

**RUMATEL 88- morantel tartrate powder**  
**Phibro Animal Health**

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**Rumatel® 88**  
**(morantel tartrate)**  
**TYPE A MEDICATED ARTICLE**

*For cattle and goats*

**Active Drug Ingredient:**

Morantel tartrate . . . . . 19.4% (88 g/lb)

**Indications for Use:**

*Cattle:* For the removal and control of mature gastrointestinal nematode infections of cattle including stomach worms (*Haemonchus* spp., *Ostertagia* spp., *Trichostrongylus* spp.), worms of the small intestine (*Cooperia* spp., *Trichostrongylus* spp., *Nematodirus* spp.), and worms of the large intestine (*Oesophagostomum radiatum*).

*Goats:* For the removal and control of mature gastrointestinal nematode infections of goats including *Haemonchus contortus*, *Ostertagia (Teladorsagia) circumcincta*, and *Trichostrongylus axei*.

**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/flock, 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 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.**

**CAUTION:**

For use in the manufacture of medicated beef, dairy, and goat feeds.

**CAUTION:** Certain components of animal feeds, including medicated premixes, possess properties that may be a potential health hazard or a source of personal discomfort to certain individuals who are exposed to them. Human exposure should, therefore, be minimized by observing the general industry standards for occupational health and safety.

Precautions such as the following should be considered: dust masks or respirators and protective clothing should be worn; dust-arresting equipment and adequate ventilation should be utilized; personal hygiene should be observed; wash before eating or leaving a work site; be alert for signs of allergic reactions—seek prompt medical treatment if such reactions are suspected.

**Mixing and Use Directions**

The following are examples in the approved range (0.44–4.4 g/lb)

lb of feed per 100 lb of body weight	lb of premix	lb of nonmedicated feed	Resulting concentration (g/lb)
1.0	10	1990	0.44
0.4	25	1975	1.10
0.2	50	1950	2.20
0.1	100	1900	4.40

### **Directions for Use of Medicated Ration**

Use a single therapeutic treatment. Medicated feed is to be fed at the rate of 0.44 grams of morantel tartrate per 100 lb of body weight. The medicated feed mix should be consumed within 6 hours. May be fed as the sole ration or mixed with 1–2 parts of complete feed or as a top dress. When used as a top dress the medication as well as the underlying feed should be evenly distributed. Animals should be grouped by size for optimum efficacy. Fresh water should be available at all times. When all medicated feed is consumed resume normal feeding. Conditions of constant worm exposure may require retreatment within 2–4 weeks. Consult your veterinarian for assistance in the diagnosis, treatment, and control of parasitism.

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

### **WARNINGS:**

Do not treat cattle within 14 days of slaughter.

Do not treat goats within 30 days of slaughter. No milk discard required following use in dairy cattle or goats.

**CAUTION:** Consult veterinarian before using in severely debilitated animals. Do not mix in feeds containing bentonite.

Restricted Drug (California) – USE AS DIRECTED

Store At or Below 25°C(77°F), Excursions Permitted Up to 40°C (104°F)

Not For Human Use

SEE BACK PANEL FOR FURTHER USE DIRECTIONS

Net Weight: 25 lb (11.3 kg)

Approved by FDA under NADA #092-444

7970000

101-8318-06B

Made in USA

# Rumatel<sup>®</sup> 88

(morantel tartrate)

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MEDICATED ARTICLE  
ANTHELMINTIC

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ANIMAL HEALTH CORPORATION TM

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Distributed by  
**Phibro**  
ANIMAL HEALTH CORPORATION  
Teaneck, NJ 07666



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morantel tartrate powder

**Product Information**

<b>Product Type</b>	OTC TYPE A MEDICATED ARTICLE ANIMAL DRUG	<b>Item Code (Source)</b>	NDC:66104-2400	
<b>Route of Administration</b>	ORAL			
<b>Active Ingredient/Active Moiety</b>				
	<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>	
	MORANTEL TARTRATE (UNII: 5WF7E9QC3F) (MORANTEL - UNII:7NJ031HAX5)	MORANTEL TARTRATE	88 g in 0.45 kg	
<b>Inactive Ingredients</b>				
	<b>Ingredient Name</b>		<b>Strength</b>	
	MINERAL OIL (UNII: T5L8T28FGP)			
	SODIUM ALUMINO SILICATE (UNII: 058TS43PSM)			
	CALCIUM CARBONATE (UNII: H0G9379FGK)			
	SOYBEAN (UNII: L7HT8F1ZOD)			
<b>Packaging</b>				
<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:66104-2400-5	11.3 kg in 1 BAG		
<b>Marketing Information</b>				
<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>	
NADA	NADA092444	03/25/2010		

**Labeler** - Phibro Animal Health (006989008)

**Registrant** - Phibro Animal Health (006989008)

Revised: 6/2019

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