

Equine Anthelmintic Feed
(pyrantel tartrate Type C medicated feed)

INDICATIONS

For the prevention of *Strongylus vulgaris* larval infestation in horses.

For control of the following parasites in horses:

Large Strongyles (adults): *S. vulgaris*, *S. edentatus*

Small Strongyles (adults and fourth-stage larvae): *Cyathostomum* spp., *Cylicocyclus* spp.,
Cylicostephanus spp., *Cylicodontophorus* spp., *Poteriostomum* spp., *Triodontophorus* spp.

Pinworms (adults and fourth-stage larvae): *Oxyuris equi*

Ascarids (adults and fourth-stage larvae): *Parascaris equorum*

ACTIVE DRUG INGREDIENT

pyrantel tartrate^a 120 to 1,200 grams/ton*
[0.06 to 0.6 grams/lb*]

GUARANTEED ANALYSIS

Crude Protein, min	_____ %
Crude Fat, min	_____ %
Crude Fiber, max	_____ %
Acid Detergent Fiber (ADF), max	_____ %
Neutral Detergent Fiber (NDF), max	_____ %
Calcium, min	_____ %
Calcium, max	_____ %
Phosphorus, min	_____ %
Copper (if added), min	_____ ppm
Selenium, min	_____ ppm
Zinc, min	_____ ppm
Vitamin A, other than the precursors of Vitamin A, (if added), min	_____ I.U./lb

INGREDIENTS

Each ingredient must be specifically named in accordance with the names and definitions adopted by the Association of American Feed Control Officials.

FEEDING DIRECTIONS

Feed on a continuous basis at the rate of 1.2 mg pyrantel tartrate per lb. of body weight daily. To achieve this dose, administer ____ lb. of pyrantel tartrate Type C medicated feed per 100 lb. of body weight.

See the table below for an examples of the amount of pyrantel tartrate Type C medicated feed (medicated grain) to be fed per 100 lb. body weight.

Pyrantel Tartrate Type C Medicated Feed (Medicated Grain) Feeding Directions

Concentration Grams/Ton	Lb. of Medicated Grain Mix per 100 lb. of Body Weight
120	2.0
160	1.5
240	1.0
480	0.5
1,200	0.2

Pyrantel tartrate Type C medicated feed should be administered for the entire period that the animal is at risk to internal parasites. Unprotected animals that have grazed may already have an established *S. vulgaris* larval infestation. Before administering pyrantel tartrate Type C medicated feed, these animals should be treated with a therapeutic dose of a larvicidal product.

Foals may be administered pyrantel tartrate Type C medicated feed as soon as consistent intake of grain mix is occurring. This is generally between 2–3 months of age.

Pyrantel tartrate Type C medicated feed may be used in mares at any stage of pregnancy or lactation. Stallion fertility is not affected by the use of pyrantel tartrate Type C medicated feed.

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.

WARNINGS

Do not use in horses intended for human consumption.

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.

Consult your veterinarian for assistance in the diagnosis, treatment, and control of parasitism.

CAUTION

Consult your veterinarian before using in severely debilitated animals and for assistance in the diagnosis, treatment, and control of parasitism. Do not mix in feeds containing bentonite.

QUESTIONS/COMMENTS

To report suspected adverse events or side effects, or for technical assistance contact Zoetis Inc. at 1-888-963-8471 or www.zoetis.com. For additional information about reporting side effects for animal drugs, contact FDA at 1-888- FDA-VETS or <http://www.fda.gov/reportanimalae>.

Net Weight _____ lb (_____ kg) on bag or bulk

Lot Number (if applicable): _____
Expiration dating not required.

Approved by FDA under NADA # 140-819

MANUFACTURED BY

Blue Bird Feed Company
Robin, Indiana 11111

*The final printed feed label should list only a single drug concentration.

^aSourced from STRONGID[®] C 2X[™] Type C medicated feed (NADA # 140-819).

Zoetis Inc. [10 October 2023]