This SDS packet was issued with item: 078914045

The safety data sheets (SDS) in this packet apply to one or more components included in the items listed below. Items listed below may require one or more SDS. Please refer to invoice for specific item number(s).

078695525 078695541 078707683 078912799 078914044



Revision date: 18-Sep-2013

Version: 3.1

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

Product Identifier

Material Name: Draxxin (Tulathromycin) Solution for Injection

Trade Name: Synonyms: **Chemical Family:**

DRAXXIN Tulathromycin injectable solution Mixture

Relevant Identified Uses of the Substance or Mixture and Uses Advised Against Intended Use: Veterinary product used as antibiotic agent

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:

Clear, colorless to slightly yellow solution in multiple-dose vials

Classification of the Substance or Mixture GHS - Classification

Serious Eye Damage/Eye Irritation: Category 2A Skin Sensitization: Category 1

Xi

EU Classification:

EU Indication of danger: Irritant

EU Symbol: EU Risk Phrases:

R43 - May cause sensitization by skin contact.

Label Elements

Signal Word:	Warning
Hazard Statements:	H319 - Causes serious eye irritation
	1017 May any a an allowing alve reaction

H317 - May cause an allergic skin reaction

Precautionary Statements:	 P261 - Avoid breathing dust/fume/gas/mist/vapors/spray P264 - Wash hands thoroughly after handling P280 - Wear protective gloves/protective clothing/eye protection/face protection P272 - Contaminated work clothing should not be allowed out of the workplace P305 + P351 + P338 - IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing P337 + P313 - If eye irritation persists: Get medical advice/attention 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 regulations
Other Hazards Short Term:	Individuals sensitive to this chemical or other materials in its chemical class may develop allergic reactions. In the event of accidental injection, an allergic reaction may occur. If an allergic reaction occurs, the worker should be removed to the nearest emergency room and the appropriate therapy instituted.
Known Clinical Effects: Australian Hazard Classification (NOHSC):	Ingestion of this material 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.
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.

3. COMPOSITION/INFORMATION ON INGREDIENTS

Hazardous

Ingredient	CAS Number	EU EINECS/ELINCS List	EU Classification	GHS Classification	%
Tulathromycin	217500-96-4	Not Listed	Xi;R36-R43	Eye Irrit. 2A (H319) Skin Sens. 1 (H317) Aquatic Acute 3 (H402) Aquatic Chronic 3 (H412)	10
Citric acid	77-92-9	201-069-1	Xi; R36	Not Listed	**
Propylene glycol	57-55-6	200-338-0	Not Listed	Not Listed	*
HYDROCHLORIC ACID	7647-01-0	231-595-7	T; R23 C; R35	Skin Corr.1B (H314) STOT SE 3 (H335)	**

Ingredient	CAS Number	EU EINECS/ELINCS	EU Classification	GHS Classification	%
		List			
Monothioglycerol	96-27-5	202-495-0	Not Listed	Not Listed	*
Water	7732-18-5	231-791-2	Not Listed	Not Listed	*

Additional Information:

** to adjust pH

* Proprietary

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

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 Effect Symptoms and Effects of Exposure: Medical Conditions Aggravated by Exposure:	cts, Both Acute and Delayed For information on potential signs and symptoms of exposure, See Section 2 - Hazards Identification and/or Section 11 - Toxicological Information. None known
Indication of the Immediate Medical Notes to Physician:	Attention and Special Treatment Needed 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** May emit toxic fumes of oxides of carbon and nitrogen. **Products:**

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.

Material Name: Draxxin (Tulathromycin) Solution for Injection Revision date: 18-Sep-2013

Environmental Precautions

Collecting:

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 / Absorb spills with non-combustible absorbent material and transfer into a labeled container for disposal. 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

Avoid accidental injection. Minimize generating airborne mists and vapors. Avoid breathing mist or aerosols. When handling, use appropriate personal protective equipment (see Section 8). Wash hands and any exposed skin after removal of PPE. 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. Refer to Section 12 - Ecological Information, for information on potential effects on the environment.

Conditions for Safe Storage, Including any Incompatibilities

Storage Conditions: Specific end use(s):

Store as directed by product packaging. No data available

8. EXPOSURE CONTROLS / PERSONAL PROTECTION

1mg/m³, Sensitizer

150 ppm 474 mg/m³ 10 mg/m^{3}

150 ppm 470 ma/m³ 10 ma/m³

7 mg/m³

7 mg/m³

2 ppm

5 ppm 7.5 mg/m³

5 ppm 8 mg/m³ 5 ppm

8 ma/m³ 8.0 ma/m³

5 ppm 5 ppm

8 mg/m³

Control Parameters

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

Tulathromycin

7		T14/4	0 1
Zoetis	OEL	IWA	8-nr

Propylene glycol

Australia TWA

Ireland OEL - TWAs

Latvia OEL - TWA Lithuania OEL - TWA

HYDROCHLORIC ACID

ACGIH Ceiling Threshold Limit: Australia PEAK

Austria OEL - MAKs

Belgium OEL - TWA

Bulgaria OEL - TWA

Cyprus OEL - TWA

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		URE CONTROLS / PERSONAL PROTECTION
	Czech Republic OEL - TWA	8 mg/m ³
	Estonia OEL - TWA	5 ppm
		8 mg/m ³
	Germany - TRGS 900 - TWAs	2 ppm
		3 mg/m ³
	Germany (DFG) - MAK	2 ppm
		3.0 mg/m ³
	Greece OEL - TWA	5 ppm
		7 mg/m ³
	Hungary OEL - TWA	8 mg/m ³
	Ireland OEL - TWAs	5 ppm
		8 mg/m ³
	Italy OEL - TWA	5 ppm
		8 mg/m ³
	Japan - OELs - Ceilings	5 ppm
		7.5 mg/m ³
	Latvia OEL - TWA	5 ppm
		8 mg/m ³
	Lithuania OEL - TWA	5 ppm
		8 mg/m ³
	Luxembourg OEL - TWA	5 ppm
		8 mg/m ³
	Malta OEL - TWA	5 ppm
		8 mg/m ³
	Netherlands OEL - TWA	8 mg/m ³
	Vietnam O EL - TWAs	5 mg/m ³
	Poland OEL - TWA	5 mg/m ³
	Romania OEL - TWA	5 ppm
		8 mg/m ³
	Slovakia OEL - TWA	5 ppm
		8.0 mg/m ³
	Slovenia OEL - TWA	5 ppm
		8 mg/m ³
	Spain OEL - TWA	5 ppm
		7.6 mg/m ³
	Switzerland OEL -TWAs	2 ppm
	Switzenand OEL -I WAS	3.0 mg/m ³
		5.0 mg/m
Expo	sure 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	Refer to applicable national standards and regulations in the selection and use of personal
	Equipment:	protective equipment (PPE).
	-4-16.000	
	Useda	We as increase increase to an except of the sector of
	Hands:	Wear impervious gloves to prevent skin contact.
	Eyes:	Wear safety glasses or goggles if eye contact is possible.
	Skin:	Wear impervious protective clothing to prevent skin contact - consider use of disposable
		clothing where appropriate.
	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.

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9. PHYSICAL AND CHEMICAL PROPERTIES

Physical State: Odor:	Solution in multiple-dose vials No data available.
Molecular Formula:	Mixture
Solvent Solubility:	No data available
Water Solubility:	No data available
pH:	5.4
Melting/Freezing Point (°C):	No data available
Boiling Point (°C):	No data available.
Partition Coefficient: (Method, pH, E	ndpoint, Value)
No data available	
Tulathromycin	
Measured 7.0 Log P -1.41	
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 (Sol	lid) (°C): No
Flammability (Solids):	No
Flash Point (Liquid) (°C):	No

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

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

Color: Odor Threshold: Molecular Weight: Colorless to slightly yellow No data available. Mixture

No data available Will not occur

10. STABILITY AND REACTIVITY

Reactivity: Chemical Stability: Possibility of Hazardous Reactions Oxidizing Properties: Conditions to Avoid: Incompatible Materials: Hazardous Decomposition Products: No data available Stable under normal conditions of use.

No data available Fine particles (such as dust and mists) may fuel fires/explosions. As a precautionary measure, keep away from strong oxidizers No data available

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)

Tulathromycin

Polymerization:

Rat Oral LDmin. > 2000 mg/kg

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Material Name: Draxxin (Tulathromycin) Solution for Injection Revision date: 18-Sep-2013

11. TOXICOLOGICAL INFORMATION

Rabbit Dermal LD50 > 2000 mg/kg

Citric acid

Rat Oral LD50 3000 mg/kg

Propylene glycol

Mouse Oral LD50 22,000 mg/kg Rat Oral LD50 20,000 mg/kg Rabbit Dermal LD50 20,800 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)

Tulathromycin

Skin Irritation Rabbit Non-irritating Eye Irritation Rabbit Positive Skin Sensitization - GPMT Guinea Pig Severe

Citric acid

Eye Irritation Rabbit Severe Skin Irritation Rabbit Mild

Propylene glycol

Skin Irritation Rabbit Mild Eye Irritation Rabbit Mild

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

Tulathromycin

1 Month(s)	Rat	Oral 50 mg/kg/day	NOAEL	Liver, Blood
3 Month(s)	Rat	Oral 15 mg/kg/day	NOAEL	Liver
1 Month(s)	Dog	Oral 15 mg/kg/day	NOAEL	Liver
3 Month(s)	Dog	Oral 5 mg/kg/day	NOEL	Liver
1 Year(s)	Dog	Oral 5 mg/kg/day	NOAEL	Liver, Male reproductive system

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

Tulathromycin

2 Generation Reproductive Toxicity Rat Oral 50 mg/kg/day NOAEL Paternal toxicity Oral 100 mg/kg/day 2 Generation Reproductive Toxicity Rat NOAEL Neonatal toxicity, Fertility Embryo / Fetal Development Oral 200 mg/kg/day Rat NOAEL No effects at maximum dose Oral 50 mg/kg/day Embryo / Fetal Development Rabbit NOAEL No effects at maximum dose

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

Tulathromycin

Bacterial Mutagenicity (Ames)SalmonellaNegativeIn Vitro Chromosome AberrationHuman LymphocytesNegativeIn Vivo Micronucleus Chromosome AberrationRatNegativeIn Vitro Chromosome AberrationChinese Hamster Ovary (CHO) cellsNegativeIn Vitro Mammalian Cell MutagenicityChinese Hamster Ovary (CHO) cellsNegative

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11. TOXICOLOGICAL INFORMATION

Carcinogen Status:

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

HYDROCHLORIC ACID IARC:

Group 3 (Not Classifiable)

12. ECOLOGICAL INFORMATION

Environmental Overview:

Environmental properties of the formulation have not been investigated. The following information is available for the individual ingredients.

Toxicity:

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

Tulathromycin

Daphnia magna (Water Flea) OECD EC50 48 Hours 64 mg/L 48 Hours Mysidopsis bahia (Mysid Shrimp) OECD LC50 20 ma/L Cyprinodon variegatus (Sheepshead Minnow) OECD LC50 48 Hours 20 mg/L Oncorhynchus mykiss (Rainbow Trout) OECD LC50 96 Hours > 982 mg/L Selenastrum capricornutum (Green Alga) OECD EC-50 72 Hours 70 ug/L

Aquatic Toxicity Comments: A greater than (>) symbol indicates that acute ecotoxicity was not observed at the maximum solubility. Since the substance is insoluble in aqueous solutions above this concentration, an acute ecotoxicity value (i.e. LC/EC50) is not achievable.

Bacterial Inhibition: (Inoculum, Method, End Point, Result)

Tulathromyc Polytox IC-5		mg/L		
Persistence and Degradability: No data available				
Bio-accumul Tulathromyc		Potenti	al:	No data available
Measured	7.0	Log P	-1.41	
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.

Material Name: Draxxin (Tulathromycin) Solution for Injection Revision date: 18-Sep-2013

Canada - WHMIS: Classifications

Class D, Division 2, Subdivision B

WHMIS hazard class:

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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

Tulathromycin **CERCLA/SARA 313 Emission reporting** Not Listed **California Proposition 65** Not Listed **EU EINECS/ELINCS List** Not Listed Monothioglycerol **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** 202-495-0 **Citric acid 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** 201-069-1 **Propylene glycol 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 200-338-0

Water

Material Name: Draxxin (Tulathromycin) Solution for Injection Revision date: 18-Sep-2013

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15. REGULATO	RY INFORMATION
CERCLA/SARA 313 Emission reporting	Not Listed
California Proposition 65	Not Listed
Inventory - United States TSCA - Sect. 8(b)	Present
Australia (AICS):	Present
REACH - Annex IV - Exemptions from the obligations of Register:	Present
EU EINECS/ELINCS List	231-791-2
HYDROCHLORIC ACID	
CERCLA/SARA 313 Emission reporting	1.0 %
CERCLA/SARA Hazardous Substances	5000 lb
and their Reportable Quantities:	2270 kg
CERCLA/SARA - Section 302 Extremely Hazardous TPQs	500 lb
CERCLA/SARA - Section 302 Extremely Hazardous Substances EPCRA RQs	5000 lb
California Proposition 65	Not Listed
Inventory - United States TSCA - Sect. 8(b)	Present
Australia (AICS):	Present
Standard for the Uniform Scheduling	Schedule 5
for Drugs and Poisons:	Schedule 6
EU EINECS/ELINCS List	231-595-7

16. OTHER INFORMATION

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

Sensitization, skin-Cat.1; H317 - May cause an allergic skin reaction Serious eye damage/eye irritation-Cat.2A; H319 - Causes serious eye irritation Skin corrosion/irritation-Cat.1B; H314 - Causes severe skin burns and eye damage Hazardous to the aquatic environment, acute toxicity-Cat.3; H402 - Harmful to aquatic life Hazardous to the aquatic environment, chronic toxicity-Cat.3; H412 - Harmful to aquatic life with long lasting effects Specific target organ toxicity, single exposure; Respiratory tract irritation-Cat.3; H335 - May cause respiratory irritation

T - Toxic C - Corrosive Xi - Irritant

R23 - Toxic by inhalation.

R35 - Causes severe burns.

R36 - Irritating to eyes.

R43 - May cause sensitization by skin contact.

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.
Prepared by:	Toxicology and Hazard Communication Zoetis Global Risk Management

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Zoetis Inc. believes that the information contained in this Material 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