

Ivermectin Paste 1.87%

Anthelmintic and Boticide

For Oral Use in Horses Only

INDICATIONS:

Consult your veterinarian for assistance in the diagnosis, treatment, and control of parasitism. Ivermectin Paste 1.87% provides effective treatment and control of the following parasites in horses.

Large Strongyles (adults) – *Strongylus vulgaris* (also early forms in blood vessels), *S. edentatus* (also tissue stages), *S. equinus*, *Triodontophorus* spp. including *T. brevicauda* and *T. serratus*, and *Craterostomum acuticaudatum*; **Small Strongyles** (adults, including those resistant to some benzimidazole class compounds) – *Coronocylus* spp. including *C. coronatus*, *C. labiatus* and *C. labratus*;

Cyathostomum spp. including *C. catinatum* and *C. pateratum*; *Cylicocylus* spp. including *C. insigne*, *C. leptostomum*, *C. nassatus* and *C. brevicapsulatus*; *Cylicodontophorus* spp.; *Cylicostephanus* spp. including *C. calicatus*, *C. goldi*, *C. longibursatus* and *C. minutus*, and *Petrovinema poculatum*; **Small Strongyles** – Fourth-stage larvae; **Pinworms** (adults and fourth-stage larvae) – *Oxyuris equi*; **Ascarids** (adults and third- and fourth-stage larvae) – *Parascaris equorum*; **Hairworms** (adults) – *Trichostrongylus axei*; **Large-mouth Stomach**

Worms (adults) – *Habronema muscae*; **Bots** (oral and gastric stages) – *Gasterophilus* spp. including *G. intestinalis* and *G. nasalis*; **Lungworms** (adults and fourth-stage larvae) – *Dictyocaulus arnfieldi*; **Intestinal**

Threadworms (adults) – *Strongyloides westeri*; **Summer Sores** caused by *Habronema* and *Draschia* spp. cutaneous third-stage larvae; Dermatitis caused by neck threadworm microfilariae, *Onchocerca* sp.

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DOSAGE:
This syringe contains sufficient paste to treat one 1250 lb horse at the recommended dose rate of 91 mcg ivermectin per lb (200 mcg/kg) body weight.

Each weight marking on the syringe plunger delivers enough paste to treat 250 lb body weight.

Do not underdose. Ensure each animal receives a complete dose based on a current body weight. Underdosing may result in ineffective treatment, and encourage the development of parasite resistance.

ADMINISTRATION:

(1) While holding plunger, turn the knurled ring on the plunger 1/4 turn to the left and slide it so the side nearest the barrel is at the prescribed weight marking. (2) Lock the ring in place by making a 1/4 turn to the right. Ensure it is locked (it should no longer slide). (3) Make sure that the horse's mouth contains no feed. (4) Remove the cover from the tip of the syringe. (5) Insert the syringe tip into the horse's mouth at the space between the teeth. (6) Depress the plunger as far as it will go, depositing paste on the back of the tongue. (7) Immediately raise the horse's head for a few seconds after dosing.

PARASITE CONTROL PROGRAM:

All horses should be included in a regular parasite control program with particular attention being paid to mares, foals and yearlings. Foals should be treated initially at 6 to 8 weeks of age, and routine treatment repeated as appropriate. Consult your veterinarian for a control program to meet your specific needs. Ivermectin Paste 1.87% effectively controls gastrointestinal nematodes and bots of horses. Regular treatment will reduce the chances of verminous arteritis caused by *Strongylus vulgaris*.

PRODUCT ADVANTAGES:
Broad-spectrum Control– Ivermectin Paste 1.87% kills important internal parasites, including bots and the arterial stages of *S. vulgaris*, with a single dose.

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Ivermectin Paste 1.87% is a potent antiparasitic agent that is neither a benzimidazole nor an organophosphate.

ANIMAL SAFETY:

Ivermectin Paste 1.87% may be used in horses of all ages, including mares at any stage of pregnancy. Stallions may be treated without adversely affecting their fertility.

WARNING:

Do not use in horses intended for human consumption. Not for use in humans. Keep this and all drugs out of reach of children.

Refrain from smoking and eating when handling. Wash hands after use. Avoid contact with eyes.

The Safety Data Sheet (SDS) contains more detailed occupational safety information. To report adverse reactions in users, to obtain more information, or to obtain an SDS, contact Dechra at (866) 933-2472.

OTHER WARNINGS:

Parasite resistance may develop to any dewormer, and has been reported for most classes of dewormers. Treatment with a dewormer used in conjunction with parasite management practices appropriate to the geographic area and the animal(s) to be treated may slow the development of parasite resistance. Fecal examinations or other diagnostic tests and parasite management history should be used to determine if the product is appropriate for the herd, prior to the use of any dewormer. Following the use of any dewormer, effectiveness of treatment should be monitored (for example, with the use of a fecal egg count reduction test or another appropriate method). A decrease in a drug's effectiveness over time as calculated by fecal egg count reduction tests may indicate the development of resistance to the dewormer administered.

Your parasite management plan should be adjusted accordingly based on regular monitoring.

PRECAUTIONS:
Ivermectin Paste 1.87% has been formulated specifically for use in horses **only**. This product should not be used in other animal species as severe adverse reactions, including fatalities in dogs, may result.

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Environmental Safety:

Ivermectin and excreted ivermectin residues may adversely affect aquatic organisms. Do not contaminate ground or surface water. Dispose of the syringe in an approved landfill or by incineration.

INFORMATION FOR HORSE OWNERS:

Swelling and itching reactions after treatment with Ivermectin Paste 1.87% have occurred in horses carrying heavy infections of neck threadworm microfilariae (*Onchocerca* sp.). These reactions were most likely the result of microfilariae dying in large numbers. Symptomatic treatment may be advisable. Consult your veterinarian should any such reactions occur. Healing of summer sores involving extensive tissue changes may require other appropriate therapy in conjunction with treatment with Ivermectin Paste 1.87%. Reinfection, and measures for its prevention, should also be considered. Consult your veterinarian if the condition does not improve.

HOW SUPPLIED:

Each syringe contains 0.21 oz (6.08 g) apple flavor ivermectin paste.

Contents will treat up to 1250 lb body weight.

NDC # 17033-455-60

STORAGE INFORMATION:

Store at 68°F - 77°F (20°C - 25°C). Excursions between 59°F - 86°F (15°C - 30°C) are permitted.

Approved by FDA under ANADA # 200-390

For additional information about adverse drug experience reporting for animal drugs, contact FDA at 1-888-FDA-VETS

or online at <http://www.fda.gov/reportanimalae>

Manufactured by:

Dechra Veterinary Products

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Overland Park, KS 66211 USA

Rev. January 2023



Penny Rousseau 18JAN2023

Product:Ivermectin Paste 1.87% - US - Leaflet	Proof:	Date:	Proof:	Date:
Dimensions:..... 5.5" x 5.8125"	1.1 (NH)	02-12-2022		
Primary brand name font size:23pt	1.2 (NH)	02-12-2022		
Established name font size:14pt	1.3 (JP)	16-12-2022		
Primary brand description font size:10pt	1.4 (NH)	04-01-2023		
Body text font size:9pt	1.5 (NH)	05-01-2023		
Item code:Rev. January 2023	1.6 (JP)	13-01-2023		
Pharmacode:N/A	1.7 (JP)	13-01-2023		

Pantone reference guide

Colours to be printed:

BLACK

Do not print

CUTTER GUIDE

STYLE DEVIATIONS

Dechra logos reduced to 80% due to them looking too large.

REGULATORY AUTHORITIES' REQUESTS

