential signs and symptoms of exposure, See Section 2 - Hazards Identification and/or Section 11 - Toxicological Information. Exposure: **Medical Conditions** People allergic to penicillins may exhibit cross reaction sensitivity. Aggravated by Exposure:

Indication of the Immediate Medical Attention and Special Treatment Needed None

Notes to Physician:

### 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: Fine particles (such as dust and mists) may fuel fires/explosions.

#### **Advice for Fire-Fighters**

Wear approved positive pressure, self-contained breathing apparatus and full protective turn out gear. Use caution in approaching fire.

# 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.

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# 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 at room temperatu

Store at room temperature in properly labeled containers. Keep away from heat, sparks and flames.

Specific end use(s):

No data available

# 8. EXPOSURE CONTROLS / PERSONAL PROTECTION

### **Control Parameters**

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

Microcry	vstalline	cellulose
MILLIOUI Y	Jamme	Centulose

Microcrystalline cellulose	
ACGIH Threshold Limit Value (TWA)	10 mg/m³
Australia TWA	10 mg/m³
Belgium OEL - TWA	10 mg/m³
Estonia OEL - TWA	10 mg/m³
France OEL - TWA	10 mg/m³
Ireland OEL - TWAs	10 mg/m³
	4 mg/m <sup>3</sup>
Latvia OEL - TWA	2 mg/m <sup>3</sup>
Vietnam OEL - TWAs	10 mg/m³
	5 mg/m³
OSHA - Final PELS - TWAs:	15 mg/m³
Portugal OEL - TWA	10 mg/m³
Romania OEL - TWA	10 mg/m³
Spain OEL - TWA	10 mg/m³
Switzerland OEL -TWAs	3 mg/m <sup>3</sup>
Colloidal silicon dioxide	
Australia TWA	2 mg/m <sup>3</sup>
Austria OEL - MAKs	$4 \text{ mg/m}^3$
	0.3 mg/m <sup>3</sup>
Czech Republic OEL - TWA	$0.1 \text{ mg/m}^3$
	$4.0 \text{ mg/m}^3$
Estonia OEL - TWA	2 mg/m <sup>3</sup>
Finland OEL - TWA	5 mg/m <sup>3</sup>
Germany - TRGS 900 - TWAs	4 mg/m <sup>3</sup>
Germany (DFG) - MAK	4 mg/m <sup>3</sup>
Ireland OEL - TWAs	6 mg/m <sup>3</sup> 2.4 mg/m <sup>3</sup>
Latvia OEL - TWA	$1 \text{ mg/m}^3$
OSHA - Final PELs - Table Z-3 Mineral D:	20 mppcf
	Listed
Slovakia OEL - TWA	4.0 mg/m <sup>3</sup>

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8. EXPO	SURE CONTROLS / PERSONAL PROTECTION
Switzerland OEL -TWAs	4 mg/m <sup>3</sup> 0.3 mg/m <sup>3</sup>
Magnesium stearate	
ACGIH Threshold Limit Val	u <b>e (TWA)</b> 10 mg/m <sup>3</sup>
Lithuania OEL - TWA	5 mg/m <sup>3</sup>
Sweden OEL - TWAs	5 mg/m <sup>3</sup>
when the available data are sufficien	posure Band (OEB) classification system is to separate substances into different Hazard categorie at to do so, but inadequate to establish an Occupational Exposure Limit (OEL). The OEB given is tly available data; as such, this value may be subject to revision when new information becomes
Amoxicillin trihydrate	
Zoetis OEB	OEB 2 - Sensitizer (control exposure to the range of 100ug/m <sup>3</sup> to < 1000ug/m <sup>3</sup> , provide additional precautions to protect from skin contact)
Potassium clavulanate	
Zoetis OEB	OEB 2 (control exposure to the range of $100 \text{ug/m}^3$ to < $1000 \text{ug/m}^3$ )
Exposure Controls	
Engineering Controls:	Engineering controls should be used as the primary means to control exposures. Keep air contamination levels below the exposure limits or within the OEB range listed above in this section. General room ventilation is adequate unless the process generates dust, mist or fumes.
Personal Protective Equipment:	Refer to applicable national standards and regulations in the selection and use of personal protective equipment (PPE).
Hands:	Not required for the normal use of this product. Impervious gloves are recommended if skin contact with drug product is possible and for bulk processing operations.
Eyes:	Not required under normal conditions of use. Wear safety glasses or goggles if eye contact is possible.
Skin:	Not required for the normal use of this product. Impervious protective clothing is recommended if skin contact with drug product is possible and for bulk processing operations.
Respiratory protection:	Whenever excessive air contamination (dust, mist, vapor) is generated, respiratory protection, with appropriate protection factors, should be used to minimize exposure. 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. If airborne exposures are within or exceed the Occupational Exposure Band (OEB) range, wear an appropriate respirator with a protection factor sufficient to control exposures to the bottom of the OEB range.

# 9. PHYSICAL AND CHEMICAL PROPERTIES

Physical State: Odor: Molecular Formula:

Solvent Solubility: Water Solubility: pH: Melting/Freezing Point (°C): Boiling Point (°C): Tablet No data available. Mixture

No data available No data available No data available. No data available No data available Color: Odor Threshold: Molecular Weight: Brown , mottled No data available. Mixture

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### 9. PHYSICAL AND CHEMICAL PROPERTIES

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

 No data available

 Decomposition Temperature (°C):

No data available.

Evaporation Rate (Gram/s):	No data available
Vapor Pressure (kPa):	No data available
Vapor Density (g/ml):	No data available
Relative Density:	No data available
Viscosity:	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.): Polymerization: No data available Will not occur

# **10. STABILITY AND REACTIVITY**

Reactivity: Chemical Stability:	No data available Stable under normal conditions of use.
Possibility of Hazardous Reactions Oxidizing Properties:	No data available
Conditions to Avoid:	Avoid dispersion as a dust cloud. Dust may form explosive mixture in air. Fine particles (such as dust and mists) may fuel fires/explosions. Keep away from heat, spark, flames and all other sources of ignition.
Incompatible Materials:	As a precautionary measure, keep away from strong oxidizers
Hazardous Decomposition Products:	Thermal decomposition products may include carbon monoxide, carbon dioxide and other toxic vapors.

### **11. TOXICOLOGICAL INFORMATION**

Information on Toxicological Effects General Information:

Toxicological properties of the formulation have not been investigated. The information included in this section describes the potential hazards of the individual ingredients.

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

#### Potassium clavulanate

Mouse Oral LD50 4526 mg/kg Rat Oral LD50 7936mg/kg

#### Amoxicillin trihydrate

MouseOralLD50> 25 g/kgRatOralLD50> 15g/kgRabbitOralLD50> 12g/kgRatSCLD50> 8g/kg

#### Microcrystalline cellulose

Rat Oral LD50 > 5000 mg/kg

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# **11. TOXICOLOGICAL INFORMATION**

Rabbit Dermal LD50 > 2000 mg/kg

#### Magnesium stearate

RatOralLD50> 2000 mg/kgRatInhalationLC50> 2000 mg/m³Acute Toxicity Comments:A green

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)

#### Microcrystalline cellulose

Skin IrritationRabbitNon-irritatingEye IrritationRabbitNon-irritating

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

#### Potassium clavulanate

26 Week(s) Dog Intravenous 20 mg/kg/day NOAEL Liver

#### Clavulanic Acid/Amoxicillin Trihydrate

4 Week(s)	Mouse	e Oral	50/500 mg/kg/a	day NOAEL	None identified
4 Week(s)	Rat	Oral	50/500 mg/kg/da	y NOAEL	None identified
28 Day(s)	Dog	Oral	90 mg/kg/day	NOEL Gas	trointestinal system
28 Week(s)	Rat	Oral	150 mg/kg/day	NOAEL	Liver, Gastrointestinal system

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

#### Amoxicillin trihydrate

Embryo / Fetal Development Pig Oral 600 mg/kg/day NOEL Not teratogenic

Carcinogen Status:

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

#### Colloidal silicon dioxide IARC:

Group 3 (Not Classifiable)

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# **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)

#### Amoxicillin trihydrate

Daphnia magna (Water Flea) EC	50 48	Hours >	2300 mg	ı/L	
Lepomis macrochirus (Bluegill Sunf	ish) EC	50 96 H	lours >	930 r	ng/L
Oncorhynchus mykiss (Rainbow Tr	out) EC5	50 96 H	ours >	1000	mg/L
Microcystis aeruginosa (Blue-green	Alga) E	C50 48	Hours 0	.0037	mg/L
Selenastrum capricornutum (Green	Alga) N	OEC 48	Hours 2	50 mg	J/L

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.

### **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**

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# **15. REGULATORY INFORMATION**

WHMIS hazard class: Class D, Division 2, Subdivision A Class D, Division 2, Subdivision B This product has been classified in accordance with the hazard criteria of the CPR and the SDS contains all of the information required by the CPR.



Microcrystalline cellulose	
CERCLA/SARA 313 Emission reporting	Not Listed
California Proposition 65	Not Listed
Inventory - United States TSCA - Sect. 8(b)	Present
Australia (AICS):	Present
<b>REACH - Annex XVII - Restrictions on Certain</b>	Use restricted. See item 9[f]. powder
Dangerous Substances:	
EU EINECS/ELINCS List	232-674-9
Americillin tribudroto	
Amoxicillin trihydrate CERCLA/SARA 313 Emission reporting	Not Listed
California Proposition 65	Not Listed
Australia (AICS):	Present
EU EINECS/ELINCS List	Not Listed
	Not Listed
Potassium clavulanate	
CERCLA/SARA 313 Emission reporting	Not Listed
California Proposition 65	Not Listed
EU EINECS/ELINCS List	262-640-9
Colloidal silicon dioxide	
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	231-545-4
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

# **16. OTHER INFORMATION**

### Text of R phrases and GHS Classification abbreviations mentioned in Section 3

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Sensitization, skin-Cat.1; H317 - May cause an allergic skin reaction Sensitization, respiratory-Cat.1; H334 - May cause allergy or asthma symptoms or breathing difficulties if inhaled

Xn - Harmful

R42/43 - May cause sensitization by inhalation and skin contact.

Data Sources:	The data contained in this SDS may have been gathered from confidential internal sources, raw material suppliers, or from the published literature.
Reasons for Revision:	New data sheet.
Prepared by:	Toxicology and Hazard Communication Zoetis Global Risk Management

Zoetis Inc. believes that the information contained in this Safety Data Sheet is accurate, and while it is provided in good faith, it is without 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