PANACUR- fenbendazole suspension
Merck Sharp & Dohme Corp.

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panacur®
(fenbendazole)
Horse & Cattle Dewormer

INDICATIONS AND DOSAGE:

Horses – 2.3 mg/lb (5 mg/kg): for the treatment and control of large strongyles (*Strongylus edentatus, S. equinus, S. vulgaris, Triodontophorus* spp.), small strongyles (*Cyathostomum* spp., *Cylicocyclus* spp., *Cylicostephanus* spp., *Cylicodontophorus* spp.), and pinworms (*Oxyuris equi*). 4.6 mg/lb (10 mg/kg): for the treatment and control of ascarids (*Parascaris equorum*).

Beef and Dairy Cattle – 2.3 mg/lb (5 mg/kg): for the treatment and control of: **Lungworms:** Adult *Dictyocaulus viviparus*; **Stomach worms:** Adult brown stomach worms (*Ostertagia ostertagi*), Adult and fourth stage larvae barberpole worms (*Haemonchus contortus* & *H. placei*) and Adult and fourth stage larvae small stomach worms (*Trichostrongylus axei*); **Intestinal worms** (Adult and fourth stage larvae): hookworms (*Bunostomum phlebotomum*), thread-necked intestinal worms (*Nematodirus helvetianus*), small intestinal worms (*Cooperia punctata* & *C. oncophora*), bankrupt worms (*Trichostrongylus colubriformis*), and nodular worms (*Oesophagostomum radiatum*).

Beef Cattle Only – 4.6 mg/lb (10 mg/kg): for the treatment and control of: **Stomach worms** (4th stage inhibited larvae): *Ostertagia ostertagi* (Type II Ostertagiasis); **Tapeworms:** *Moniezia benedeni.*

Do not use in dairy cattle at 10 mg/kg.

Shake well before use.

Store at or below 25°C (77°F)

Protect from freezing.

DIRECTIONS:

Horses and Beef and Dairy Cattle:

Administer orally by suitable dosing syringe. Insert nozzle of syringe through the interdental space and deposit the drug on the back of the tongue by depressing the plunger. The drug may also be administered by stomach tube.

EXAMPLES:

Animal Weight	Dose (5 mg/kg)	Dose (10 mg/kg)

100 lb	**2.3 mL**	**4.6 mL**
200 lb	**4.6 mL**	**9.2 mL**
300 lb	**6.9 mL**	**13.8 mL**
400 lb	**9.2 mL**	**18.4 mL**
500 lb	**11.5 mL**	**23.0 mL**
1000 lb	**23.0 mL**	**46.0 mL**
1500 lb	**34.5 mL**	**69.0 mL**

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: NOT FOR USE IN HUMANS. KEEP OUT OF REACH OF CHILDREN. The Safety Data Sheet (SDS) contains more detailed occupational safety information. For customer service, adverse effects reporting, and/or a copy of the SDS, call 1-800-211-3573. For additional information about adverse drug experience reporting for animal drugs, contact FDA at 1-888-FDAVETS, http://www.fda.gov/reportanimalae.

OTHER 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.

Fenbendazole (active ingred.) made in: see imprint.
Formulated in France.
Distributed by:
Intervet Inc. (d/b/a Merck Animal Health) Madison, NJ 07940
Approved by FDA under NADA # 104-494
Approved by FDA under NADA # 128-620

Rev. 03/22

API
Made in:

LOT
NUMBER:

EXPIRATION
DATE:

312522 R11

PRINCIPAL DISPLAY PANEL - 1,000 mL Bottle Label

panacur®
(fenbendazole)

Horse & Cattle
Dewormer

Suspension 10%
(100 mg/mL)

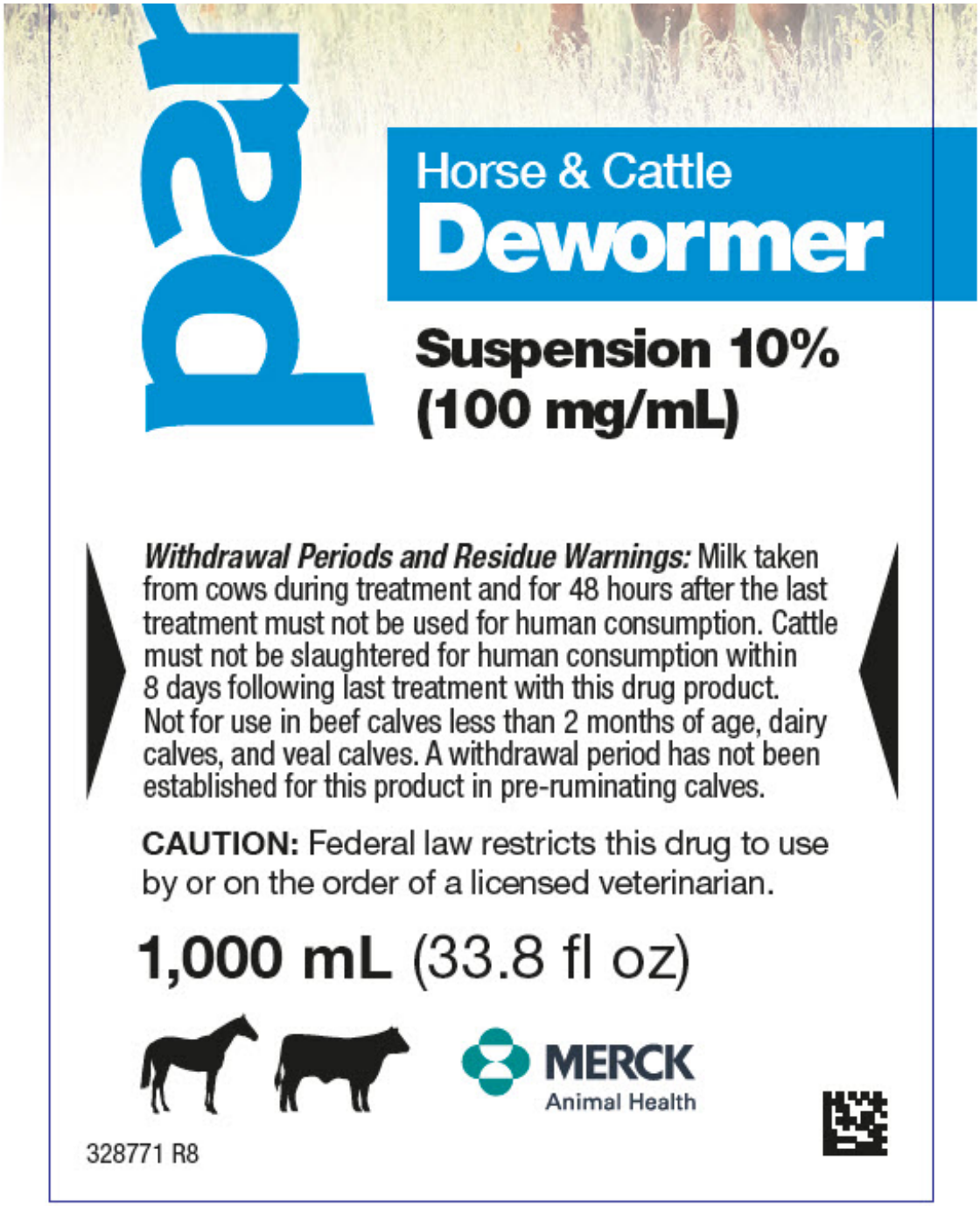
Withdrawal Periods and Residue Warnings: Milk taken from cows during treatment and for 48 hours after the last treatment must not be used for human consumption. Cattle must not be slaughtered for human consumption within 8 days following last treatment with this drug product. Not for use in beef calves less than 2 months of age, dairy calves, and veal calves. A withdrawal period has not been established for this product in pre-ruminating calves.

CAUTION: Federal law restricts this drug to use by or on the order of a licensed veterinarian.

1,000 mL (33.8 fl oz)

MERCK
Animal Health

328771 R8



PANACUR

fenbendazole suspension

Product Information			
Product Type	PRESCRIPTION ANIMAL DRUG	**Item Code (Source)**	NDC:57926-087
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Fenbendazole (UNII: 621BVT9M36) (Fenbendazole - UNII:621BVT9M36)	Fenbendazole	100 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
methylparaben (UNII: A2I8C7HI9T)	
propylparaben (UNII: Z8IX2SC1OH)	
silicon dioxide (UNII: ETJ7Z6XBU4)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
trisodium citrate dihydrate (UNII: B22547B95K)	
citric acid monohydrate (UNII: 2968PHW8QP)	
water (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:57926-087-01	1000 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NADA	NADA128620	11/15/1977	

Labeler - Merck Sharp & Dohme Corp. (001317601)

Establishment

Name	Address	ID/FEI	Business Operations
Intervet Production S.A.		771867553	ANALYSIS, MANUFACTURE

Establishment

Name	Address	ID/FEI	Business Operations
Zhejiang Apeloa Kangyu Pharmaceutical Co., Ltd		420823163	API MANUFACTURE

Revised: 11/2023 Merck Sharp & Dohme Corp.