

EQUIMAX®

(ivermectin 1.87%/praziquantel 14.03%)
Paste

1. IDENTIFICATION

Product Name EQUIMAX®

(ivermectin 1.87%/praziquantel 14.03%) Paste

Recommended use of the chemical and

restrictions on use

Identified usesWorm control for horses and ponies

Restrictions on Use Not for human use.

Company Identification Virbac AH, Inc.
P.O. Box 162059

Fort Worth, Texas 76161

(800) 424-9300

Customer Information Number (800) 338-3659

Emergency Telephone Number

Chemtrec Number

Other Emergency Number: Human Toll-free 833-224-2009 Animal Toll-free 833-224-2013

Issue DateMarch 3, 2020Supersedes DateDecember 21, 2015

Safety Data Sheet prepared in accordance with OSHA's Hazard Communication Standard (29 CFR 1910.1200) and the Globally Harmonized System of Classification and Labelling of Chemicals (GHS)

2. HAZARDS IDENTIFICATION

Hazard Classification

Acute Hazards to the Aquatic Environment – Category 1 (This classification not adopted by OSHA.)

Label Elements

Hazard Symbols



Signal Word: Warning

Hazard Statements

Very toxic to aquatic life.

Precautionary Statements

Prevention

Avoid release to the environment.

Response

None .

Storage

None

Disposal

Dispose of contents/container in accordance with local regulation.

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2. HAZARDS IDENTIFICATION

Other Hazards

None

Specific Concentration Limits

The values listed below represent the percentages of ingredients of unknown toxicity.

Acute oral toxicity 0%
Acute dermal toxicity <10%
Acute inhalation toxicity <10%
Acute aquatic toxicity 10 - 20%

3. COMPOSITION/INFORMATION ON INGREDIENTS

Synonyms:

This product is a mixture.

 Component Name
 CAS Number
 Concentration

 Praziquantel
 55268-74-1
 14.03%

 Ivermectin
 70288-86-7
 1.87%

 Titanium Dioxide
 13463-67-7
 1 - 5%

4. FIRST AID MEASURES

Description of necessary first-aid measures

Eyes

Immediately flood the eye with plenty of water for at least 15-20 minutes, holding the eye open. Obtain medical attention if soreness or redness persists.

Skin

Wash skin thoroughly with soap and water. Obtain medical attention if redness or soreness persists.

Ingestion

Call a poison control center or doctor immediately for treatment advice. Do not induce vomiting unless told to do so by a poison control center or doctor. Do not give anything by mouth to an unconscious person.

Inhalation

Remove person to fresh air. Seek medical attention if symptoms persist.

Most important symptoms/effects, acute and delayed

Aside from the information found under Description of necessary first aid measures (above) and Indication of immediate medical attention and special treatment needed, no additional symptoms and effects are anticipated.

Indication of immediate medical attention and special treatment needed Notes to Physicians

Treat symptomatically.

5. FIRE - FIGHTING MEASURES

Extinguishing Media

Use extinguishing media appropriate for surrounding materials.

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5. FIRE - FIGHTING MEASURES

Unusual Fire and Explosion Hazards

Can release hazardous vapors during a fire.

Protective Equipment for Fire-Fighting

Wear full protective clothing and self-contained breathing apparatus.

6. ACCIDENTAL RELEASE MEASURES

Personal precautions, protective equipment and emergency procedures

Wear appropriate protective clothing.

Environmental Precautions

Prevent the material from entering drains or watercourses.

Methods and materials for containment and cleaning up

Wipe up and transfer into suitable containers for recovery or disposal.

7. HANDLING AND STORAGE

Precautions for safe handling

Wear appropriate protective clothing. Wash hands after dispensing to dog and before eating, drinking or smoking.

Conditions for safe storage

Store in original container at room temperature 25°C/77°F with excursions permitted between 59°F and 86°F (15°C - 30°C). Keep out of sunlight. Keep away from children.

8. EXPOSURE CONTROLS/PERSONAL PROTECTION

Control parameters

Exposure limits are listed below, if they exist.

Praziquantel

None established

Ivermectin

None established

Titanium Dioxide

ACGIH TLV: 10 mg/m3 TWA

OSHA PEL: 15 mg/m³ TWA (Total dust)

Appropriate engineering controls

No specific measures necessary. Good general room ventilation is expected to be adequate to control airborne levels.

Individual protection measures

Respiratory Protection

Not required under normal conditions of use.

Skin Protection

Protective gloves.

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8. EXPOSURE CONTROLS/PERSONAL PROTECTION

Eye/Face Protection

Not required under normal conditions of use.

Body Protection

Normal work wear.

9. PHYSICAL AND CHEMICAL PROPERTIES

Appearance

Physical State Solid (paste)

Color White

Odor None

Odor Threshold No data available pН Not applicable Density No data available **Boiling Range/Point (°C/F)** Not applicable Melting Point (°C/F) Not applicable Flash Point (PMCC) (°C/F) Not flammable Vapor Pressure Not applicable **Evaporation Rate (BuAc=1)** Not applicable Solubility in Water No data available Vapor Density (Air = 1) Not applicable VOC Not applicable Partition coefficient (n-Not applicable

octanol/water)

Viscosity

Auto-ignition Temperature

Decomposition Temperature
Upper explosive limit
Lower explosive limit
Flammability (solid, gas)

Not applicable
No data available
No data available
No data available

10. STABILITY AND REACTIVITY

Reactivity

Data is not available

Chemical Stability

Stable under normal conditions.

Possibility of hazardous reactions

Hazardous polymerization will not occur.

Conditions to Avoid

Heat - high temperatures

Incompatible Materials

Strong oxidizers

Hazardous Decomposition Products

Oxides of carbon - organic compounds

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11. TOXICOLOGICAL INFORMATION

See product insert and/or packaging for additional information.

Acute Toxicity

Ivermectin:

Oral LD50 (rat) 50 mg/kg
Oral LD50 (human) >15mg/kg
Dermal LD50 406 mg/kg (rabbit)
Praziquantel
Oral LD50 (rat) 2840 mg/kg
Dermal LD50 (rat) >2000 mg/kg

Specific Target Organ Toxicity (STOT) - single exposure

Ivermectin: At high doses in humans and animals vomiting, tachycardia, blood pressure fluctuation, CNS effects (somnolence, ataxia) and visual disturbances have been observed. Higher doses may cause death due to respiratory depression.

Praziquantel: Acute overexposure (ingestion) may cause dizziness, headache, drowsiness, malaise, abdominal pain.

Specific Target Organ Toxicity (STOT) - repeat exposure

Ivermectin: No data available

Praziquantel: Chronic overexposure may cause nausea, lethargy, diarrhea, itching, fever and rash.

Serious Eye damage/Irritation

Ivermectin: Slightly irritating to eyes in rabbit tests. Praziquantel: Not irritating to eyes in rabbit studies.

Skin Corrosion/Irritation

Ivermectin: Non-irritating in animal studies.
Praziquantel: Not irritating to skin in rabbit studies.

Respiratory or Skin Sensitization

Ivermectin: Hypersensitivity reactions have been reported in humans.

Praziquantel: Not sensitizing to skin in guinea pig study.

Carcinogenicity

<u>Titanium Dioxide:</u> IARC Overall Evaluation is 2B (Possibly carcinogenic to humans) IARC evaluation guidelines consider the generation of tumors, in 2 different studies within the same animal species, to be adequate criteria for an assessment of sufficient evidence. The conclusions of several epidemiology studies on more than 20000 TiO₂ industry workers in Europe and the USA did not suggest a carcinogenic effect of TiO₂ dust on the human lung. Mortality from other chronic diseases, including other respiratory diseases, was also not associated with exposure to TiO₂ dust. Based upon these studies, titanium dioxide is not expected to cause lung cancer or chronic respiratory diseases in humans at concentrations experienced in the workplace.

Praziquantel: Long term studies in rats and golden hamsters did not reveal any carcinogenic effect.

Germ Cell Mutagenicity

Ivermectin: Negative in the AMES Assay, and in a mouse lymphoma mutation assay. In addition, it did not induce unscheduled DNA synthesis in a human fibroblast cell culture, suggesting that it does not damage DNA.

Praziguantel: Negative results for in vivo and in vitro animal studies.

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11. TOXICOLOGICAL INFORMATION

Reproductive Toxicity

Ivermectin: Teratogenic in rats, rabbit and mice at or near materno-toxic dose levels. The abnormalities are limited mainly to cleft palate.

Praziquantel: Studies in rats and rabbits have shown no evidence of impaired fertility or harm to the fetus. An increase of the abortion rate was found in rats at three times the single human therapeutic dose.

Aspiration Hazard

Not an aspiration hazard.

12. ECOLOGICAL INFORMATION

Ecotoxicity

Ivermectin:

LC50 (Trout) 0.003mg/l 96hr

LC50 (Daphnia magna) 0.000025mg/l 48hr

Mobility in soil

Ivermectin: Ivermectin is metabolized in the soil. Water solubility is limited and it binds to soil very tightly. It does not bioconcentrate in fish and is not taken up from soil into plants.

Persistence/Degradability

Ivermectin: Slow biodegradation. Photodegrades rapidly.

Bioaccumulative Potential

No relevant studies identified.

Other adverse effects

No relevant studies identified.

13. DISPOSAL CONSIDERATIONS

Dispose of the syringe in an approved landfill or by incineration.

14. TRANSPORT INFORMATION

Contact supplier for transport information.

15. REGULATORY INFORMATION

United States TSCA Inventory

This product is excluded from the US EPA Toxic Substance Control Act and is regulated under the Food, Drug and Cosmetic Act.

Canada DSL Inventory

All ingredients have not been verified for listing on the Domestic Substance List (DSL).

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15. REGULATORY INFORMATION

SARA Title III Sect. 311/312 Categorization

None

SARA Title III Sect. 313

This product does not contain any chemicals listed in Section 313 at or above de minimis concentrations.

16. OTHER INFORMATION

Legend

ACGIH: American Conference of Governmental Industrial Hygienists

BOD: Biological Oxygen Demand

CAS#: Chemical Abstracts Service Number

DSL: Domestic Substances List ECHA: European Chemicals Agency EPA: Environmental Protection Agency

FIFRA: Federal Insecticide, Fungicide and Rodenticide Act

HDT: Highest Dose Tested

HMIS: Hazardous Materials Identification System IARC: International Agency for Research on Cancer

LC50: Lethal Concentration 50%

LD50: Lethal Dose 50%

LOEL: Lowest Observed Effect Level

N/A: Denotes no applicable information found or available

NFPA: National Fire Protection Association

NOEL: No Observed Effect Level

OSHA: Occupational Safety and Health Administration

PEL: Permissible Exposure Limit TSCA: Toxic Substances Control Act

SARA: Superfund Amendments and Reauthorization Act

SDS: Safety Data Sheet

STEL: Short Term Exposure Limit

WHMIS: Workplace Hazardous Materials Information System

TLV: Threshold Limit Value

TSCA: Toxic Substance Control Act

Revision Date: March 3, 2020 Replaces: December 21, 2015

Changes made: Update to emergency numbers and section 15.

Information Source and References

This SDS is prepared by Hazard Communication Specialists based on information provided by internal company references.

Prepared By: EnviroNet LLC.

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16. OTHER INFORMATION

The information and recommendations presented in this SDS are based on sources believed to be accurate. Virbac AH, Inc. assumes no liability for the accuracy or completeness of this information. It is the user's responsibility to determine the suitability of the material for their particular purposes. In particular, we make NO WARRANTY OF MERCHANTABILITY OR ANY OTHER WARRANTY, EXPRESS OR IMPLIED, with respect to such information, and we assume no liability resulting from its use. Users should ensure that any use or disposal of the material is in accordance with applicable Federal, State, and local laws and regulations.

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