

## **DURAMECTIN- ivermectin paste**

**Durvet**

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**For Oral Use in Horses Only**

**DuraMectin**

**(ivermectin paste)1.87%**

***Anthelmentic & Boticide***

**Removes worms and bots with a single dose**

**Net Wt. 0.21 oz (6.08g)**

**Contents will treat up to 1250 lb body weight**

**Approved by FDA under ANADA # 200-326**

**INDICATIONS:** Consult your veterinarian for assistance in the diagnosis, treatment, and control of parasitism. DURAMECTIN (ivermectin paste) 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.

**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.

**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. DURAMECTIN (ivermectin paste) 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** - DURAMECTIN kills important internal parasites, including bots and the arterial stages of *S. vulgaris*, with a single dose. DURAMECTIN is a potent antiparasitic agent that is neither a benzimidazole nor an organophosphate.

**ANIMAL SAFETY:** DURAMECTIN (ivermectin paste) may be used in horses of all ages, including mares at any stage of pregnancy. Stallions may be treated without adversely affecting their fertility.

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

**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, aligning the arrow on the ring to the line between the weight and lbs, as shown in the pictogram.**

**(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.



**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 suspected adverse drug events, for technical assistance, or to obtain a copy of the SDS contact Durvet, Inc. at 1-800-821-5570. For additional information about adverse drug experience reporting for animal drugs, contact FDA at 1-888-FDA-VETS or [www.fda.gov/reportanimalae](http://www.fda.gov/reportanimalae).

**PRECAUTIONS:** DURAMECTIN (ivermectin paste) 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.

**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.

**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 DURAMECTIN (ivermectin paste) have occurred in horses carrying heavy infections of neck threadworm (*Onchocerca* sp.) microfilariae. 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 DURAMECTIN. Reinfection, and measures for its prevention, should also be considered. Consult your veterinarian if the condition does not improve.

**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 suspected adverse drug events, for technical assistance or to obtain a copy of the SDS, contact Durvet, Inc. at 1-800-981-5578. For additional information about adverse drug experience reporting for animal drugs, contact FDA at 1-888-FDA-VETS or www.fda.gov/reportanimal. **PRECAUTIONS:** DURAMECTIN (ivermectin paste) 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. **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. **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 DURAMECTIN (ivermectin paste) have occurred in horses carrying heavy infections of neck threadworm (*Onchocerca* sp.) microfilariae. 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 DURAMECTIN. Reinfection, and measures for its prevention, should also be considered. Consult your veterinarian if the condition does not improve.

**durvet**

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(ivermectin paste) 1.87%

Anthelmintic and Boticide

Removes worms and bots with a single dose

Net Wt. 0.21 oz (6.08 g)  
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Manufactured by: BiMeda-MTC Animal Health Inc.  
Cambridge, Ontario, Canada N3C 2W4  
Distributed by: Durvet Inc., Blue Springs, MO 64014, www.Durvet.com  
Rev. 02.21 00UR000

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**DOSE:** This syringe contains sufficient paste to treat one (1250 lb) horse at the recommended dose rate of 51 mcg ivermectin per lb (100 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.

**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. DURAMECTIN (ivermectin paste) effectively controls gastrointestinal nematodes and bots of horses. Regular treatment will reduce the chances of verminous arteritis caused by *Strongylus vulgaris*.

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

**ADMINISTRATION:**

- (1) **White Anting plunger:** Turn the knurled ring on the plunger 1/4 turn to the left and slide it to the side nearest the barrel so the prescribed weight markings, aligning the arrow on the ring to the line between the weight and lbs, as shown in the pictogram.
- (2) Lock the ring in place by moving a 1/8 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.

## DURAMECTIN

ivermectin paste

### Product Information

Product Type

OTC ANIMAL DRUG

Item Code (Source)

NDC:30798-837

**Route of Administration** ORAL

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
IVERMECTIN (UNII: 8883YP2R6D) (IVERMECTIN - UNII:8883YP2R6D)	IVERMECTIN	18.7 mg in 1 g

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:30798-837-81	6.08 g in 1 SYRINGE		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANADA	ANADA200326	01/17/2007	

**Labeler** - Durvet (056387798)

**Registrant** - Bimeda Inc. (060492923)

### Establishment

Name	Address	ID/FEI	Business Operations
Bimeda-MTC		256232216	manufacture

Revised: 4/2022

Durvet