

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

### SUBSTANCE:

Trade Name	AIVLOSIN® (62.5% w/w tylvalosin) Water Soluble Granules		
Chemical Family:	Antibiotic, macrolide class		
Synonyms:	Tylvalosin tartrate (approved non-proprietary name) 3-O-acetyl-4"-O-isovaleryltylosin tartrate Acetylisovaleryltylosin tartrate AIVT tartrate AIV tartrate		
Therapeutic Use:	Veterinary antibiotic powder for preparation of medicated drinking water for animals.  Not for human use.		

### **CONTACT INFORMATION:**

# FOR EMERGENCIES IN USA OR CANADA CONTACT:

**ASPCA Animal Product Safety Service** 

Telephone: 1-800-345-4735

Hours: 24 Hours a Day/ 7 Days a week

# Section 2. COMPOSITION/INFORMATION ON INGREDIENTS

COMPONENT 1	
Common Name:	Tylvalosin tartrate (approved non-proprietary name)
Chemical Name:	(-)- $(4R,5S,6S,7R,11E,13E,15S,16R)$ - $15$ -[[(6-deoxy-2,3-di- $O$ -methyl- $\beta$ -D- $allo$ pyranosyl)oxy]methyl]-6-[[3,6,-dideoxy-4- $O$ -[2,6-dideoxy-3- $C$ -methyl-4- $O$ -(3-methylbutanoyl)- $\alpha$ -L- $ribo$ -hexopyranosyl]-3-(dimethylamino)- $\beta$ -D- $gluco$ pyranosyl]oxy]-4-acetoxy-16-ethyl-5,9,13-trimethyl-2,10-dioxooxacyclohexadeca-11,13-diene-7-acetaldehyde( $2R,3R$ )-tartrate
CAS Registry No.:	63409-12-1 (free base) 63428-13-7 (tartrate)
Percent by Weight:	96 % w/w
COMPONENT 2	Non-Hazardous Inert Ingredients
Percent by Weight:	~ 4 % w/w

# Section 3. HAZARDS IDENTIFICATION

# PRIMARY ROUTES OF EXPOSURE:

SKIN CONTACT:	Probable, if protective clothing, gloves are not worn during handling.
EYE CONTACT:	Possible, if goggles are not worn during handling
INHALATION:	Probable, if dust mask is not worn during handling.
INGESTION:	Low likelihood. Avoid eating, chewing of gum and smoking during handling.
EFFECTS OF	Overexposure to dust may cause respiratory effects including coughing, sneezing or
OVEREXPOSURE	wheezing.





# Section 4. FIRST AID MEASURES

SKIN CONTACT:	Wash hands and exposed skin after working and before eating			
EYE CONTACT:	Wash eyes immediately with water. Seek medical attention if irritation persists.			
INHALATION:	Remove individual from area of exposure; supportive treatment.			
INGESTION:	Treat symptomatically; induction of vomiting may not be needed.			
NOTE TO PHYSICIAN:	N: AIVLOSIN ® Water Soluble Granules has a low toxicity. In cases of			
	idiosyncratic reactions following exposure or accidental ingestion, treat			
	symptomatically.			
ANTIDOTES:	No data available.			

# Section 5. FIRE FIGHTING MEASURES

# FIRE CONTROL:

Extinguisher Media:	Water
Fire Fighting Instructions:	Use spray
Extinguisher Media To Avoid:	None known

### FIRE AND EXPLOSION HAZARDS:

Maximum rate of Pressure Rise (Pmax)	7.5 barg
Specific Material Constant (Kst)	247 bar.m/sec
<b>Dust Explosion Class</b>	St2
Minimum Ignition Energy (MIE)	3-10 mJ.

# Section 6. ACCIDENTAL RELEASE MEASURES

General Measures:	Review Sections 3, 8 and 12 before proceeding with clean-up.
Small Spill:	Wear dust mask, overalls, gloves and sweep up spilled material.
Large Spill:	Wear dust mask, overalls, gloves and sweep up spilled material.

# Section 7. HANDLING AND STORAGE

General Handling	Minimize creation of dust; Close bag tightly after use.
Storage Conditions	Cool, dry conditions
Temperature Range for Storage	At or below 25°C (77°F)

# Section 8. EXPOSURE CONTROLS / PERSONAL PROTECTION

Ventilation	Ventilation recommended to reduce respirable dust
Respiratory Protection:	Dust Mask
Eye Protection:	Goggles
Skin Protection:	Overalls
Hand protection:	Gloves (e.g., latex, polyethylene)
Other Protective	None
Equipment:	



The information in the following sections apply to Tylvalosin Tartrate.

# Section 9. PHYSICAL AND CHEMICAL PROPERTIES

Physical Form / Appearance	White to off-v	vhite powder (AIVLO	OSIN® Water So	oluble Granules)
Odor Threshold	No information available			
Molecular Weight	1042.27 (free base)			
	1192.27 (tartr	ate)		
Molecular Formula	C53H87NO19C4H6O6			
pH	3.5 – 5.0			
Melting Point	125-129°C			
<b>Boiling Point</b>	Not applicable			
Water Solubility	948 g/L at 25 ± 2°C			
Organic Solvent Solubility	Methanol	387 g/L at 25°C	Ethanol	220 g/L at 25 °C
		331 g/L at 25°C	Ethyl acetate	
		301 g/L at 25°C	Ether	
	·	274 g/L at 25°C		
Vapor Pressure	No information available. As a high molecular weight solid material, this			
	material is not expected to possess a significant vapor pressure.			
Octanol:Water Partition	3.82			
Coefficient (log Pow)				

# Section 10. STABILITY AND REACTIVITY

Reactivity:	None known		
Conditions To Avoid:	See Explosive Properties		
Incompatibility:	None known		
Flammability:	Not applicable (see Explosive Properties)		
Explosive Properties:	The creation of a dust cloud of organic material of any origin may form an explosive mixture with air. See Section 5 for more details.		
Hazardous Polymerization:	None known – unlikely		
Hazardous Decomposition	None known – unlikely		
Products:			

# Section 11. TOXICOLOGY INFORMATION

ORAL TOXICITY:	Mouse oral LD <sub>50</sub> >750 mg/kg bodyweight (respiratory disturbance, salivation, locomotor effects, reduced growth rate)	
	Rat oral LD <sub>50</sub> >3016 mg/kg bodyweight (respiratory disturbance, salivation,	
	locomotor effects, reduced growth rate)	
INHALATION TOXICITY:	No information available. Particle size analyses using Aivlosin® premix	
	suggest that less than 5.7 % of dust particles are inhalable and less than about	
1	2.3% could penetrate the lungs.	
SKIN IRRITATION:	In an occluded 4 hr exposure test in rabbits, a 1% saline solution of Aivlosin	
	tartrate was non-irritating. A 10% saline solution was mildly irritating to the	
	skin.	



### Section 11. TOXICOLOGY INFORMATION (Continued)

SENSITIZATION:	In a Guinea pig maximization assay (Magnusson-Kilgmann) 12 of 20 animals showed discreet or patchy erythema following challenge. Aivlosin tartrate is considered to be a mild sensitizer.
GENOTOXICITY:	No evidence of mutagenic potential in bacterial and mammalian cell gene mutation assays. In <i>in vitro</i> cell culture assays structural chromosomal damage was observed in two studies, however, in three <i>in vivo</i> tests, no evidence of chromosomal damage was observed.
CARCINOGENICITY:	Studies were not required as short term tests showed no relevant adverse effects.
REPRODUCTIVE EFFECTS:	There were no adverse effects on parameters of reproduction in rats fed tylvalosin in the diet up to 10,000 ppm, a dose which produced maternal toxicity. Fetuses from the 10,000 ppm dose group had slightly reduced body weights. NOEL 400 ppm (approx 18 mg tylvalosin/kg bodyweight). Developmental toxicity studies in rats and mice showed slightly reduced fetal body weights, but only at doses which were maternally toxic. There was no evidence of other developmental effects and no evidence of teratogenic effects in either species.
IMMUNOTOXICOLOGIC EFFECTS:	No evidence in several repeated dose studies that tylvalosin has any adverse effect on the components of the immune system (e.g. thymus, spleen, lymph nodes). There was no evidence of increased incidences of infections.
PHARMACOLOGIC EFFECTS:	Macrolides interfere with protein synthesis by binding reversibly to the 50S ribosome subunit. They bind to the donor site and prevent the translocation necessary to maintain the growth of peptide chains.
TARGET ORGAN EFFECTS:	None reported for this antimicrobial drug.

# Section 12. ECOLOGICAL INFORMATION

### **ENVIRONMENTAL FATE:**

**Mobility:** Poor, tylvalosin has a very strong affinity for soil **Persistence / Degradability:** DT<sub>50</sub> (soil half life) <20 days

**Bioaccumulative Potential: Low** 

### **ECOTOXICITY:**

**Toxicity to Fish:**  $LC_{50} > 100$  mg tylvalosin/L (rainbow trout) **Toxicity to Daphnids:** 48 hr  $EC_{50} = 617$  mg tylvalosin/L; **Toxicity to Plants:**  $LC_{50} > 1090$  mg tylvalosin/kg soil

### Section 13. DISPOSAL CONSIDERATIONS

Observe all federal, state and local regulations when disposing of this material. May be disposed of by landfill.





### Section 14. TRANSPORT INFORMATION

DOT Considerations: None established IMO Considerations: None established IATA Considerations: None established State Regulations: None established Local Considerations: None established

### Section 15. REGULATORY INFORMATION

This material is for the treatment of animals only

Keep out of the reach of children.

**OECD Harmonized Classification**: Xi; R43

### Section 16. OTHER INFORMATION

**HAZARD LABEL:** May cause skin irritation. Tylvalosin tartrate has been shown to cause hypersensitization in laboratory animals. Persons with known hypersensitivity should avoid contact with the product. In case of accidental ingestion, seek medical advice.

#### OTHER INFORMATION:

EINECS NUMBER Component 1: 264-132-2 (free base). Tartrate salt not listed

**EINECS NUMBER Component 2:** Not listed **EINECS NUMBER Component 3:** Not listed

# ECO ANIMAL HEALTH

344 Nassau Street
Princeton, NJ 08540, USA
Tel: 1-609-683-9030 Fax: 1-609-683-3727
email: nate.manco@ecoanimalhealth.com

The Grange, 100 The High Street
Southgate, London N14 6BN, UK
Tel: +44 (0) 20 8447 8899 Fax: +44 (0) 20 8447 9292

www.ecoanimalhealth.com

email: sales@ecoanimalhealth.com

This information contained herein is based on data considered to be accurate. However, no warranty is expressed or implied regarding the accuracy of these data or the results to be obtained from the use thereof. The publisher, its parent, subsidiaries and affiliates assume no responsibility for personal injury or property damage to vendees, users, or third parties caused by the material. Such persons assume all risks associated with the use of the material.